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Exploring the dietary intake and eating patterns of New Zealand European women aged 16-45 years

A thesis presented in partial fulfilment of the requirements for the degree of

Masters of Science

in

Nutrition and Dietetics

at Massey University, Albany New Zealand

> Jenna Kate Schrijvers 2015

Abstract

Background/Aim: Analysing dietary intakes gives insight to an individual or groups nutritional status. Investigating dietary patterns provides an alternative measure to identify combinations of foods that are related to excess adiposity. The aim of this study is to investigate dietary intakes and eating patterns of New Zealand European (NZE) women with different body composition profiles, participating in the women's EXPLORE (Examining the Predictors Linking Obesity Related Elements) study.

Methods: Post-menarche, pre-menopausal NZE women (16-45 years) (n=231) completed a validated 220-item, self-administrated, semi-quantitative food frequency questionnaire (FFQ) assessing dietary intake over the previous month. Quetelet's body mass index (BMI) was calculated (kg/m²) from height and weight measurements; body fat percentage (BF%) was measured using air displacement plethysmography (BodPod). Participants were categorised into one of three body composition profile (BCP) groups: normal BMI (18.5-24.9 kg/m²), normal BF% (\geq 22%, <30%) (HH); normal BMI, high BF% (\geq 30%)(NH); high BMI (\geq 25 kg/m²), high BF% (HH). Dietary intakes, macronutrient profiles and diet quality for the total NZE women and the BCP groups were analysed. Dietary patterns were identified using principal component factor analysis and broken into tertiles (T1, T2, T3). Associations between dietary patterns, age, BMI and BF% were investigated.

Results: Many NZE women consumed insufficient vitamin D (55%), iron (82%), calcium (28.5%), folate (48%) and dietary fibre (28%) intakes. Mean±SD percentage of energy intake for carbohydrate (41.9±7%) was below and for saturated fat (13.9±3.5%) above the acceptable macronutrient distribution range for the total NZE women. The top 40 food items consumed by the NZE women included water, bread, tea, coffee, milk and yoghurt. Diet soft drinks were only present in the HH BCP group. Four dietary patterns were identified: P1: 'Snacking' pattern; P2: 'Energy-dense meat' pattern; P3: 'Fruit and vegetable' pattern; P4: 'Healthy' pattern, which explained 6.9, 6.8, 5.6 and 4.8% of variation in food intake, respectively. Younger (16-24 years) (P=0.035) and overweight (26.4±26.7kg/m²) (P=0.036) women were significantly associated with P2, loading highly in T3. No significant associations were found with BF%. Intakes of vitamin A, E, D, and zinc were comparable between normal BF% and high BF% BCP groups.

I

Conclusion: NZE women consume inadequate iron, vitamin D, folate, calcium and dietary fibre intakes irrespective of body fatness. Dietary patterns of NZE women can be linked to specific body compositions, specifically, women with a high BMI high BF% were associated with a diet characteristic of meat, high fat sauces, puddings and fried foods. Regardless of BF%, NZE women follow a diet low in carbohydrate and high in saturated fat. Diet quality of vitamin A, D, E, iron, and zinc in women with a high BF% is comparable to that of women with normal BF%'s showing good diet quality. Targeted interventions can be developed based on these findings to increase nutrient intakes of NZE women and improve the health status of those with excess adiposity.

Key words: Dietary intake, dietary patterns, factor analysis, food frequency questionnaire

Acknowledgements

There are a number of people I would like to thank for their involvement in this thesis project.

Firstly, I would like to thank all of the women who participated in this research, who were part of the wider EXPLORE study, without you this would not have been possible.

Thank you to my supervisors, Rozanne Kruger and Sarah McNaughton, for your extensive knowledge, wisdom, dedication and support.

Thank you to Wendy O'Brien and Shakeela Jayasinghe for your commitment and all the work you did with recruiting, screening and testing.

Thank you to the EXPLORE team Sarah Philipsen, Owen Mugridge, PC Tong, Pam Von Hurst, Kathryn Beck, Cath Conlon, Richard Swift, AJ Hepburn and Zara Houston. It was a pleasure working with you all.

To Maria Casale, we got through the early mornings and the endless hours of recruitment, screening, phone calls and data entry. Thank you for your on-going support, encouragement and friendship throughout these past two years.

To my second family in Auckland: Sue, Robert and Merryn. Thank you so much for your love, support, generosity and delicious home cooked meals.

Lastly, thank you to all of my friends and family who have provided me with their on-going support. I would not have been able to do this without you.

Table of Contents

Abstract	I
Acknowledgements	
Table of Contents	IV
List of Tables	VII
List of Figures	IX
List of Appendices	X
Abbreviation List	XI
Chapter 1 Introduction	1
1.1 Background	
1.2 Purpose of the Study	6
1.2.1 Aim	6
1.2.2 Objectives	6
1.3 Thesis Structure	6
1.4 Researcher's Contributions to the Study	7
Chapter 2 Literature Review	8
2.1 Obesity and Health	
2.1.1 Obesity definition	
2.1.2 Obesity prevalence in New Zealand	
2.1.3 Causes of obesity	9
2.1.4 Obesity and chronic disease	
2.1.5 Assessing body fatness	
2.2 Dietary Patterns	15
2.2.1 Energy dense diets	
2.2.2 Sugar sweetened beverages	
2.2.3 Fast-food consumption	
2.3 Macronutrient Distribution	43
2.3.1 Compliance with recommendations	
2.4 Diet Quality	45
2.4.1 Micronutrient status	

2.4.2 Vitamin A
2.4.3 Vitamin D
2.4.4 Iron57
2.4.5 Vitamin E61
2.4.6 Zinc61
2.4.7 Vitamin C62
2.5 Assessing dietary intake
2.5.1 Misreporting of energy intake67
2.5.2 Recovery markers
2.5.3 Goldberg cut-off

Chapter 3 Methods	9
3.1 EXPLORE Study Design	
3.2 Ethical Approval70	
3.3 Study Population70	
3.3.1 Participants	
3.3.2 Recruitment70	
3.4 Procedures72	
3.4.1 Phase 172	
3.4.2 Phase 272	
3.5 Body Composition73	
3.6 Dietary Questionnaires74	
3.6.1 Food frequency questionnaire74	
3.7 Data Analysis75	
3.8 Food Groupings76	
3.9 Data Accuracy	
3.10 Statistical Analysis	
3.10.1 Frequency of food item consumption80	
3.10.2 Food pattern derivation81	

Chapter 4.0 Results	
4.1 Study Population	.83
4.2 Participant Characteristics	.85
4.3 Dietary Analysis	.86

4.3.1 Total NZE population	5
4.3.2 Body composition profile groups	3
4.4 Top 40 Food Items	3
4.5 Dietary Pattern Analysis)
4.6 Diet Quality	3

Chapter 5 Discussion	109
5.1 Participant Characteristics	. 109
5.2 Dietary Intake Analysis	. 110
5.2.1 Total population	. 110
5.2.2 BCP groups	. 116
5.2.3 Macronutrient distribution	. 119
5.3 Top 40 Food Items	. 123
5.4 Dietary Patterns	. 126
5.5 Diet Quality	. 133

Chapter 6 Conclusions	
6.1 Aim of the research	
6.2 Main findings and Conclusions	
6.3 Study strengths	
6.4 Study limitations	
6.5 Recommendations	
6.6 Conclusion	

References	
Appendices	

List of Tables

Table 1.2 Researchers contributions to the study. 7
Table 2.1 Mean BMI and prevalence of overweight and obesity in New Zealand European
females and the total population from 2006/07 to 2013/149
Table 2.2 Body Mass Index (BMI) classification
Table 2.3 Body fat percentage (BF%) categories for women over 18 years of age13
Table 2.4. Strengths and weaknesses of the main body composition assessment methods 14
Table 2.5 Studies investigating dietary patterns and body fatness. 18
Table 2.6 Studies investigating the relationships between dietary characteristics and body
fatness
Table 2.7 Studies investigating sugar-sweetened beverage (SSB) consumption
Table 2.8 Studies investigating the effects of fast-food consumption. 40
Table 2.9 Studies investigating the relationship between diet quality and body composition46
Table 2.10 Studies investigating the relationship between vitamin A status and body
composition
Table 2.11 Studies investigating the relationship between vitamin D status and body
composition55
Table 2.12 Studies investigating the relationship between iron status and body composition. 59
Table 2.13 Studies investigating vitamin E, zinc, vitamin C and their relationships with body
composition63
Table 2.14. Strengths and weaknesses of the five main dietary assessment techniques
Table 3.1 Body composition profile (BCP) groups69
Table 3.2 Frequency translations. 76
Table 3.3 Food groups 77
Table 3.4 Scholfield equations. 80
Table 4.1 Study population. 84
Table 4.2 Characteristics of all New Zealand European study participants (n = 231)85
Table 4.3 Characteristics of participants compared between body composition profile (BCP)
groups
Table 4.4 Mean daily dietary intakes from the NZWFFQ (n = 207) vs. New Zealand National
recommendations and percentage of NZE women below the recommendations87
Table 4.5 Nutrient analysis comparison between body composition profile (BCP) groups and
with the estimated average nutrient requirements (EAR)

Table 4.6 Mean macronutrient percentage intakes (%) compared between body composition
profile (BCP) groups
Table 4.7 Top 40 foods consumed for the total population
Table 4.8 Top 40 foods compared between the three main body composition profile (BCP)
groups
Table 4.9 Factor loading matrix for the four dietary patterns identified in NZE women (n=231).
Table 4.10 Cronbach's α score for each of the dietary patterns
Table 4.11 Age, body mass index (BMI) and body fat percentage (BF%) characteristics of the
tertiles in the dietary patterns for the NZE women (n = 231)
Table 4.12 Socio-demographic characteristics of the NZE women in the tertiles of each dietary
pattern
Table 4.13 Nutrient intakes within the dietary pattern tertiles identified from the NZWFFQ
among NZE women
Table 4.14 Macronutrient distribution between dietary patterns tertiles. 108

List of Figures

Figure 1.1 The relationship between diet, body fat percentage and health outcomes
Figure 2.1 The Socio-Ecological Model of factors influencing increased adiposity10
Figure 2.2 The medical complications associated with obesity11
Figure 2.3 The prevalence of vitamin D deficiency by body size in adults over 15 years of age
(unadjusted prevalence)53
Figure 3.1 Methodological overview of the study73
Figure 3.2 Food Frequency Questionnaire (FFQ) example questions explained to participant
prior to completion
Figure 4.1 Outline of the recruitment process
Figure 4.2 Mean (SD) macronutrient intakes compared between the body composition profile
groups
Figure 4.3 Mean (SD) vitamin D intakes of the body composition profile (BCP) groups91
Figure 4.4 Mean (SD) vitamin B12 intakes of the body composition profile (BCP) groups91
Figure 4.5 Mean (SD) calcium intakes of the body composition profile (BCP) groups91
Figure 4.6 Mean (SD) iron intakes of the body composition profile (BCP) groups91
Figure 4.7 Mean (SD) sodium intakes compared between the body composition profile (BCP)
groups91
Figure 4.8 Mean (SD) intake of dietary fibre compared between the body composition profile
(BCP) groups91
Figure 4.9 Mean (SD) vitamin A intakes of the body composition profile (BCP) groups92
Figure 4.10 Mean (SD) vitamin C intakes of the body composition profile (BCP) groups92
Figure 4.11 Mean (SD) vitamin E intakes of the body composition profile (BCP) groups92
Figure 4.12 Mean (SD) total folate intakes of the body composition profile (BCP) groups92
Figure 4.13 Mean (SD) macronutrient intakes as percentage of energy intake compared
between the body composition profile (BCP) groups

List of Appendices

Appendix A. Pre-screening health and demographic questionnaire	173
Appendix B. Standard operating procedures for the food frequency questionnaire	177
Appendix C. Food frequency questionnaire	181
Appendix D. Assumptions made when entering the New Zealand Women's Food Freque	ency
Questionnaire (NZWFFQ)	198

Abbreviation List

ADP	Air Displacement Plethysmography
BCP group	Body Composition Profile Group
BF%	Body Fat Percentage
ВР	Blood Pressure
BMR	Basal Metabolic Rate
BIA	Bioelectrical Impedance Analysis
BMI	Body Mass Index
CHD	Coronary Heart Disease
CVD	Cardiovascular Disease
СНО	Carbohydrate
DASH	Dietary Approach to Stop Hypertension
DQ	Diet Quality
DXA	Dual Energy X-ray Absorptiometry
EXPLORE study	Examining Predictors Linking Obesity Related Elements
FFQ	Food Frequency Questionnaire
GI	Glycaemic Index
GL	Glycaemic Load
НС	Hip Circumference

HFCS	High Fructose Corn Syrup
нн	High Body Mass Index; High Body Fat Percentage
HN	High Body Mass Index; Normal Body Fat Percentage
КМО	Kaiser-Meyer-Olkin
LDL-c	Low Density Lipoprotein Cholesterol
МАМС	Mid Arm Muscle Circumference
NN	Normal Body Mass Index; Normal Body Fat Percentage
NH	Normal Body Mass Index; High Body Fat Percentage
NL	Normal Body Mass Index; Low Body Fat Percentage
NZE	New Zealand European
NZWFFQ	New Zealand Women's Food Frequency Questionnaire
NZWFFQ PCA	New Zealand Women's Food Frequency Questionnaire Principal Component Analysis
РСА	Principal Component Analysis
PCA SSB	Principal Component Analysis Sugar- Sweetened Beverage
PCA SSB T2DM	Principal Component Analysis Sugar- Sweetened Beverage Type 2 Diabetes Mellitus
PCA SSB T2DM TC	Principal Component Analysis Sugar- Sweetened Beverage Type 2 Diabetes Mellitus Total Cholesterol

Chapter 1 Introduction

1.1 Background

The 2008/09 New Zealand Adult Nutrition Survey identified 32.8% and 27.8% of females over the age of 15 years were overweight and obese, respectively (University of Otago and Ministry of Health, 2011). Prevalence of obesity has significantly risen since 1997 (females 20.6%) in both New Zealand and worldwide (University of Otago and Ministry of Health, 2011, World Health Organisation, 2015). It is well-established that obesity is a major risk factor for many health conditions including but not limited to cardiovascular disease, type 2 diabetes, hypertension, insulin resistance, dyslipidaemia, osteoarthritis, sleep apnoea, psychological and social problems and some cancers (University of Otago and Ministry of Health, 2011, World Health Organisation, 2015, Stein, 2004, Oliveros, 2014, Poirier *et al.*, 2006, Pi-Sunyer, 2002, McMichael, 2008).

To determine obesity, Body Mass Index (BMI) is calculated. This is an easy non-invasive calculation using height (m) and weight (kg) measurements. According to the Ministry of Health overweight and obesity are defined as having a BMI score between 25 kg/m² and 29.9 kg/m^2 , and greater than 30 kg/m^2 , respectively (Ministry of Health, 2008). Although routinely used in epidemiological studies and by health professionals, BMI is an imperfect measure of body fatness due to indirectly measuring body composition and assuming excess body fat to be present in overweight and obese individuals (University of Otago and Ministry of Health, 2011). It speculates that at any given height, a higher weight correlates to a larger body fat percentage (BF%) and consequently a higher risk of morbidity and mortality (Gallagher et al., 2000). The BMI calculation has good specificity but poor sensitivity and as such, some individuals for example athletes, who have a high lean muscle mass are incorrectly classed as overweight, whereas others are classed as having a normal BMI where a large BF% may be present (Gallagher et al., 2000, Romero-Corral et al., 2008a). In this case, large body fatness is indicated as greater than 30% (Oliveros, 2014, De Lorenzo et al., 2006). It is estimated that more than half of individuals are misclassified, and thus it is important to determine how we classify them into the correct category to accurately acknowledge their body fatness and consequently what, if any specific dietary recommendations we should be advising (Gallagher et al., 2000, Oliveros, 2014).

As a more accurate measure of body composition, the emerging gold standard method of air displacement plethysmography (ADP) can be used (Fields *et al.*, 2002, Collins and McCarthy, 2003). This method commercially known as the BODPOD calculates whole body volume and density which is then used to determine the percentage proportions of body fat and lean mass (Lowry and Tomiyama, 2015). It is a quick, safe, non-invasive method which provides highly reproducible results and is accommodating for all participants (e.g. children, adults, elderly, disabled, overweight, and obese) (Fields *et al.*, 2002, Lowry and Tomiyama, 2015, Ginde *et al.*, 2005). As a result, compliance and acceptability of the method is high due to minimal participant burden (Collins and McCarthy, 2003). The highly accurate and precise body composition results can be compared to the individuals BMI, which enables us to identify those who have a normal BMI with either a high or normal BF% (Collins and McCarthy, 2003, Fields *et al.*, 2002, Lowry and Tomiyama, 2015). Reviewing this in combination with both lifestyle and dietary factors allows the chronic disease risk to be assessed.

Body composition is influenced by lifestyle and dietary factors. The types of foods we habitually consume impact the ratio of fat mass to lean muscle mass in the body. Although many studies have investigated the effects of specific macronutrients on body composition, dietary patterns have had less attention (McNaughton *et al.*, 2007, McNaughton *et al.*, 2008, Hu, 2002). Assessing dietary patterns and diet quality addresses the diet as a whole and are two important determinants to consider.

Dietary patterns

The western dietary pattern has been associated with a higher BMI and an increased risk of cardiovascular disease, type 2 diabetes and cancer (Drewnowski, 2004, Schulze, 2006, Cha *et al.*, 2012, Rodríguez-Monforte *et al.*, 2015). It is characterised by a habitual intake of high glycaemic index (GI), refined grains, breads and cereals, red and processed meats, frequent consumption of fast-food, sugar-sweetened beverages (SSB), sweets and desserts (Schulze, 2006, Murtaugh *et al.*, 2007, Hu, 2002). New Zealanders adherence to this diet is increasing and as a consequence the prevalence of chronic disease is rising (Cha *et al.*, 2012).

Specific components of the western diet such as fast-food and SSB consumption are important factors of increased body fatness (Nicklas, 2001, Te Morenga *et al.*, 2013). The frequent consumption of SSB has consistently been associated with excess caloric consumption, increased body fatness and a diet lacking in essential nutrients (Nikpartow *et al.*, 2012, Schulze, 2004, DiMeglio, 2000, Johnson *et al.*, 2009, Tordoff and Alleva, 1990, Vartanian *et al.*, 2007). Individuals who consume primarily SSB have been found to have an 85% increased risk

of being overweight, and twice as likely to be overweight or obese compared to individuals who drink a combination of beverages (Nikpartow *et al.*, 2012). It is thought that the energy provided as a liquid carbohydrate is not compensated for throughout the day and this promotes a positive energy balance compared to consuming carbohydrate in a solid food form (DiMeglio, 2000, Mourao, 2007). Further to this, diet SSB have also been associated with an increased BMI due to increasing the desire to eat thus leading to a positive energy balance (Liebman *et al.*, 2006).

A positive energy balance can also be linked with fast-food consumption. Dietary patterns characterised by regular fast-food intakes have been shown to significantly increase the risk of obesity, especially in women (Liebman *et al.*, 2006, French, 2000, Schroeder *et al.*, 2007). Fast-food outlets are significantly increasing and are popular choices by many due to their convenience (Rosenheck, 2008). The food options available are typically energy-dense, nutrient poor, low in fibre, high in saturated fatty acids, trans fatty acids and salt, and served in excessive portion sizes (Rosenheck, 2008). Frequent consumption increases the energy and fat content of a diet and subsequently leads to increased body fatness and an increased BMI (Liebman *et al.*, 2006, French, 2000, Bowman and Vinyard, 2004). This is true, especially for women who have lower energy requirements compared to men and in conjunction with the low satiety factor of fast-food, the ability to overeat is enhanced (Liebman *et al.*, 2006). Although a strong relationship between obesity and fast-food consumption has been well documented (Prentice and Jebb, 2003, Rosenheck, 2008, Pereira *et al.*, 2005, Bowman and Vinyard, 2004), there is still controversy surrounding the causal relationship with some studies, questioning the role of physical inactivity (Rosenheck, 2008).

In comparison, a diet composed of fruit, vegetables, whole grain bread and cereals, lean unprocessed meats, poultry, fish and low fat dairy products is independently associated with a higher dietary adequacy, lower BMI, and lower waist circumference, thus reducing the risk of excess adiposity and consequent chronic disease (Schulze, 2006, Murtaugh *et al.*, 2007, He, 2004, Newby *et al.*, 2003, Liu *et al.*, 2003).

Diet quality

Diet quality can be referred to as dietary adequacy; the extent to which ones diet fits within the acceptable macronutrient distribution ranges (AMDR) and meets the recommended daily intake (RDI) for all micronutrients (Steyn, 2013). Dietary adequacy can be determined by an individual's dietary intake and eating patterns, which influence macronutrient distribution, body composition and health outcomes (figure 1.1) (Wirt and Collins, 2009). Women with poor

diet quality have been shown to consume diets lower in carbohydrates and micronutrients and higher in saturated fat and alcohol. These women are also at a higher risk of obesity (Wolongevicz, 2010).

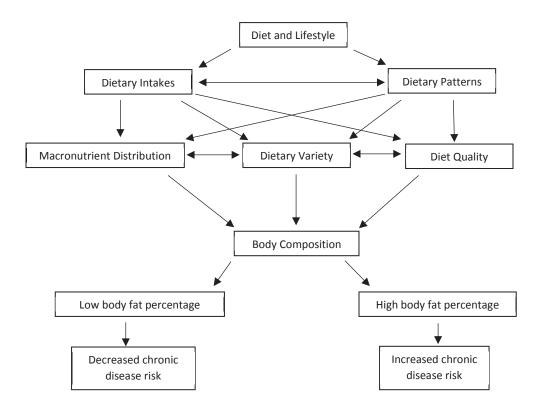


Figure 1.1 The relationship between diet, body fat percentage and health outcomes.

Studies investigating the macronutrient distribution ranges have found that when the BF% of women increased, the percentage of fat derived from their diet also increased (35%) and the carbohydrate content decreased (46%) compared to lean individuals (29% and 53%, respectively) (Miller *et al.*, 1990). Furthermore, compliance to dietary recommendations is low in obese individuals with less than 10% complying with the daily fruit and vegetable recommendations (De Abreu, 2013). This suggests that the dietary macronutrient composition may be an important determinant in dietary adequacy and resulting body composition.

Dietary intakes in obese individuals have routinely shown to be inadequate. Individuals with excess body fat consistently have a low quality diet, compared to those within the healthy BMI range who generally have high dietary adequacy (Wolongevicz, 2010). Micronutrients such as vitamin A, zinc, calcium and iron have previously been identified as low or deficient in individuals with excess body fat (Eckhardt, 2008, García *et al.*, 2009, Tidwell and Valliant,

2011). Study results have indicated that obesity may exacerbate dietary inadequacy or cause an adequate diet to become inadequate (Chai, 2010). When adjusted for nutrient density, obese individuals had a significantly lower serum level of 25-hydroxyvitamin D and carotenoids particularly pro-vitamin A compared to the normal weight women (Chai, 2010). This difference is thought to be due to the increased chronic inflammation seen in those with excess body fat which may lead to a higher requirement of some nutrients (Oliveros, 2014, De Lorenzo *et al.*, 2007). Low micronutrient status may contribute to the development of obesity and thus dietary recommendations different to that of individuals with healthy BF%'s may be required.

1.2 Purpose of the Study

Obesity is a multi-factorial worldwide health problem where its prevalence is increasing at an alarming rate. The dietary intake of individuals is a significant factor in determining their body composition and while obesity is important to consider, the dietary intakes and how they influence body fat content in women may be of value in assessing an individual's risk of chronic disease. It may be that those who have a normal BMI with high body fat require different dietary advice to those of normal BMI, normal body fat or those of high BMI, high body fat. While efforts have been made to address the role of dietary intake on BMI values, few attempts have investigated how the current dietary intakes and patterns of women relate to BF%, their subsequent BMI's and how we can tailor our dietary advice according to these body fat profiles. Investigating these factors will enable targeted interventions to be developed in an attempt to improve the morbidity and mortality outcomes in these women.

1.2.1 Aim

To explore the dietary intake and eating patterns of New Zealand European women with predetermined body composition profiles aged 16-45 years.

1.2.2 Objectives

- To explore the dietary intakes of women in the different body composition profile (BCP) groups.
- 2. To identify the eating patterns which are the strongest predictors of increased body fat.
- 3. To explore the differences in the macronutrient profile intakes in women with different body composition profiles.
- 4. To explore the micronutrient intake in each BCP group and compare the differences in diet quality.

1.3 Thesis Structure

This study had been structured into six sections. Chapter 1 introduces the research and highlights its importance. Chapter 2 is a review of the literature, including sections on obesity and health, dietary patterns, macronutrient distribution, diet quality, and assessing dietary intake. Chapter 3 outlines the methods and materials used in this study. This is then followed by chapter 4 which presents the results of the study. Chapter 5 discussed the findings of this study. Finally, chapter 6 draws conclusions and highlights the study strengths, limitations and recommendations for future research.

1.4 Researcher's Contributions to the Study

Researchers	Contributions to the thesis
Jenna Schrijvers	Main researcher, recruited and screened participants,
	supervised participant testing, data entry and analysis,
	conducted statistical analysis, interpretation and
	discussion of results, author of thesis
A/Prof Rozanne Kruger	Main academic supervisor, primary investigator of the
	EXPLORE study, application for ethics, development of
	study design, data analysis, and interpretation of
	results, thesis revision and approval
A/Prof Sarah McNaughton	Academic co-supervisor, advised and checked
	statistical analysis, interpretation of methods and
	results, reviewed thesis
Dr Kathryn Beck	Academic co-supervisor, advised on data analysis
Zara Houston, Chelsea Symons, Alex	Assistance with data entry of the NZWFFQ
Lawn	
Wendy O'Brien, Shakeela Jayasinghe	Coordination of participant recruitment, screening
	and testing
Maria Casale, Andrea Fenner, Zara	Recruitment and screening of participants
Houston, Adrianna Hepburn, Sarah	
Philipsen, Rozanne Kruger	
Pam Von Hurst, Cathryn Conlon,	Facilitated testing of the participants which included
Kathryn Beck, Richard Swift, Owen	eight stations: health screening questionnaire, blood
Mugridge, Maria Casale, Andrea	sample, blood pressure, BODPOD, DEXA scan, three
Fenner, Adrianna Hepburn, Sarah	dietary questionnaires (NZWFFQ, eating habits, eating
Philipsen, Rozanne Kruger	behaviour)
	(Note: participants were recruited as part of the wider
	EXPLORE study and not all data collected is used
	within this thesis)

NZWFFQ = New Zealand Women's Food Frequency Questionnaire;

DEXA scan = Dual Energy X-ray Absorptiometry

Chapter 2 Literature Review

2.1 Obesity and Health

2.1.1 Obesity definition

Overweight and obesity are defined as the presence of abnormal or excessive body fat which poses a health risk to the individual (World Health Organisation, 2014b). In general, a good indicator of obesity is the Body Mass Index (BMI), defined by using weight (kg) divided by height (m) squared (kg/m²). Individuals are classified into one of four categories: underweight (< 18.50 kg/m²), normal weight (18.50-24.99 kg/m²), overweight (25.00–29.99 kg/m²), and obese (\geq 30.00 kg/m²) (World Health Organisation, 2014b). A BMI greater than 25.0 kg/m² predisposes the individual to chronic disease, including diabetes, hypertension, dyslipidaemia, stroke and cardiovascular disease (World Health Organisation, 2014b). Overweight and obesity are the leading causes of death in the world with around 3.4 million adults dying each year due to complications associated with excessive body fat (World Health Organisation, 2014b). Globally in 2014, 1.4 billion adults (39%) over 18 years old were classified as overweight and of these, 600 million (13%) were obese (World Health Organisation, 2015).

2.1.2 Obesity prevalence in New Zealand

In New Zealand the 2012/13 National Health Survey identified 31.3% of adults over the age of 15 years as obese and a further 34% as overweight (Ministry of Health, 2014c). Specifically in New Zealand females, there has been a progressive increase in the prevalence of obesity (Ministry of Health, 2014c, University of Otago and Ministry of Health, 2011). An increase of both BMI and obesity prevalence was seen in the New Zealand European women and the total New Zealand population from 2006 to 2013/14. Although the percentage of overweight individuals decreased in both groups, this indicates an increase in the prevalence of significant excess adiposity in the New Zealand population (table 2.1).

Year	New Z	ealand Europea	an women	Total N	ew Zealand pop	oulation
	Mean BMI	Overweight	Obese	Mean BMI	Overweight	Obese
	(kg/m ²)	(%)	(%)	(kg/m ²)	(%)	(%)
2006/07	26.9	32.2	24.6	27.4	36.0	26.5
	(26.7-27.2)	(30.7-33.8)	(22.9-26.4)	(27.3-27.5)	(35.0-36.9)	(25.5-27.5)
2011/12	27.2	32.9	26.2	27.7	35.4	28.5
	(27.0-27.5)	(31.0-34.9)	(24.5-28.0)	(27.5-27.8)	(34.4-36.4)	(27.3-29.8)
2012/13	27.5	31.7	28.7	27.9	34.1	30.6
	(27.3-27.7)	(30.0-33.5)	(27.2-30.3)	(27.7-28.0)	(32.8-35.4)	(29.5-31.7)
2013/14	27.5	31.3	27.5	27.9	35.0	29.9
	(27.2-27.7)	(29.7-32.8)	(25.9-29.1)	(27.8-28.1)	(33.9-36.1)	(28.9-30.9)

Table 2.1 Mean BMI and prevalence of overweight and obesity in New Zealand Europeanfemales and the total population from 2006/07 to 2013/14 (Ministry of Health, 2014b).

BMI = Body mass index;

^a = Unadjusted mean BMI (95% confidence interval);

^b = Unadjusted prevalence percentage (95% confidence interval)

2.1.3 Causes of obesity

The aetiology of obesity is varied and complex. The alleged fundamental cause is energy imbalance, where the amount of energy consumed exceeds that which is expended over several months or years (Stein, 2004). Factors such as genetics, age, sex, hormones and other individualised features influence the risk of fat deposition, nevertheless the rapid increase in obesity prevalence suggests it is how the individual reacts to the environment which predicts excess adiposity (Caterson and Gill, 2002, Stein, 2004). The socio-ecological model is a theoretical framework which explains the complex interwoven relationship between environmental factors and obesity in developed countries (figure 2.1) (Blanchard, 2005). Intra and interpersonal factors such as a sedentary lifestyle, energy-dense nutrient-poor food choices (e.g. sugar-sweetened beverages (SSB) and fast-food), large portion sizes along with heavy marketing from food companies all play a part in the rising obesity levels (Swinburn, 2004).

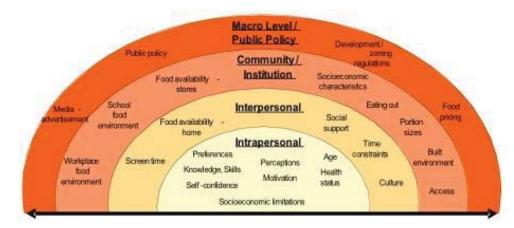


Figure 2.1 The Socio-Ecological Model of factors influencing increased adiposity (Fitzgerald, 2009).

Today, constant changes in the physical, socio-cultural, economic and political environments make it easy to constantly be in a positive energy balance (figure 2.1) (Caterson and Gill, 2002). Modernisation and economic restructuring such as changes in transportation, energy efficient machinery, and increasing urbanisation contribute to the increased availability, variety and consumption of energy-dense foods as well as the increasingly sedentary lifestyles of many individuals (World Health Organisation, 2014b, James *et al.*, 2004, Swinburn, 2004, Caterson and Gill, 2002, Stein, 2004). Advances in technology mean that machines have replaced what once were labour intensive jobs, leaving a sedentary desk job in its place. Domestic tasks such as grocery shopping have been replaced with the internet or smart phone applications, where food can be ordered and delivered straight to your door. This replaces many tasks contributing to our daily incidental physical activity levels and although many of these may indeed save us time, this time is not being put to good use and we are now more sedentary than ever (Caterson and Gill, 2002, Stein, 2004).

In parallel to this, advances in technology have also contributed to increased production of highly processed convenience foods designed to save us even more time. These products are often cheap, readily available, energy-dense and high in fat, salt and sugar; replacing the more traditional diet of what once was high in micronutrients, complex carbohydrates and fibre (Swinburn, 2004). These products in conjunction with cheap sugar sweetened beverages (SSB) and readily available fast-food largely contribute to the increase in obesity rates seen in New Zealand and around the world (Caterson and Gill, 2002).

Additionally, the marketing of these convenience foods is frequently targeted at young adults (Utter *et al.*, 2006). Typically during early adulthood, family influence on dietary habits is

reduced, where individuals start making their own food choices (Demory-Luce *et al.*, 2004). Young adults often move out of home to get further education, travel overseas or obtain additional life skills through flatting. During this time individual eating and lifestyle habits are created, therefore if healthy eating habits are not maintained or adopted, this is likely to contribute to increased adiposity in later adulthood (McNaughton *et al.*, 2008).

2.1.4 Obesity and chronic disease

In 2001, 60% of the 56.5 million deaths in the world were caused by chronic disease, with cardiovascular disease (CVD) contributing to almost half, and obesity and diabetes following close behind (World Health Organisation, 2003). Obesity is one of the most modifiable risk factors in the prevention of chronic disease, yet it is also one of the most common nutritional disorders in the developed world, and is becoming increasingly prevalent in developing countries alongside that of undernutrition (James *et al.*, 2004). The World Health Organisation (WHO) predicts that chronic disease will contribute to almost three quarters of all deaths worldwide by 2020 (World Health Organisation, 2003).

Vast amounts of literature have documented the increased risk associated with obesity, particularly abdominal obesity and the development of chronic diseases (figure 2.2) which include but are not limited to CVD, type 2 diabetes, hypertension, insulin resistance, dyslipidaemia, osteoarthritis, sleep apnoea, metabolic syndrome, and some cancers (University of Otago and Ministry of Health, 2011, Nicklas, 2001, Guh, 2009, Kannel *et al.*, 1996, Poirier *et al.*, 2006, Schenck-Gustafsson, 2009, Oliveros, 2014, McMichael, 2008).

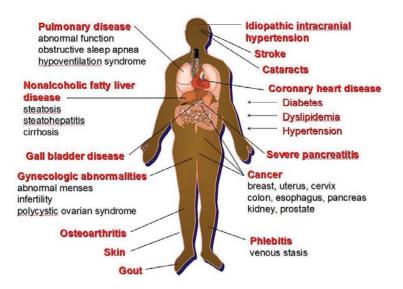


Figure 2.2 The medical complications associated with obesity (Mahan, 2012).

As adiposity increases, many individuals develop more than one of these diseases and thus are particularly susceptible to the metabolic syndrome (James *et al.*, 2004). An independent positive correlation between weight and coronary heart disease (CHD), stroke, congestive heart failure and CVD in women has been found, and those that are overweight by at least 30% are thought to have a stroke risk four times that of their leaner counterparts (Hubert, 1983, Kannel *et al.*, 1996). In addition, weight gain during adulthood is thought to increase the risk of CVD independent of initial weight or other risk factors that may be present (Hubert, 1983). Manson *et al* (1990) reported women to be two to five times more likely to develop hypertension, diabetes, hyperlipidaemia and CHD as adiposity increased with age. This was at least double the risk found in their leaner counterparts. Likewise, women who were moderately overweight or obese (BMI 25-30 kg/m²) had an 80% increased risk of developing CHD compared to their leaner counterparts (Manson, 1990). Conversely, weight loss has been found to improve or prevent chronic disease associated with excess body fat (Poirier *et al.*, 2006, Klein *et al.*, 2004). A five to 10% loss of body weight significantly improves CVD risk factors, with larger weight loss increasing these benefits (Wing *et al.*, 2011).

2.1.5 Assessing body fatness

Body mass index was first presented in the mid nineteenth century by Adolphus Quetelet based on the observation that body weight was proportional to the squared height in adults with normal body fat (Romero-Corral *et al.*, 2008b). It is the most widely used indicator to quickly identify an individual as overweight or obese, where excess body fat is assumed to be present (table 2.2) (Caterson and Gill, 2002).

Classif	ication	BMI (kg/m²)
Underv	weight	< 18.50
Norma	l weight	18.50 - 24.99
Overw	eight	25.00 - 29.99
Obese		≥ 30.00
-	Obese class I	30.00 - 34.99
-	Obese class II	35.00 - 39.99
-	Obese class III	≥ 40

Table 2.2 Body Mass Index (BMI) classification (World Health Organisation, 2014a, ObesityEducation Initiative Expert Panel on the Identification, 1998).

Although BMI is routinely used in many epidemiological studies and by health professionals, it is an imperfect measure of body fatness. The major assumption of the BMI guidelines is that at any given height, a higher weight correlates to a larger BF% and consequently a higher risk of morbidity and mortality (Flegal *et al.*, 2009). Some individuals who are overweight do not have excess adiposity, and in reverse those who have a normal BMI may have a high percentage of hidden body fat (Gallagher *et al.*, 2000). This is the main limitation of BMI; its inability to differentiate between lean muscle mass and fat mass (Oliveros, 2014). For example, an individual with a normal lean and healthy BF% and an individual who has a low lean mass with a high BF% may both have a BMI within the healthy range, however they have different susceptibility to disease risk. The latter individual may have a greater unidentified risk of morbidity and individuals such as these commonly present with metabolic dysregulation, thus BMI can be considered a crude measure of an individual's body fatness (Oliveros, 2014).

Body fat percentage is another way to identify obesity. Although less practical to determine, its accuracy allows us to better identify true excess adiposity, with a BF% greater than 30% correlating to obesity in women (table 2.3) (Oliveros, 2014, De Lorenzo *et al.*, 2006).

Body fat classification	BF% (women)	Explanation
High body fat (risky)	> 40%	Body fat needs to be lowered
Excess fat	31-40%	Excess body fat is present
Moderately lean	23 - 30%	Acceptable range for good health
Lean	19 – 22%	In range for excellent health and longevity
Ultra-lean	15 – 18%	Body fat levels found in elite athletes
Low body fat (risky)	< 15%	Body fat needs to be increased

Table 2.3 Body fat percentage (BF%) categories for women over 18 years of age (LifeMeasurement Inc, 2005).

Applies to women over 18 years of age

Commonly used methods to assess body composition include underwater weighing, skinfolds, dual x-ray absorptiometry and bio-electrical impedance. Their strengths and weaknesses are outlined in table 2.4.

Body composition assessment method	Strengths	Weaknesses	Assumptions
Underwater weighing	 Gold standard for determining body density and body fat Highly accurate 	 Impractical for a large sample size Subject must be willing to be submerged underwater Not suitable for children, the sick or elderly High degree of subject cooperation required Special equipment required Expensive 	• The fat-free compartment has a constant density
Skinfolds	 Equipment is inexpensive and portable Measurements are quick and easily obtained 	 A trained researcher is required to administer Strict adherence to standardised techniques are required Poor repeatability 	 Thickness of subcutaneous adipose tissue reflects a constant proportion of total body fat Skinfold sites selecte for measurement represent average thickness of the entire subcutaneous adipose tissue.
Dual x-ray absorptiometry (DEXA)	 Highly precise Fast and safe Low participant burden Requires little cooperation from patients Can be used for the old and sick Results highly correlated to underwater weighing 	 Cannot be performed on pregnant women due to radiation Differences in hydration and the presence of bone or calcified tissues may affect results 	• Hydration of fat free mass remains constant
Bio-electrical impedance (BIA)	 Safe Convenient Portable Fast Non-invasive 	 Expensive to purchase Dehydration can result in overestimation of fat mass Not as accurate as other methods 	 Assumes subjects are normally hydrated
Air displacement plethysmography	Highly accurateGood repeatability	 Subject must wear tight clothing and be 	• Fat free tissue has a constant density

Table 2.4. Strengths and weaknesses of the main body composition assessment methods (Lee,2010, Gibson, 2005)

Body composition assessment method	Strengths	Weaknesses	Assumptions
(ADP) (BODPOD)	 Quick and easy Can be used for children, adults, elderly and the disabled Can be used for large sample sizes 	comfortable in confined spaces • Not portable	

Romero-Corral *et al* (2008b) investigated the validity of BMI in relation to an individual's body composition and found that more than half of the participants with a BF% relating to obesity were not classified as overweight or obese using BMI as an indicator, while greater than 50% of individuals with excess body fat were classed as having a normal body weight (Romero-Corral *et al.*, 2008b). Confirmation of this is provided by Frankenfield *et al* (2001) who found that 46% of women identified as obese through bioelectrical impedance were not classified as obese using the BMI calculation. It has been suggested that using BMI with those who have a normal body weight is limited because it does not take into consideration differences in body fat and lean muscle mass (Romero-Corral *et al.*, 2008b). This indicates that the current obesity epidemic may in fact be larger than once thought, with many normal weight individuals carrying hidden body fat, increasing susceptibility to chronic disease.

2.2 Dietary Patterns

Nutritional epidemiology has traditionally focused on the effects of specific nutrients and foods on health. This approach requires researchers to distinguish the effects of one nutrient from another which is difficult due to complex nutrient combinations within foods (Schulze *et al.*, 2003). Although still an important part of research, large randomised controlled trials have been unsuccessful in showing the predicted health outcomes of individual nutrients (Jacques and Tucker, 2001). Recently, dietary patterns have emerged as a more comprehensive way to assess disease risk in relation to overall eating patterns (Mishra, 2006, McNaughton *et al.*, 2007). This approach recognises that nutrients and foods are consumed in combination with each other, rather than in isolation (Newby *et al.*, 2003, Hu, 2002, Hu *et al.*, 1999, Smith *et al.*, 2011, Schulze *et al.*, 2003, Mishra *et al.*, 2010). Foods have different nutrient compositions and the relationship between nutrients and different foods is an important determinant of disease risk. Particular combinations of foods are likely to be stronger predictors of health compared to considering only one nutrient in the diet (Michels, 2005). For example, a dietary pattern consisting of refined grains, high fat and high added sugar foods may elicit different health outcomes compared to a diet high in fruit, vegetables and whole grains. Assessing specific

dietary patterns may therefore provide a more comprehensive approach when identifying obesogenic trends and chronic disease risk because they focus on the diet as a whole and highlight these relationships and their resulting health outcomes (Jacques and Tucker, 2001, Fung *et al.*, 2001, Kerver *et al.*, 2003, Hu, 2002).

There are three main statistical methods of assessing dietary patterns commonly used in the literature. They include dietary indices, cluster analysis and factor analysis (Hu, 2002). Dietary indices assess overall diet quality using pre-determined criteria to assess diets, whereas cluster and factor analysis used the actual data to derive dietary patterns from. Cluster analysis aggregates individuals into groups based on commonalities between their eating patterns (Hu, 2002), while factor analysis measures the common variance between food items or food groups to identify combinations of foods that are regularly consumed together (Hu, 2002, Smith *et al.*, 2011). There are two main types of factor analysis: exploratory and confirmatory. Exploratory factor analysis investigates the underlying relationship between the constructs influencing a data set, whereas confirmatory factor analysis tests whether the constructs are related in the predicted way (DeCoster, 1998). The generic term 'factor analysis' encompasses both principal components analysis (PCA) and common factor analysis (Hu, 2002). Principal components analysis is considered to be a type of exploratory factor analysis, however it is commonly reported in the literature as being technically different (Hu, 2002, Field, 2009). Both are variable reduction techniques however factor analysis estimates factors from a derived mathematical model, whereas PCA identifies relationships between the linear components derived from the data set (Field, 2009). In pattern analysis, PCA is commonly referred to as 'factor analysis' and is the primary method used as it is psychometrically sound and is less complex than other factor analysis techniques (Field, 2009).

Literature has shown dietary patterns to be valuable in assessing diet-disease relationships. For this literature review, the most relevant studies using dietary pattern analysis and those that have been conducted recently involving women have been reviewed. Several examples of these studies are presented in table 2.5. Dietary patterns were used in the Dietary Approach to Stop Hypertension (DASH) study, where a dietary pattern was trialled in an attempt to reduce blood pressure in a population (Sacks *et al.*, 1999). The DASH diet significantly lowered participant's blood pressure after following the dietary pattern for 8 weeks, signifying the importance of dietary pattern changes rather than specific nutrients (Sacks *et al.*, 1999). The DASH diet success highlights the potential for this method to be expanded to other areas of health, such as obesity, CVD, type 2 diabetes and the resulting metabolic syndrome.

The western dietary pattern has been identified in literature as being a significant contributor to the increasing body fatness of the population. This pattern is characterised by a high intake of refined grains, red and processed meats, frequent consumption of fast-food, SSB, sweets and desserts which have been associated with a high BMI and an increased risk of chronic disease (Schulze, 2006, Nikpartow et al., 2012, Murtaugh et al., 2007, Murakami et al., 2013, Liu et al., 2003, Liebman et al., 2006). Literature shows the dietary components most strongly associated with weight gain were increased consumption of potato chips, potatoes (french fries, boiled, baked or mashed), SSB, red and processed meats, refined grains, sweets, desserts and alcohol (table 2.5). Strong positive correlations have also been found between weight gain and starches, refined grains and processed foods (Mozaffarian, 2011). This dietary pattern is common in obese individuals where these food items are likely to be in their top 10 favourite foods, with women preferring foods high in sugar and fat (Lahti-Koski et al., 2002). Consuming an empty calorie, energy dense diet (e.g. high in animal and vegetable fats, sweets, desserts, SSB and low in fruit, vegetables and fibre) has been shown by Quatromani et al (2002) to be associated with a 40% increased risk of obesity. These foods are energy dense and have a low satiety factor thus they are easy to over consume and contribute to excess energy intake (Mozaffarian, 2011).

Author(s) and location	Study design	Purpose	Population	Methods	Findings and conclusions
Suliga, 2015	Cross-	To identify the	2479 men	 Validated FFQ 	 Four dietary patterns were identified:
	sectional	dietary patterns of	and women,	 Principal component factor 	1. Healthy
	study	individuals with a	37-66 years;	analysis – food groups	2. Fat, meat and alcohol
		normal BMI and	BMI 18.5 –	 Height, weight, WC, BP 	3. Prudent
		which dietary	24.9 kg/m ² .	 Fasting blood samples (cholesterol, 	4. Coca-cola, hard cheese and french fries
Poland		patterns are		triglycerides, glucose levels)	 A high intake of fish and whole grains ('healthy'
		associated with the			pattern) and a low intake of refined grains, sugar and
		metabolic syndrome.			cold processed meat ('prudent' pattern) were
					associated with a lower risk of the metabolic normal
					weight obesity syndrome, higher HDL-cholesterol and
					better blood glucose control.
McNaughton <i>et al.</i> ,	Cross-	To investigate the	1086 girls	 108-item self-administered FFQ 	 Three dietary patterns were identified:
2008	sectional	association between	and boys, 12-	 Principal component factor 	1. Fruit, salad, cereals and fish pattern
	study	dietary patterns and	18 years.	analysis - food items	2. High fat and sugar
		socioeconomic		 24 hour food recall 	3. Vegetables
Australia		factors, nutrient		 Height, weight, WC, BP, BMI 	 Dietary patterns were not associated with
		intakes and health		(kg/m²)	socioeconomic factors, WC or BMI after adjusting for
		outcomes in			age, sex, and physical activity.

Table 2.5 Studies investigating dietary patterns and body fatness.

Author(s) and location	Study design	Purpose Po	Population	Methods	Findings and conclusions
		adolescents.			 Frequent intake of the 'fruit, salad, cereals and fish'
					pattern may be associated with diastolic BP in
					adolescent's ≥ 16 years.
McNaughton <i>et al.</i> ,	Longitudinal	To investigate the	569 men and	 Three 5-day food records at 36, 43 	 Four dietary patterns identified:
2007	study	relationship between	696 women,	and 53 years.	1. Ethnic foods and alcohol
		dietary patterns and	assessed at	 Exploratory factor analysis – 	2. Meat, potatoes and sweet foods
		the risk factors for	36, 43, and	food items	3. Fruit, vegetables and dairy
		chronic disease in	53 years.	 Height, weight, WC, BMI (kg/m²), 	4. Mixed
England, Scotland,		adulthood.		BP	ullet The 'fruit, vegetable and dairy' pattern was inversely
Wales				 Non-fasting blood samples (lipids, 	associated with BMI, WC and BP.
				HbA1c)	 A high consumption from the 'meat, potatoes and
					sweet foods' pattern may increase HbA1c.
					 Dietary patterns can predict chronic disease risk in
					adulthood.
Murtaugh <i>et al.</i> , 2007	Cross-	To investigate the	871 Hispanic	 Interviewed diet history 	 Five dietary patterns were identified:
	sectional	association between	and 1,599	questionnaire	1. Western
	study	dietary patterns, diet	non-Hispanic	 Factor analysis - food groups 	2. Native Hispanic
		composition, and	women, 25-	 Interviewed physical activity, 	3. Prudent
America		overweight and	64 years.	medical and reproductive	4. Mediterranean

Author(s) and location	Study design	Purpose P	Population	Methods	Findings and conclusions
		obesity.		questionnaire	5. Dieter
				 Weight, height, BMI (kg/m²) 	The 'western' dietary pattern (high fat dairy foods,
					refined grains, fast-food, processed meat, red meat,
					desserts) increases the risk of overweight and obesity.
					 Following a 'prudent' dietary pattern (low fat dairy,
					whole grains, legumes, vegetables, and nuts)
					decreases the risk of obesity by 50% in women.
Newby <i>et al.</i> , 2006	Longitudinal	To determine	33,840	 97-item, interviewed FFQ 	 Four dietary patterns were identified:
	Study	whether changes in	women, 49-	Principal components factor	1. Healthy
		dietary patterns	55 years.	analysis – food groups	2. Western/Swedish
		were associated with		• Weight, height, BMI (kg/m ²)	3. Alcohol
		changes in BMI in		 Data collected twice three years 	4. Sweets
Sweden		women.		apart	• The 'healthy' pattern was inversely associated with
					BMI.
					• An increase in the 'healthy' dietary pattern score was
					associated with a decrease in BMI in obese individuals.
					• Changes in eating patterns are significantly related to
					changes in BMI over several years.
Schulze, 2006	Prospective	To determine the	51,670	 133-item self-administered semi- 	 Two dietary patterns were identified:

Author(s) and location	Study design	Purpose	Population	Methods	Findings and conclusions
	cohort study	relationship	women, 26-46	quantitative FFQ	1. Prudent
		between dietary	years.	Principal component factor	2. Western
		patterns and		analysis – food groups	 Frequent consumption of fruit, vegetables, whole
America		weight changes in		 Demographic and physical activity 	grains, fish, poultry and salad dressing ('prudent'
		women.		questionnaire	pattern) is positively associated with a lower BMI
				 Height, weight, BMI (kg/m²) 	 Frequent consumption of red and processed meat,
				 Data collected at baseline, at year 	refined grains, sweets and desserts ('western' pattern)
				5 and year 8.	may lead to long term weight gain.
Newby, 2004	Cross-	To determine	459 healthy	 7 day food record 	 Six dietary patterns were identified:
	sectional	whether dietary	men and	 Exploratory factor analysis – 	1. Reduced fat dairy products, fruit and fibre
	study	patterns are	women, 22-	food groups	2. Protein and alcohol
		associated to	88 years.	 Demographic and physical activity 	3. Sweets
		anthropometric		questionnaire	4. Vegetable fats and vegetables
America		changes.		 Height, Weight, WC, BMI (kg/m²) 	5. Fatty meats
					6. Eggs, bread and soup
					 Frequent consumption of food items in the 'reduced fat
					dairy, fruit, and fibre' pattern leads to smaller gains in
					BMI and WC compared to other dietary patterns.
Newby <i>et al.,</i> 2003	Prospective	To investigate the	459 healthy*	 7 day food record 	 Five dietary patterns were found:
				ç	

Author(s) and location	Study design	Purpose Po	Population	Methods	Findings and conclusions
	study	dietary patterns	men and	 Cluster analysis – food 	1. Healthy
		leading to changes in	women, 30-	groups	2. White bread
		body mass index and	80 years.	 Height, weight, BMI (kg/m²) 	3. Alcohol
		waist circumference.		Data collected every 12-24 months	4. Sweets
				for 7 years.	5. Meat and potatoes
					 Fruit, vegetables, low fat dairy and whole grains
America					('healthy' pattern) are associated with smaller gains in
					BMI and WC compared to a high intake of red and
					processed meats, fast-food and SSB's ('meat and
					potatoes' pattern).
					 Analysing dietary patterns looks at the combination of
					foods and nutrients and is therefore a useful method to
					identify causes associated with obesity.
Quatromani <i>et al.</i> ,	Longitudinal	To determine the	737 women,	 145-item semi-quantitative FFQ 	 Five dietary patterns were identified:
2002	study	relationship between	30-98 years,	 Cluster analysis – food 	1. Heart healthy
		dietary intake and	BMI 18.5 -	groups	2. Light eating
		the development of	24.9 kg/m ² .	 Height, weight, BMI (kg/m²) 	3. Wine and moderate eating
		overweight (BMI ≥25		 Physical activity questionnaire 	4. High fat
		kg/m²).		 Data collected every 4 years for 	5. Empty calorie

Author(s) and location	Study design	Purpose P	Population	Methods	Findings and conclusions
				12 years.	 Women who consumed a diet reflecting the 'empty
					calorie' dietary pattern had a poor micronutrient intake
					and were at a higher risk of being overweight, compared
America					to those who consumed a 'heart healthy' dietary
					pattern.
					 Changes in dietary patterns and diet quality may
					influence weight management in women.
Maskarinec <i>et al.</i> ,	Cross-	To investigate the	514 women,	 209-item self-administered 	 Four dietary patterns were identified:
2000	sectional	relationship between	35-85 years.	validated FFQ	1. Meat
	study	dietary patterns and		 Principal components factor 	2. Vegetables
		body mass index in		analysis – food items	3. Bean
		women		 A one page questionnaire with 10 	4. Cold foods
Hawaii				soy food items	• The 'meat' dietary pattern was positively associated
				 Height, weight, BMI (kg/m²) 	with a higher BMI compared to the 'vegetable', 'bean'
					and 'cold foods' patterns.
					Assessing dietary patterns may be important in
					determining weight management.
*Healthy = free of chronic c GI = Glycaemic index; GL = Glycaemic load; WC = Waist circumference;	*Healthy = free of chronic disease at recruitment; GI = Glycaemic index; GL = Glycaemic load; WC = Waist circumference;		BP = Blood pressure; BMI = Body mass index (kg/m ²); HDL cholesterol = High density lipop. FFQ = Food frequency questionnaire	BP = Blood pressure; BMI = Body mass index (kg/m²); HDL cholesterol = High density lipoprotein cholesterol; FFQ = Food frequency questionnaire	

In comparison, fruit, vegetables, whole grains, nuts, seeds and yoghurt were associated with less weight gain, which is thought to be due to their nutrient dense, high fibre properties (Mozaffarian, 2011). For this literature review, the most relevant studies addressing different dietary characteristics have been reviewed. Several examples of these that have been recently conducted and included women in the analysis are presented in table 2.6. Individuals who consume a diet composed of the items above generally have a lower BMI and waist circumference with fewer weight related morbidities (Newby *et al.*, 2003, He, 2004, Murtaugh *et al.*, 2007). Women obtaining a favourable nutrient intake profile by following the nutritional guidelines and typically consuming a diet low in saturated fat, high in fruit, vegetables, whole grains and fibre are more likely to have a lower BMI and adequate nutrient intakes compared to those who routinely consume a poor nutrient dense diet (O'Neil, 2010, Quatromani *et al.*, 2002, Wirfalt, 1997).

Specific properties of these foods such as their nutrient density and high fibre content are thought to contribute to their effect on body mass (table 2.6). Fibre slows digestion and increases satiety, thus a larger consumption of these foods would displace other energy-dense foods thereby decreasing overall energy consumption (Mozaffarian, 2011). The fibre content of the foods also decreases the glycaemic index (GI) which is another important factor to consider. Murakami *et al* (2013) found that a large intake of high GI foods (e.g. processed breads and potatoes) and a low intake of lower GI foods (e.g. fruits, whole grain cereals and dairy products) was independently associated with an increased risk of obesity. This has also been shown in a study by Liu *et al* (2003) who also found a positive relationship between increased intake of refined grains and increased body weight in female participants.

Additional dietary patterns of milk and dairy food consumption and snacking behaviours have shown controversial results on body composition in the literature. Interestingly, some studies show no significant weight changes when dairy products or different types of milk are consumed (Mozaffarian, 2011, Gunther *et al.*, 2005). In contrast, a large meta-analysis concluded that dairy intake was associated with an overall healthier eating pattern and lifestyle, both of which are protective of chronic disease (Benatar *et al.*, 2013). Conversely, Lahti-Koski *et al* (2002) observed a positive relationship between whole fat milk consumption and obesity. Additionally, participants who preferred skim milk were more likely to become obese compared to those who did not drink milk at all. Reasons for this finding may be that obese subjects are giving socially acceptable answers or trying to offset their weight gain by choosing low fat items; unaware that they can be laden with sugar, which then may give them license to eat larger portions or more of other foods (Stein, 2004, Nielsen *et al.*, 2002). This

dietary habit may be true for either main meals or snacking throughout the day, as foods such as these are especially prevalent in items which are frequently consumed as snacks.

Snacking between meals can significantly contribute to excess energy intake, leading to weight gain. Basdevant *et al* (1993) found that those who snacked between meals generally consumed more energy from fat and less from protein throughout the day. Snacks tended to be items such as biscuits and pastries during the day, and cheese, beef and pork products as entrees before dinner. Additionally, food cravings were reported by 68% of the snackers and those that craved carbohydrate rich foods ate significantly more snacks per day compared to other snackers. Frequent snacking may lead to an increased dietary variety of undesirable food items. Dietary variety has been found to impact body fatness with a study finding increased food variety from entrees, sweets, snacks, condiments and high carbohydrate foods led to an increased energy intake and consequently an increased BF% (McCrory, 1999). Comparatively, they found that an increase in the variety of vegetable consumption (excluding potatoes) decreased the risk of obesity (McCrory, 1999).

Author(s)	Study design	Purpose	Population	Methods	Findings and conclusions
Murakami <i>et al.</i> ,	Cross-	To investigate the	678 men and 809	 7 day weighed food record 	A high GI and GL diet is independently associated
2013	sectional	relationship between	women, 19-65 years.	 Dietary GI and GL 	with an increased risk of general and central obesity.
	study	dietary GI and GL and		calculated	ullet Dietary GL had a positive relationship with general
		obesity.		 Height, weight, BMI (kg/m²), 	obesity in women.
United Kingdom				WC, W:H ratio	 Breads and potatoes were positive predictors of
				 7 day physical activity diary 	dietary GI. Fruit, cereals, and dairy products were
					negative predictors of dietary GI.
Mozaffarian,	Prospective	To investigate the	98, 320 healthy*	 Validated semi-quantitative 	Dietary and lifestyle factors were independently
2011	cohort study	relationship between	women and 22,557	FFQ	associated with long term weight gain.
		lifestyle changes and	healthy* men, 37-52	 Self-reported weight and 	 Frequent consumption of potato chips, french fries,
		long term weight gain.	years, BMI < 30.0	height	SSB, meat and processed meats, refined grains,
			kg/m²	 Medical history and lifestyle 	sweets and desserts were associated with weight
America				questionnaires	gain.
				 Data collected every 4 years 	 Weight gain was inversely associated with
				for 20 years.	consumption of fruit, vegetables, whole-grains, nuts
					and yoghurt.
Liu <i>et al.,</i> 2003	Prospective	To investigate the	74,091 healthy	61-item semi-quantitative	A higher intake of whole grains was associated with
	cohort studv	relationship between	females, 38-63 vears.	validated FFO's	less weight gain and a lower BMI.

Table 2.6 Studies investigating the relationships between dietary characteristics and body fatness.

Author(s)	Study design	Purpose	Population	Methods	Findings and conclusions
		refined grains and fibre		 Height, weight, BMI (kg/m²) 	Women with the highest whole grain intakes had a
		intake on long term		 12 year study 	49% lower risk of major weight gain.
America		weight gain.		 Dietary data collected every 3 	 Weight gain was negatively associated with high
				and anthropometric data	fibre, whole grain food consumption and positively
				collected every 2 years over	associated with refined grain consumption.
				12 years.	
* Hec	* Healthy = free of chronic disease at recruitment;	sease at recruitment;			

BMI = Body mass index (kg/m²); WC = Waist circumference; W:H ratio = Waist to hip ratio

2.2.1 Energy dense diets

Energy density can be defined as the amount of metabolisable energy per gram of food (Crowe *et al.*, 2004). The energy density of a particular food item is determined by its composition; with water and fat being the primary factors (Ledikwe *et al.*, 2005). Fat provides more than twice as much as energy per gram (9 kcal/g; 37.6 kJ/g) than protein and carbohydrate (4 kcal/g; 16.7 kJ/g), thus this coupled with the common low water and fibre content of high fat foods leads to a highly palatable and energy-dense item. Examples include foods such as potato chips, chocolate and doughnuts as they have a relatively low water content compared to fruit and vegetables (Drewnowski, 2004).

The energy density of a diet is closely correlated to its macronutrient distribution. An energydense diet is likely to obtain a higher percentage of energy coming from fat (Drewnowski, 2004). Several studies have shown a correlation between a high saturated fat diet (e.g. regular fast-food consumption) and a high BMI (Tremblay, 1989, Miller, 1990). It is thought that individuals consuming a large proportion of saturated fat are more likely to over-consume on any one day and thus may lead to unhealthy weight gain. Crowe *et al* (2004) found a positive relationship between energy density and the fat content of a diet. Passive overconsumption is likely when high fat foods are consumed due to their increased palatability, thus contributing to a positive energy balance. In addition to food, beverages, especially those sugar-sweetened have the potential to significantly increase the amount of energy consumed during the day. Food and fluids may have differential physiological effects on satiety and ad libitum food intake, thus it is important to consider the impact of fluids on energy intake separately (Swinburn, 2004, Mourao *et al.*, 2007).

2.2.2 Sugar sweetened beverages

Specific dietary patterns such as frequent consumption of SSB may be an important factor in increased body fatness (Nicklas, 2001). The term SSB encompasses all sweetened beverages such as soft drinks, fruit juice, cordials, sports drinks and iced teas. In the 2008/09 New Zealand Adult Nutrition Survey, 34.5% of the female population aged 15 years and over consumed fruit juice or drinks three or more times per week, and 34.7% of females consumed one or more soft drink per week (University of Otago and Ministry of Health, 2011).

The frequent consumption of SSB has been associated with excess caloric intake, increased body fatness and a diet lacking in essential nutrients (Nikpartow *et al.*, 2012, Schulze, 2004, DiMeglio, 2000, Johnson *et al.*, 2009, Van Wymelbeke *et al.*, 2004). A vast amount of literature has demonstrated a positive relationship between frequent SSB consumption and excess

energy intake leading to increased adiposity. Despite this, there is still much controversy surrounding the causal relationship and the role SSB's play in the development of overweight or obesity (Trumbo and Rivers, 2014). For this literature review, recent studies that included women and investigated the effects of SSB's on body composition were reviewed. Several examples of similar studies are presented in table 2.7 and merely showcase literature relevant to this thesis.

Author	Study design	Purpose	Population	Methods	Findings and conclusions	
Trumbo &	Systematic	To review the	59 Observational	 Literature search using key 	Inconsistent evidence was found for an association	ound for an association
Rivers., 2014	review	strength of	and 17	words related to obesity	between SSB intake and ob	between SSB intake and obesity risk when adjusted for
		evidence for the	intervention		energy balance.	
		relationship	studies in			
America		between SSB	children,			
		intake obesity	adolescents and			
		risk.	adults			
Malik <i>et al.,</i>	Systematic	To summarise	32 randomised	 Literature search (PubMed, 	The systematic review and I	The systematic review and meta-analysis both provided
2013	review and	the evidence on	controlled trials	EMBASE, Cochrane databases)	evidence that SSB consumption promotes weight gain	tion promotes weight gain
	meta-analysis	SSBs and body	(20 children, 12	 Prospective cohort studies and 	in children and adults.	
		weight in	adults)	randomised controlled trials		
America		children and		 Search conducted in March 2013 		
		adults.		 Random and fixed effect models 		
				used for the meta-analysis		
Nikpartow et	Cross-	To investigate	6,814 men and	 24 hour food record 	Seven clusters were identified:	ied:
<i>al.</i> , 2012	sectional	the relationship	7,463 women,	 Cluster analysis 	1. Soft drink	5. Beer
	study	between SSB	18-65 years.	 Height, weight, BMI (kg/m²) 	2. Fruit drink	6. Plain milk
		intake and BMI			3. Tea	7. Mix of beverages

Table 2.7 Studies investigating sugar-sweetened beverage (SSB) consumption.

Author	Study design	Purpose	Population	Methods	Findings and conclusions
					4. Coffee
					After adjusting for energy intake, a high consumption
					of fruit drink was a predictor of overweight and obesity
					in women.
Canada					Those whose primary beverage of choice was fruit
					juice (as a type of SSB) were 84% more likely to be
					overweight than those who drank a combination of
					beverages.
					A strong association was found between SSB
					consumption and overweight and obesity.
Chen, 2009	Prospective	To examine the	810 men and	 Telephone interviewed 24 hour 	Decreasing caloric intake from beverages was more
	study	relationship	women, 25 to 79	recalls	significantly associated with weight loss compared to a
		between	years	 Two recalls (one week day and 	decrease in calories from solid foods.
		beverage		one weekend day) were collected	A positive relationship was seen between the
		consumption		at baseline, 6 and 18 months.	reduction of SSB intake and weight loss at both 6 and
America		and weight		 Height, weight, BMI (kg/m²) 	18 months.
		changes in		obtained via weekly group	 Limiting calorie intakes from beverages, especially
		adults.		meetings.	SSB's aids in weight loss and preventing weight gain
				 Socio-demographic questionnaire 	

Author	Study design	Purpose	Population	Methods	Findings and conclusions
Liebman <i>et al.,</i>	Cross-	To investigate	883 men and	 Eating behaviour and general 	A high BMI was associated with frequent diet soft
2006	sectional	the association	1030 women, 18-	dietary questionnaires –	drink consumption.
	study	between BMI	96 years.	focussing on markers of a	 Individuals consuming diet soft drinks were more
		and dietary		healthy diet	likely to eat fast-food, order super-sized meals and
America		intake, eating		 Self-reported height and weight 	beverages, eat while doing other tasks and were less
		behaviours and		 Demographic questionnaire 	likely to participant in regular physical activity.
		lifestyle factors.			 Individuals who drank diet beverages were less likely
					to engage in other key lifestyle changes that help to
					facilitate a healthy body weight.
DellaValle <i>et</i>	Intervention	To determine	44 healthy	 Six test meals - ad libitum lunch 	Caloric beverages increased the energy intake at lunch
<i>al.</i> , 2005	study	whether specific	women, 18-60	with compulsory consumption	compared to non-caloric beverages.
		properties of	years.	of one of five beverages (water,	 Food intake was not decreased to compensate for the
		beverages		diet cola, regular cola, orange	increased energy provided from the caloric beverage.
		consumed with a		juice, and milk) or no beverage	A caloric or non-caloric beverage with lunch increased
		meal influence		as the control.	ratings of fullness compared to that of no beverage
America		dietary intake.		 Subjects rated their fullness, 	consumption.
				hunger, thirst, prospective	 Caloric beverages add a significant amount of energy
				consumption and nausea using	to a meal without affecting satiety ratings.
				100-mm visual analog scales.	

Author	Study design	Purpose	Population	Methods	Findings and conclusions
				 One meal per week for 6 	
				weeks.	
Schulze, 2004	Prospective	To investigate	51603 healthy	 133 item self-administered 	Frequent SSB consumption provides excess calories
	cohort	the relationship	women, 24-44	semi-quantitative FFQ.	leading to weight gain and an increased T2DM risk.
	analysis	between SSB's,	years.	 Biennial questionnaire (weight, 	Women who increased SSB consumption from 1 to 1 or
America		weight gain and		height, age, smoking,	more drinks per day gained more weight over a 4 year
		the risk of type 2		pregnancies).	period than those who consumed less.
		diabetes.		 Data collected every 2 	Women consuming 1 or more SSB per day were nearly
				years for 8 years.	twice as likely to develop T2DM.
Almiron-Roig &	Intervention	To compare the	14 men and 18	 Weight, height 	Inconclusive results to determine whether liquid and
Drewnowski,	study	effects of	women, 18-35	 Four beverages: orange juice, 	solid foods have differential effects on satiety.
2003		specific	years.	1% milk, cola, carbonated water	 Orange juice, milk and cola all increased fullness and
		beverages on		 Participants consumed a 	satiety and reduced hunger.
		hunger, thirst,		beverage with a standardized	Orange juice, milk and cola differed from sparkling
		satiety and		breakfast.	water but not from each other in their effects on
America		energy intakes		 Ratings of hunger, thirst, 	hunger and satiety ratings.
		at the next meal.		fullness and desire to eat were	Energy intakes at lunch did not compensate for the
				recorded.	beverage consumption at breakfast.
				 Food consumption at lunch was 	

Author	Study design	Purpose	Population	Met	Methods	Findings and conclusions
					measured.	
				•	One session per week for 4	
				-	weeks	
Raben <i>et al.</i> ,	Long-term	To investigate	6 healthy* men	•	Intervention groups: 1. F&D	Overweight individuals who consumed large amounts
2002	intervention	the long term	and 35 healthy*	- /	supplemented with sucrose 2.	of sucrose mainly as SSB had increased weight, BMI,
	study	effect of sucrose	women, 20-50	_	F&D supplemented artificial	body fat and BP.
		and artificial	years, BMI 25-30	- /	sweeteners.	 Ad libitum energy intake significantly increased in the
		sweetener	kg/m²	•	7-day food diaries completed at	sucrose group but stayed the same in the artificial
		supplementation		-	weeks 1, 5 and 10.	sweetener group.
		on ad libitum		•	A three-factor eating	• The weight gain in the sucrose group was thought to
Denmark		food intake and		-	questionnaire on dietary habits	be due to liquids being less satiating, inability to
		body weight		-	completed prior and post the	compensate for liquid calories, and the inability of
		regulation in			intervention period.	overweight individuals to regulate their energy intakes
		overweight		•	Weight, height, and body	efficiently.
		individuals.		-	composition (BIA and DXA	
				-,	scans) was measured every two	
				-	weeks.	
				•	24 hour urine samples collected	
				-	during the 6 th day and night	

Author	Study design	Purpose	Population	Methods	Findings and conclusions
				during weeks 0, 5 and 10.	
				10 week trial	
DiMeglio, 2000	Cross-over	To determine	7 males and 8	 Weight (kg) 	Liquid carbohydrate promotes a positive energy
	study	the differential	women, mean	 Liquid (soda) or solid 	balance.
		effects of liquid	age 22.8 years ±	(jellybeans) carbohydrate load	 Solid carbohydrates promote a balanced energy intake.
		and solid	2.73, mean BMI	consumed.	 Consuming solid carbohydrates resulted in a decrease
		carbohydrates	21.9 kg/m ² ± 2.2.	 24-hr phone dietary recalls: 3x 	in ad libitum food intake, whereas liquid carbohydrate
		on food intake		at baseline and 6x during	consumption did not.
America		and body weight		treatment and washout	 Body weight and BMI significantly increased over the 4
				periods.	week liquid carbohydrate groups.
				Hunger ratings measured twice	
				during intervention.	
				 2x 4 week periods separated by 	
				a 4 week washout period.	
De Castro, 1993	Observational	To investigate	124 male and	 7-day food diary 	Energy consumption is elastic and independent of any
	study	the effect of	199 females, 17-	 Rating of hunger and satiety 	single dietary component.
		consumption of	75 years, BMI	before and after eating,	• The consumption of caloric beverages, diet soda, tea
Atlanta		food and	17.7 - 38.1 kg/m²	respectively.	or coffee did not significantly change the amount of
		beverages on			energy consumed.

Author	Study design	Purpose	Population	Methods	Findings and conclusions
		nutrient intakes			When subtracting the energy of the caloric drink from
		throughout the			overall food intake, the energy from food was
		day			equivalent to days where the drink was not consumed.
Tordoff &	Cross-over	To investigate	21 males and 9	 Consumption of a 1135 g 	Frequent consumption of large volumes of SSB
Alleva, 1990	study	whether	females, mean	beverage sweetened with high	contributes to weight gain compared to the
		artificial	age 22.9 years ±	fructose corn syrup, aspartame	consumption of artificially sweetened beverages.
		sweeteners aid	0.8 for males and	or no drink each day for 3	The consumption of the SSB significantly increased
		in long-term	28.2 ± 2.7 for	weeks each.	participant weights by 0.66 kg compared to the
		weight	females.	 40-item eating attitudes 	aspartame beverage.
America		management		questionnaire	 Artificially sweetened beverages may aid in weight
		and food intake.		• 51-item restrained eating	control by reducing sugar and calorie intake.
				questionnaire	
				 Dietary record kept for 9 	
				weeks.	
				 Weight, height, BMI (kg/m²) 	
SSB = SL $BMI = B$ $BMI = B$ $FQ = FQ$ $*Healthy$ $T2DM =$ $F&D = Fc$ $BP = Bloc$	SSB = Sugar-Sweetened Beverage; BMI = Body Mass Index; FFQ = Food frequency questionnaire; *Healthy = Free of chronic disease at recruitment; T2DM = Type 2 diabetes mellitus; F&D = Food and Drink; BP = Blood pressure	age; naire; ase at recruitment; us;			

Women who increase their SSB consumption from one or less to one or more servings per day has been shown to have a larger weight gain over a four year period (Schulze, 2004). This correlates with findings from Nikpartow *et al* (2012) who found women that primarily choose fruit juice as a beverage are 84% more likely to be overweight, 2.55 times more likely to be obese and 2.05 times more likely to be either overweight or obese compared to those who drank a combination of beverages. Conversely, several studies have reported that unlike soft drinks, fruit juice has been associated with smaller increases in excess adiposity (Schulze, 2004, Mozaffarian, 2011). This is postulated to be due to the lower glycaemic index of fruit juice, the presence of phytochemicals, soluble fibre or the smaller quantities of fruit juice that is frequently consumed leading to lesser caloric intakes (Schulze, 2004).

The exact mechanism for the effects of SSB consumption on body weight is relatively unknown, however several theories have been postulated. Simple sugars such as those found in SSB elicit a hedonic effect by up regulating the expression of hunger signals such as neuropeptide Y and dopamine and decreasing satiety signals such as cholecystokinin and peptide YY, thereby increasing energy intake and leading to overconsumption (Erlanson-Albertsson, 2005). It has been proposed by Schulze et al (2004) that those who consume SSB have higher total energy intakes, specifically by 358 kcal/d (1,496.4 kJ/d) on average. In a cross-over study, Tordoff and Alleva (1990) compared the consumption of a soft drink sweetened with high fructose corn syrup (HFCS) to one sweetened with aspartame on body weight. When participants consumed the HFCS beverage they consumed 335 kcal/d (1,400.3 kJ/d) more and significantly increased their weight by 0.66 kg compared to when the aspartame beverage was consumed where energy intake and weight decreased nonsignificantly by 179 kcal/d (748.2 kJ/d) and 0.17 kg, respectively. Similarly, several studies have also shown energy intakes to be significantly increased in subjects who consume a SSB such as coca-cola, fanta, sprite or fruit juice compared to days when they did not or when a diet beverage was consumed (De Castro, 1993, Raben et al., 2002). This suggests that energy provided as a liquid carbohydrate is not compensated for by solid food intake during the day, thus promoting a positive energy balance (Schulze, 2004, DiMeglio, 2000, Mourao, 2007, De Castro, 1993). This is true, especially for overweight individuals who are less sensitive to dietary manipulation and cannot efficiently adjust their energy intakes accordingly (Raben et al., 2002).

The overconsumption seen with SSB intake may also be explained by its poor satiation effect. Both DellaValle *et al* (2005) and Almiron-Roig & Drewnowski (2003) found that consuming caloric beverages with a meal contributed to the higher overall energy intake without

significantly influencing overall satiety. The result was overconsumption, therefore SSB were deemed to be less satisfying than calories obtained from solid foods. Interestingly, one study found that this may also be true for diet SSB, where energy intake may also be higher due to the increased desire to eat during and following a meal (Liebman *et al.*, 2006). Various theories have been proposed in an attempt to explain the poor satiating properties of SSB. One involves the absence of mastication when they are consumed compared to solid foods. Chewing increases saliva and signals pancreatic and endocrine secretions some of which act to signal satiety such as cholecystokinin, peptide YY and ghrelin. Beverages are also cleared from the stomach faster than solid foods which may also decrease satiety and increase hunger sooner (Chen, 2009).

Additionally, large SSB intake tends to be accompanied by poor dietary and lifestyle habits as they are unlikely to be consumed in isolation to other foods. Large population studies have demonstrated that a high energy intake tends to be paralleled by a high total sugar intake, thus leading to excess caloric consumption, an increased risk of obesity and poor diet quality (Swinburn, 2004, Kvaavik *et al.*, 2005). Sugar- sweetened beverages tend to be consumed in large portion sizes complimented by high fat, sugar and salt laden meals from fast-food restaurants (Brown, 2008). A sedentary lifestyle, smoking and poor diet quality have also been linked to individuals who habitually consumed SSB and thus this group of individuals have a higher prevalence of morbidities such as obesity, type 2 diabetes and CVD (Schulze, 2004). Although SSB's have been shown to lead to increased adiposity by much literature (Malik *et al.*, 2013), the topic remains controversial with more recent literature questioning this relationship (Trumbo and Rivers, 2014, Pereira, 2014).

2.2.3 Fast-food consumption

Fast-food can be defined as food that is easily prepared and readily available from food outlets and restaurants or immediate consumption (Collins English Dictionary, 2015). The frequent consumption of fast-food has been shown to significantly increase the risk of obesity, especially in women (Bowman and Vinyard, 2004, French, 2000, Pereira *et al.*, 2005, Prentice and Jebb, 2003). Fast-food is a popular choice by many as it provides a fast and convenient meal option which is compatible with busy lifestyles (Rosenheck, 2008). Although the variety of fast-food outlets are becoming more diverse with many promoting healthier options, the industry leaders remain those selling items typically energy-dense, nutrient poor, and served in excessive portion sizes (Rosenheck, 2008, Swinburn, 2004). The frequent consumption of these energy-dense items such as burgers, deep fried chips and takeaway curries increases the energy content of the diet contributing to increased adiposity, especially abdominal obesity

which is known to increase the risk of chronic disease (Liebman *et al.*, 2006, Swinburn, 2004, French, 2000). This literature review explored studies that investigated the effect of fast food on body composition. Studies most relevant to this topic, those that included women and used a dietary analysis technique such as a FFQ, eating habits questionnaire or diet history were reviewed. Several examples of recent studies are presented in table 2.8.

Author	Study design	Purpose	Participants	Methods	Findings and conclusions
Wilcox et al.,	Cross-	To examine the	230 women, 25 -	3x telephone interviewed	Women who eat more fast-food have a higher
2013	sectional	association between	50 years, BMI >25	24 hr recalls within 15	energy intake as a result of consumption of energy
	study	fast-food consumption	kg/m2, waist	days of each other.	dense foods.
		and dietary intake and	circumference ≥88	 Alternative Healthy 	 Fast-food consumption was significantly associated
		quality in overweight	cm.	Eating Index calculated	with a higher energy intake, from meat, grains, SSB,
		and obese women.		for diet quality	dairy, added sugar, total, saturated and trans fats.
Columbia				 Weight, height, BMI 	 Fast-food consumption was negatively associated
				(kg/m²)	with the consumption of fruit, vegetables and whole
				 An ActiGraph 	grains.
				accelerometer was worn	The mean nutrient intakes showed a poor diet
				for 7 consecutive days.	quality score of 29.9, also increasing the risk of
					chronic disease.
Smith <i>et al.</i> ,	Cross-	To examine the socio-	1277 men and	127-item FFQ	Consuming fast-food was associated with a poor
2009	sectional	economic, lifestyle	1585 women, 26-	 Food habits 	diet quality in both men and women.
	study	factors and diet quality	36 years	questionnaire	 Women eating fast-food ≥2x per week were less
		associated with fast-		 Self-administered 	likely to meet the recommended intakes for fruit,
Australia		food consumption.		demographic and	vegetables, dairy, lean meat and legumes.
				lifestyle questionnaire	 Consuming fast-food ≥2x per week was associated

Table 2.8 Studies investigating the effects of fast-food consumption.

	Juuy ucaigii	Purpose	Population	Methods	Findings and conclusions
				 Weight, height, WC, BMI 	with a 25% higher prevalence of obesity in women.
				(kg/m ²)	
Schroeder <i>et</i>	Prospective	To identify the	1491 men and	165-item FFQ	There is a positive relationship between energy
al., 2007	study	relationship between	1563 women, 25-	 Mediterranean diet score 	intake, energy density and fast-food consumption.
		fast-food consumption,	74 years.	and healthy eating index	 An inverse relationship was seen with fast-food
		BMI, energy intake and		calculated for diet	intake and diet quality.
Spain		diet quality.		quality.	 An increased frequency of fast-food consumption
				 Weight, height and BMI 	increased BMI and the risk of obesity.
				(kg/m ²)	
Liebman <i>et al.</i> ,	Cross-	To determine the	883 men and 1030	 Socio-demographic 	BMI was positively associated with fast-food
2006	sectional	association between	women, 18-96	questionnaire (age,	consumption, large portion sizes and diet soft drink
	study	dietary intake,	years.	ethnicity, education,	consumption.
		behaviour and physical		employment status, PA)	
America		activity and a high BMI.		 Dietary intake and eating 	
				habits questionnaire	
				 Self-reported height 	
				 Self-reported weight 	
Pereira <i>et al.</i> ,	Prospective	To investigate the	3031 men and	 Qualitative FFQ 	Fast-food is independently and positively associated
2005	study	reported fast-food	women, 18-30	 Diet history of usual 	with weight gain and insulin resistance.

Author(s)	Study design	Purpose	Population	Methods	Findings and conclusions
		consumption habits and	years.	 dietary practices 	Frequent fast-food consumption was associated
		changes in body weight		Structured interview	with higher intakes of total energy, total fat,
		and insulin resistance		obtaining information on	saturated fatty acids, SSB, refined grains, meat and
		over a 15 year period.		dietary practices, food	low fibre intakes.
America				preparation methods and	 Frequent fast-food consumption was associated
				location of meals.	with lower intakes of fruit, non-starchy vegetables,
				 Standardised 	whole grains and reduced fat dairy products.
				demographic,	
				behavioural and PA	
				questionnaires.	
French, 2000	Prospective	To examine the	891 healthy*	 60-item validated FFQ 	Fast-food intake was associated with an increased
	study	relationship between	women, 20-45	TFEQ	energy intake, percentage energy from fat and a
		demographics,	years, mean BMI	 PA questionnaire 	higher body weight.
		behavioural and dietary	27.0 kg/m ²	 Weight, height, BMI 	 Consumption of vegetables per day was inversely
America		factors and frequency of		(kg/m²)	associated with fast-food consumption.
		fast-food consumption.		Data obtained once per	 An increase of one serving of fast-food per week
				year for 3 years	was associated with a 55 kcal increase per day and a
					weight gain of 1.68 kg over 3 months.
FFQ = Food frequency PA = Physical activity,	FFQ = Food frequency questionnaire; PA = Physical activity;	naire; WC: Waist circumference ; *Healthy = Free of chronic disease at recruitm	; c disease at recruitment;	TFEQ = Three factor eating questionnaire	nnaire

Liebman *et al* (2006) showed a positive association in women between frequent fast-food consumption and the likelihood of increased adiposity. Pereira *et al* (2005) also found an independent relationship between fast-food consumption and weight gain. In addition, a positive relationship between total energy, fat intake and fast-food consumption was shown by French *et al* (2000). They also found that increasing fast-food consumption by one serving per week led to an increase of 55 kcal per day, which correlated to a weight gain of 1.68 kg over three months (French, 2000). Likewise, consuming fast-food two or more times per week has been reported to lead to a 25% higher prevalence of obesity in women (Smith *et al.*, 2009).

The weight gain observed during frequent fast-food consumption is thought to be due to the lower energy requirements of women in conjunction with a low satiety factor (Liebman *et al.*, 2006). The inability for fast-food to satiate coupled by its high energy density properties means that it is easy to overeat (Liebman *et al.*, 2006). Sensory aspects of fat such as its 'mouth feel' make it pleasant and desirable to eat and thus is sought out in foods (Drewnowski, 1995). Many foods which have a high fat content are also high in energy and this coupled with low fibre and low satiety often leads to passive overconsumption, thus contributing to calories in excess of what is physiologically required (Blundell and Macdiarmid, 1997).

Eating in such a pattern means that the recommended daily intake of the food groups outlined by the Ministry of Health (e.g. fruit, vegetables, dairy, lean meat and legumes, breads and cereals) are unlikely to be met leading to a low nutrient dense diet (Smith *et al.*, 2009, Wilcox *et al.*, 2013). Individuals who frequently consume fast-food typically have a diet characterised by high total energy, saturated fat, salt, sugar, and refined grains, and this is thought to be due to these food items displacing healthier food options such as fruit, vegetables, whole grains and low fat dairy products (Schroeder *et al.*, 2007, Wilcox *et al.*, 2013, Pereira *et al.*, 2005, Paeratakul *et al.*, 2003).

2.3 Macronutrient Distribution

The ratio of carbohydrate, protein and fat which make up total energy intake may be a significant factor in excess adiposity. The Ministry of Health currently recommends 45-65%, 15-25%, and 20-35% of energy come from carbohydrate, protein and fat, respectively (National Health Medical Research Council, 2005). Several studies investigated the macronutrient ratio in regards to excess body fat with aligning results. Miller *et al* (1990) found that as the BF% of women increased, the percentage of fat derived from their diet also increased (35%) whereas the carbohydrate content decreased (46%) compared to lean individuals (29% of energy from

fat and 53% of energy from carbohydrate, respectively). In agreement with this Merchant *et al* (2009) found that women with lower BF%'s tended to consume diets higher in carbohydrates, where a diet composed of 47-64% of energy from carbohydrates was associated with a low risk of obesity. Tidwell & Valliant (2011) found that women with a BF% of at least 37.9% consumed on average 45.8% of energy intake as carbohydrates which was lower than those with less body fat. Fat intakes were shown to vary among women with different BF%'s, with those less and greater than 37.9% body fat having a mean intake of 21.6% \pm 6.0 and 32.6% \pm 6.2 of total energy intake, respectively.

2.3.1 Compliance with recommendations

The majority of NZE women meet the dietary macronutrient distribution range recommendations. On average, 16.4% of energy is obtained from protein, 33.6% of energy is from fat, and 47.1% of energy is from carbohydrates (University of Otago and Ministry of Health, 2011). The largest provider of protein was from the bread group (11%), butter and margarine provided the most energy from fat (9%) and bread was the largest contributor of carbohydrates (17%) (University of Otago and Ministry of Health, 2011). New Zealand European females consumed on average 73 g of protein, 69 g of total fat, and 211 g of carbohydrate per day. The Ministry of Health recommend no more than 15% of total energy come from sugar per day (National Health Medical Research Council, 2005). In the 2008/09 National Nutrition Survey, females consumed on average 101 g of total sugar per day, which equals 22.6% of total energy intake. Of this, 21.3% came from sugar and sweets and 15.2% came from non-alcoholic beverages (University of Otago and Ministry of Health, 2011). In addition, the Ministry of Health recommend no more than 10% of the energy allowance for fat should come from saturated and trans fats (National Health Medical Research Council, 2005). The National New Zealand Nutrition Survey found that on average, 13.1% of females' energy intakes were coming from saturated fats (University of Otago and Ministry of Health, 2011). These discrepancies of sugar and fat intakes with the recommendations imply that although the majority of New Zealand women meet the main macronutrient recommendations for protein, carbohydrate and total fat, the quality of carbohydrate and fat consumed is the issue driven by undesirable food sources.

Various studies have found that compliance to dietary recommendations is very low in obese individuals. De Abreu *et al* (2013) found that less than 10% of obese participants complied with the daily fruit and vegetable recommendations. Gonelevu *et al* (1997) investigated the fruit, vegetable and cereal intakes in New Zealand European women living in Auckland. They found that the obese women ate more broccoli and less carrot, apricots, kumara and boiled rice than

their non-obese counterparts; nevertheless, they concluded that neither group was meeting the '5+ a day' guideline for fruit and vegetable intake set out by the Ministry of Health (2013). Similarly, overweight and obese individuals have been shown to consume less than the recommended intakes of fruit, vegetables and dairy products and this may be due to the regular consumption of SSB, fast-food, sweets and processed foods which make it harder to conform to the dietary guidelines (Mozaffarian, 2011, Wilcox *et al.*, 2013). This is also demonstrated by Schroeder (2007) where an inverse relationship between fast-food consumption and meeting the daily dietary recommendations was seen.

2.4 Diet Quality

Diet quality is a term used to describe the nutrient adequacy of one's diet (Steyn, 2013, Streppel, 2014). It is the extent to which the diet meets all of the individuals daily energy and nutrient requirements (Steyn, 2013). To obtain high diet quality it is important to eat a variety of food items as no one food contains all of the nutrients needed for good health, thus low dietary variety leads to a poor diet quality (Steyn, 2013, Eckhardt, 2008). Poor diet quality has been associated with lower intakes of fibre, carbohydrates, protein and micronutrients and higher amounts of saturated fat and alcohol (Wolongevicz, 2010). When a diet with low variety is consumed, it tends to be based on starchy foods rather than fruit, vegetables and lean animal products (Steyn, 2013).

Individuals who have poor diet quality tend to have a higher risk of obesity, thus obese individuals appear to be at greater risk of suboptimal micronutrient levels such as vitamin A, vitamin D, zinc, calcium and iron (Wolongevicz, 2010, Eckhardt, 2008, García *et al.*, 2009). Consuming a poor quality diet may exacerbate the risk of chronic disease already experienced by obese and overweight individuals due to excessive adiposity (Ruxton, 2011). Several studies have investigated the relationship between diet quality and obesity. For this literature review, studies most relevant to this topic, specifically those investigating diet quality and body composition in women were reviewed. Several examples of these are presented in table 2.9. Wolongevicz *et al* (2010) found women to have a two-fold increased risk of obesity when consuming a poor quality diet. Women with the lowest diet quality consumed diets low in energy, carbohydrate and micronutrients and higher in total fat, saturated fat and alcohol. These women also gained 2.3-3.6 kg over eight years compared to those consuming a higher quality diet (1.4 kg weight gain). In addition, Sundararajan *et al* (2014) also found an inverse relationship between diet quality and BMI which is strongest among women.

Author	Study design	Purpose	Participants	Methods	Findings and conclusions
Sundararajan <i>et</i>	Cross-sectional	To assess the	6325 men and	24-hour interviewed dietary recall	DQ was associated with a lower BMI in
<i>al.</i> ,2014	study	relationship between	7211 women, 18-	 DQ: Healthy eating index 	obese individuals.
		diet quality and BMI.	65 years	 Weight, height, BMI (kg/m²) 	The relationship between DQ and BMI
Canada					was strongly related in women.
Wolongevicz et	Longitudinal	To determine the	1265 healthy*	3-day food record	Women with poor diet quality were
<i>al.</i> ,2010	study	association between	women, 28-62	DQ: Dietary risk index score	twice as likely to be overweight or
		diet quality and	years	 Weight, height, BMI (kg/m²), WC 	obese.
		abdominal obesity		 Fasting blood samples (blood lipids 	Women with the lowest DQ had low
America		risk in women.		and HbA1c)	intakes of fibre, calcium, protein,
				• BP	carbohydrates and higher consumption
				 Data obtained once every 4 years 	of energy, alcohol, and total, saturated,
				for 12 years.	polyunsaturated and monounsaturated
					fats.
Wolongevicz,	Prospective	To investigate the	590 women, 25-	3-day dietary records per week	Women with lower DQ were more likely
2010	study	relationship between	71 years, BMI <25	(two weekdays, one weekend day)	to become overweight or obese.
		diet quality and the	kg/m²	 DQ: Framingham nutritional risk 	Women with low DQ had lower energy,
		development of		score	carbohydrate, fibre and all
		overweight/obesity.		 Food habits questionnaire 	micronutrient intakes and higher

Table 2.9 Studies investigating the relationship between diet quality and body composition.

Author	Study design	Purpose	Participants	Methods	Findings and conclusions
				 Weight, height, BMI (kg/m²), WC 	alcohol, total, saturated and
				 Fasting blood samples (lipids and 	monounsaturated fat intakes.
America				HbA1c)	Weight gain was on average 1.4 kg for
				• BP	those with higher DQ, compared to 2.3-
				Data obtained once every 4 years	3.6 kg with low diet quality, over 8
				for 16 years.	years.
McNaughton <i>et</i>	Cross-sectional	To identify the	7441 men and	74-item FFQ developed for use in	A higher DQ was associated with lower
<i>al.</i> ,2009	study	relationship between	women, aged ≥	Australian adults	systolic BP and fasting blood glucose
		diet quality and the	25 years old	DQ: Dietary guideline index	levels and higher insulin sensitivity in
		presence of pre-		 Weight, height, BMI (kg/m²), WC, 	women.
Australia		diabetes, newly		• BP	A high DQ was inversely associated with
5		diagnosed diabetes		 75 g oral glucose tolerance test 	the prevalence of pre diabetes among
		and cardiovascular		 Fasting blood samples (lipids and 	women.
		risk factors.		HbA1c)	 High DQ was characterised by higher
					wholegrain, fruit, vegetables, lean
					protein, and reduced fat dairy products.
DQ = Diet quality; BMI = Body mass index;		BP = Blood pressure; WC = Waist circumference			

Poor diet quality may help to explain the consistent finding of low or deficient levels of several micronutrients in obese individuals. The reasons behind these deficient levels is relatively unknown, however several theories have been postulated. Firstly, obese individuals tend to have adequate intakes of energy, thus their intakes of micronutrient rich food may be low due to poor food choices, or they may have increased requirements due to changes in the absorption, excretion or metabolism of various nutrients (Ruxton, 2011). Secondly, the primary principle in weight loss strategies is to restrict energy intake. As micronutrient intakes are already low in these individuals, further restriction of food intake without increasing the consumption of fruit, vegetables and whole grains may further exacerbate the low micronutrient levels (Ruxton, 2011). Thirdly, studies suggest that individuals who follow a low fat diet (which is thought to be healthy) do not necessarily have a nutrient rich diet (Wirfalt, 1997). In one study, two clusters (soft drink and skim milk clusters) were identified both of which had low fat intakes. The micronutrient intakes between each group however differed significantly, with the skim milk cluster having a higher micronutrient intake compared to the soft drink cluster. The soft drink cluster's micronutrient intake was consistent with that seen in individuals who consume a relatively high fat diet (e.g. pastries, meat and white bread clusters) (Wirfalt, 1997).

It is thought that the diet quality of these individuals can be increased by incorporating more fruits, vegetables and whole grains into their diet, while eliminating the consumption of fastfoods and SSB (Bowman and Spence, 2002). When whole grain consumption is increased, micronutrient levels improve with the exception of vitamin B12 and sodium (O'Neil, 2010). The concentrations of both vitamin A and C also increase regardless of whole grains being a poor source of these nutrients. This suggests that individuals who increase their whole grain consumption also increase the quality of other parts of their diet (e.g. fruit and vegetable intake), thus overall consuming a healthier diet (O'Neil, 2010).

2.4.1 Micronutrient status

Excess adiposity disrupts the normal metabolic actions of the body, increasing the risk of hypertension, insulin resistance, diabetes, dyslipidaemia and CVD (Wisse, 2004). The risk depends on the extent to which excess adipose tissue is present, regardless of the individuals BMI, among other factors such as genetic disposition, ethnicity, age and lifestyle (Wisse, 2004). As BF% increases, so too does oxidative stress in the body caused by the secretion of various proteins, hormones and other factors by the excess adipocytes. This oxidative stress leads to tissue damage putting the body in a chronic low-grade systemic inflammatory state due to the

innate immune response, which is thought to contribute to the pathogenesis of insulin resistance (Wisse, 2004, Capurso and Capurso, 2012, Cazettes *et al.*, 2011, Garcia-Bailo, 2011).

Actions of modifiers such as micronutrients on inflammation may assist in reducing the damage of oxidative stress on the body (Garcia-Bailo, 2011). In reverse, inflammation may play a crucial role in determining the micronutrient levels in the blood stream (Chai, 2010). Individuals with excess adiposity have been shown to have low micronutrient levels regardless of their diet quality (Chai, 2010). Although a relatively new area of research, the available evidence routinely shows that overweight and obese individuals who have a higher percentage of fat mass compared to lean mass are more likely to have low levels of various micronutrients compared to that of normal weight women (tables 9, 10, 11, 12). This lower level of micronutrient status is also thought to be independent of dietary intake, even after adjusting for nutrient density of the diet, thus obesity may have a direct influence on micronutrient status in these individuals (Chai, 2010). Various mechanisms have been suggested to explain the low levels of micronutrients in overweight and obese women, and one popular theory is that the cutaneous and dietary fat soluble vitamins become sequestrated by the adipose tissue decreasing its availability for absorption (Martini and Wood, 2006). Another theory states it is due to early stage development of obesity related diseases such as diabetes and CVD, where the body requires more of these nutrients (Ruxton, 2011, Kimmons, 2006).

2.4.2 Vitamin A

Controversy surrounds the correlation between vitamin A levels and BF%. Literature surrounding the relationship between vitamin A status and BF% was explored. Recent studies most relevant to this topic and those that included women between the ages of 16 and 45 years were reviewed. Several examples of these studies are shown in table 2.10. An inverse relationship between vitamin A levels and obesity in women has been found in some studies. Obese premenopausal women to have significantly lower vitamin A levels compared to their normal weight counterparts (table 2.10) (Chai, 2010).

Garcia, 2012b Cr sti Mexico	Cross-sectional)
		To determine the	580 women, 37 ±	 24 hr recall (1x week day, 	Vitamin A levels were associated with women who had
Mexico	study	association of zinc,	7.5 years	1x weekend day)	higher leptin concentrations.
Mexico		vitamin A, C and E		 Weight, height, WC, HC 	High leptin levels were associated with a high body fat
Mexico		concentrations		 Body composition (DXA) 	percentage.
)		with BMI, central		 Fasting blood samples 	No significant difference in vitamin A levels between
		adiposity, body fat		(glucose, lipids, leptin, zinc,	overweight/obese and normal weight individuals.
		and leptin levels.		vitamin A, C and E)	
Chai, 2010a Cr	Cross-sectional	To determine the	180 women, 35-46	 Validated FFQ for dietary 	The metabolism of pro-vitamin A may contribute to the
sti	study	association	years. Stratified by	assessment.	differences seen in different body compositions.
		between serum	BMI: normal (18.5-	 Height, weight, BMI 	Both pro and non-pro vitamin A carotenoid serum levels
		soluble	24.9 kg/m²);	(kg/m ²)	were significantly lower in the obese BMI subgroup
		micronutrients and	overweight (25-	 Fasting blood samples 	compared to the normal BMI group.
		BMI.	29.9 kg/m ²); obese	(pro-vitamin A (α-	Pro-vitamin A, lutein and total energy intakes were not
Hawaii			(>30 kg/m ²)	carotene, β-carotene, β-	significantly different between BMI groups.
				cryptoxanthin), non-pro	
				vitamin A (lycopene,	
				lutein/zeaxanthin, α-	
				cryptoxanthin,	

Table 2.10 Studies investigating the relationship between vitamin A status and body composition.

Author(s)	Study design	Purpose	Population	Methods	Findings and conclusions	conclusions
				anhydrolutein), tocopherols		
				(α-tocopherol, γ-tocopherol).		
Viroonudomp	Cross-sectional	To examine the	Intervention group:	 Weight, height, BMI 	 Low concentratior 	Low concentrations of antioxidant vitamins may increase
hol <i>et al.</i> , 2003	study	relationship	16 males and 56	(kg/m ²), WC, HC, W:H	the risk of coronary artery disease.	ry artery disease.
		between	females, BMI ≥	ratio, MAMC	 Weight, BMI, HC, i 	Weight, BMI, HC, and MAMC were negatively correlated
		anthropometric	25.0 kg/m², 16-60	Fasted blood sample	with serum retino	with serum retinol and serum $lpha$ -tocopherol in both
		measurements,	years.	(serum retinol, α-	overweight and obese females.	bese females.
Thailand		antioxidant status,	Control group: 14	tocopherol, lipid profiles)	 Lower serum retir. 	Lower serum retinol levels were seen in overweight (1.8%)
		and lipid profiles.	males and 58		and obese (10.7%,	and obese (10.7%) individuals, although not significant.
			females, BMI 18.5-			
			24.9 kg/m², 16-60			
			years.			
FFQ = F BMI = L WC = N	FFQ = Food frequency questionnaire; BMI = Body mass index; WC = Waist circumference;		DXA = Dual-energy x-ray absorptiometry; HC = Hip circumference; W:H ratio = Waist to hip ratio;	netry; MAMC = Mid arm muscle circumference; LDL-c = Low density lipoprotein cholesterol; HDL -c= High density lipoprotein cholesterol;	ircumference; itein cholesterol; otein cholesterol;	TC = Total cholesterol; TGs = Triglycerides

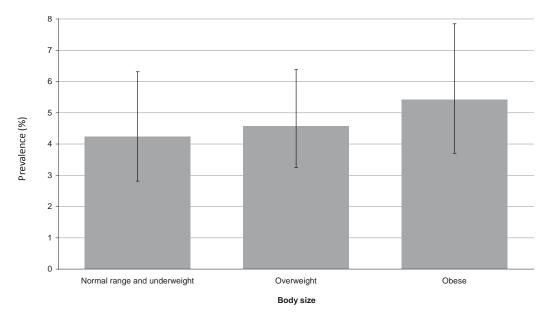
In contrast Garcia *et al* (2012) found that all overweight, obese and normal weight women had adequate vitamin A levels, however the levels in normal weight women being higher than those with excess BF. Viroonudomphol *et al* (2003) also found no significant differences in serum retinol levels between overweight and normal weight subjects. A mere 1.8% of overweight and 10.7% of obese women were found to have low retinol levels, however this was higher than those with lower BF%'s. Seventy-five percent of the studies in table 10 used serum retinol as the measure of vitamin A status. Serum retinol is the measure of adequate vitamin A levels which in normal individuals is tightly regulated by the liver within a narrow window, and it is only when the vitamin A stores start to become depleted that these levels also drop (Viroonudomphol *et al.*, 2003). It is possible that the difference observed in these studies is due to the participants being at different stages of vitamin A liver store depletion at the time of the study.

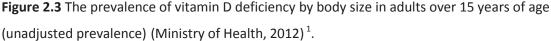
The effect of obesity on micronutrient status may be independent of dietary intake. After adjustment for nutrient density, Chai et al (2010) found the obese subgroup had on average a 45% lower pro-vitamin A concentration (α -carotene, β -carotene and β -cryptoxanthin), whereas non-pro vitamin A (lycopene, lutein/zeaxanthin α -crytoxanthin and anhydrolutein) was only reduced by 15% in the obese women. Due to no differences seen in the serum carotenoid levels after adjustment for nutrient density, it is unlikely that this difference is due to dietary factors. It was suggested that this is due to the increased chronic inflammation which may lead to a higher requirement of some nutrients in this subgroup of the population. Vitamin A has antioxidant properties, and also works with the immune system to prevent infections and cell damage (Kimmons, 2006). Low carotenoid levels have been associated with increases in oxidative stress, insulin resistance, impaired glucose metabolism, some cancers, and age-related macular degeneration (Kimmons, 2006). A high adiposity leads to a high inflammation level in the body and thus more oxidative stress which may lead to a faster metabolism of pro-vitamin A and increased oxidation of carotenoids causing the smaller decrease in non-pro-vitamin A (Chai, 2010). Larger requirements of total vitamin A would therefore be required to counteract and manage the detrimental effects this has on body tissues (Chai, 2010).

2.4.3 Vitamin D

Vitamin D is one of few micronutrients which have been extensively studied in relation to obesity. An inverse relationship has been shown to exist between increased adiposity and vitamin D levels (Chai, 2010, Blum *et al.*, 2008, Samuel and Borrell, 2013, Lagunova, 2009, Ardawi *et al.*, 2011). The New Zealand 2008/09 Adult Nutrition Survey showed an increase in

the prevalence of vitamin D deficiency as excess adiposity increased in individuals over 15 years of age (figure 2.3). The prevalence of vitamin D deficiency was 4.2 percent in individuals in the normal range and underweight group, and 5.4 percent in those who were obese. This difference was no longer significant however once adjusted for age, sex and ethnicity.





This positive relationship is also seen in females where low intakes of vitamin D and calcium have been associated with obesity (Moy and Bulgiba, 2011). Blum et al (2008) found that obese women had a significantly lower 25-hydroxyvitamin D level compared to normal weight individuals. This is backed up by Lagunova *et al* (2009) who found a negative association between BMI and both 1,25(OH)₂D₃ and 25(OH)D₃ concentrations with 32% (1 in 3) women with a BMI greater or equal to 40 kg/m² to have a vitamin D deficiency. Literature surrounding the relationship between vitamin D status and BF% was explored. Studies most relevant to this topic and those that included women between 16 and 45 years old were reviewed. Few studies investigating this topic were relevant to this thesis and those that were are presented in table 2.11. It was concluded that BMI has a negative impact on vitamin D deficiency,

¹ Vitamin D deficiency was defined as a serum 25-OHD level less than 25 nmol/L. Underweight was defined as a body mass index (BMI) of less than 18.5 kg/m², normal body size was defined as a BMI of 18.5-24.9 kg/m², overweight was defined as a BMI of 25.0-29.9 kg/m², obesity was defined as a BMI score of greater or equal to 30 kg/m².

 $25(OH)D_3$ level and seasonal variance. Further confirmation of this is provided by Samuel & Borrell (2013) who found that overweight and obese individuals were 24% and 55% more likely to have low vitamin D levels compared to their normal weight counterparts, respectively.

Author	Study design	Purpose	Participants	Methods	Findings and conclusions
Ardawi <i>et al.,</i>	Epidemiological	To determine	1172 healthy*	 Standardized socio- 	Vitamin D deficiency was attributed to the presence of
2011	study	factors influencing	women, 20-79 years	demographic	obesity, poor sunlight exposure and poor dietary
		vitamin D status in	(501 pre-	questionnaire (lifestyle,	supplementation.
		pre and post-	menopausal, 671	smoking habits, PA, tea	 Obesity was the main risk factor for vitamin D deficiency.
		menopausal	post-menopausal),	and coffee consumption,	• BMI was the best predictor of vitamin D deficiency.
		women.	mean BMI 30.6 ± 6.6	vitamins and medication)	38.3% of premenopausal women had moderate to severe
Saudi Arabia			kg/m²	 Weight, height, BMI 	vitamin D deficiency (serum 25(OH)D < 25 nmol/L).
				(kg/m ²), W:H ratio	A negative relationship between vitamin D status and BMI
				 Fasting blood samples 	was seen in pre-menopausal women.
				(serum 25(OH)D, calcitrol,	Only 16.4% of premenopausal women had vitamin D levels
				Ca, PO_4 , Mg, creatinine	≥ 75.0 nmol/L.
				and albumin).	
				 Sun exposure index 	
				calculated.	
				 BMD measured using DXA 	
Tidwell and	Cross-sectional	To determine the	100 African	 24-hour dietary recall 	Women with lower calcium and vitamin D intakes are
Valliant, 2011	study	relationship	American women,	 Body composition (DXA) 	likely to have excessive adiposity compared to those with
		between excessive	18-40 years, Mean		a higher intake.

Table 2.11 Studies investigating the relationship between vitamin D status and body composition.

	Study design	Purpose	Population	Methods	Findings and conclusions
		body fat and levels	BMI 29.8 kg/m ²		• Women with \ge 37.9% body fat had calcium and vitamin D
		of vitamin D and			intakes of 528.6 mg/d and 3.8 µg/d, respectively.
America		calcium			• Women with \leq 37% body fat had calcium and vitamin D
					intakes of 911.5 mg/d and 5.0 µg/d, respectively.
					An inverse relationship was seen between calcium, vitamin
					D and BF%.
Lagunova, Epidemiological	iological	To investigate the	1737 women, < 50	 Weight, height, BMI 	• The prevalence of vitamin D deficiency, the 25(OH)D ₃ level
2009 study		seasonal variation	years , mean BMI	(kg/m 2), body fat mass.	and seasonal variation are all dependent on BMI.
		and the prevalence	31.8 ±6.67 kg/m²;	 Blood samples (serum 	 The prevalence of vitamin D deficiency was significantly
		of vitamin D	805 women, >50	1,25(OH) ₂ D ₃ and 25(OH)D ₃	higher in women with a BMI \ge 40 kg/m ² .
		deficiency in a	years, mean BMI	levels).	
Norway		Norwegian	31.8±6.28 kg/m²;		
		population.	205 men, < 50 years,		
			mean BMI 33±6.91		
			kg/m2; 184 men, >		
			50 years, mean BMI		
			32.3± 5.92 kg/m².		

BMI = Body mass index; BMD: Bone mineral density; DXA = Dual-energy x-ray absorptiometry

Several mechanisms are thought to be behind this lower vitamin D level. Firstly, the amount of vitamin D obtained through dietary sources determines the serum concentration in the body and also reflects diet quality. Tidwell & Valliant (2011) have shown that African-American women with a BF% less than or equal to 37.9% had a higher dietary intake of vitamin D and calcium as well as lower intakes of carbohydrates, energy and fat compared to those with higher BF%'s. The dietary intakes in those women with higher adiposity levels revealed that they consumed more SSB, fruit juice and non-fat smoothies. This suggests that a higher diet quality is associated with higher levels of these nutrients (Tidwell and Valliant, 2011).

Secondly, vitamin D is thought to be sequestrated by adipose tissue, thus decreasing the bioavailability of cutaneous and dietary vitamin D sources (Wortsman *et al.*, 2000, Chai, 2010). A study by Blum *et al* (2008) found a positive correlation between vitamin D and storage in adipose tissue. This correlation is thought to be due to the enhanced uptake and reduced availability of vitamin D once deposited in body fat compartments.

Thirdly, it is thought that obese individuals may not be exposed to sunlight as often as their normal weight counterparts and thus may lead to lower serum 25(OH)D levels in this subgroup of women (Chai, 2010). Nevertheless, Wortsman *et al* (2000) found that obese women taking a vitamin D supplement and being exposed to sunlight still had significantly lower vitamin D levels compared to those of normal weight thus suggesting that vitamin D is either metabolised by different metabolic pathway's or sequestrated in individuals with higher BF%'s.

For most people, the main source of vitamin D is skin exposure from sunlight; however dietary intake is also important in maintaining vitamin D status, especially in New Zealanders during the winter months. Small amounts of vitamin D can be found in oily fish (salmon, tuna, sardines), eggs, liver, and fortified foods such as margarines, milk and milk products, and yoghurt (Ministry of Health, 2015c). The fortification of food items such as milk, yoghurt and margarine is voluntary in New Zealand, however in Australia, England, Scotland, Wales and Ireland it is mandatory. Vitamin D deficiency is still highly prevalent in Australia, especially in those who are obese, indicating the sequestration of vitamin D and / or the different dietary patterns between these groups (Australian Government, 2011, Department of Health, 2011, Daly, 2012).

2.4.4 Iron

Poor iron absorption and the resulting iron deficiency are commonly seen in obese individuals, with the literature continuously showing an inverse relationship between iron status and excess adiposity (table 2.12). Literature surrounding the relationship between iron status and

BF% was reviewed. Studies that included pre-menopausal healthy obese women were reviewed. The most relevant studies exploring this topic are presented in table 2.12. Yanoff et al (2007) found 26.9% of obese subjects to have iron deficiency based on their serum transferrin receptor, compared to 15.7% in the non-obese group. A negative relationship was seen between serum iron, transferrin saturation, C-reactive protein, BMI and fat mass independent of dietary iron intake. It can therefore be speculated that iron deficiency seen in obese individuals is not related to poor diet quality and more so through other metabolic alterations caused by the increase in adiposity. In correlation with this, Cepeda-Lopez et al (2011) concluded that low iron levels in obese individuals is likely to be due to inflammatory mediators influencing iron availability and / or metabolism such as an increase in hepcidin rather than dietary factors. Furthermore, Zimmermann et al (2008) investigated the effect of an iron supplement on iron status in those with excess adiposity, and found a negative correlation between iron status and BMI, showing that an iron supplement is insufficient to prevent iron deficiency in these individuals. This study looked at serum ferritin levels after only one iron supplemented breakfast meal. It implies that inflammatory mediators may play an important part in iron status in these individuals; however a longer supplementation period would be beneficial to determine the long term effects.

A mechanism thought to contribute to poor iron status is the inflammatory-mediated sequestration of iron in these individuals (Yanoff *et al.*, 2007). The chronic inflammation causes increased levels of pro-inflammatory cytokines; interleukin-6 and tumor necrosis factor α thus increasing the expression of hepcidin which may reduce iron absorption (García *et al.*, 2009). Iron requirements in overweight individuals may also be increased due to larger blood volume and higher basal iron losses with increased BF%, but this has not been directly measured (Yanoff *et al.*, 2007).

Author	Study design	Purpose	Participants	Methods	Findings and conclusions
Cepeda-Lopez	Cross-sectional	To examine the	621	Data from the 1999	The prevalence of iron deficiency was significantly higher
<i>et al.,</i> 2011	analysis	relationships	premenopausal	Mexico Nutrition Survey	in the obese women.
		between BMI,	healthy* women,	were analysed.	 Obese women had significantly lower mean serum iron
		dietary iron and	18-50 years; 1174	 Validated 24 hour dietary 	concentrations (62.6 \pm 29.5 $\mu g/dl)$ compared to the
		dietary factors	children, 5-12	recall interview	normal-weight group (72.4 ± 34.6 μg/dl).
Mexico		influencing iron	years.	 Weight, height, BMI 	 Dietary iron intakes were not significantly different
		status.		(kg/m ²)	between BMI groups.
				 Blood and urine samples 	
				(HB, serum iron, TIBC).	
Zimmermann	Cross-sectional	To determine the	92 healthy*	 Intervention group: 4 mg 	Excess adiposity predicts poor iron absorption in women
<i>et al.</i> , 2008	study	relationship	premenopausal	ferrous sulfate consumed	independent of dietary intake.
		between iron	women, 18-50	in the morning $(n = 67)$;	• 4% of participants were anaemic, 20% were iron deficient
		absorption, status	years, maximum	 Control group (n = 27). 	and 22% were overweight.
Thailand		and response to	weight of 70 kg	 Weight 	A negative correlation was found between BMI and iron
		fortification and		 Fasting blood sample (Hb, 	absorption.
		current BMI.		serum ferritin, zinc,	
				transferrin).	
Yanoff <i>et al.</i> ,	Cross-sectional	To examine the	172 healthy* non-	 7-day food record 	Serum iron and transferrin saturation was negatively

Table 2.12 Studies investigating the relationship between iron status and body composition.

Author(s)	Study design	Purpose	Population	Methods	Findings and conclusions
2007	study	relationship	obese (BMI < 30	Diet history questionnaire	correlated with BMI and fat mass.
		between obesity,	kg/m^2) and 234	 Weight, height, BMI 	 Mean transferrin receptor, mean ferritin and c-reactive
		serum iron, iron	obese (BMI ≥ 30	(kg/m²)	protein were higher in obese subjects.
America		stores and	kg/m²) men and	 Body composition (DXA) 	 Adjusted mean daily intake of dietary iron was not
		inflammation.	women, 18 and 70	 10 hour fasted blood 	different between obese and non-obese subjects.
			years.	sample (serum iron,	 26.9% of obese adults had iron deficiency based on serum
				transferrin, Hb).	transferrin receptor compared to 15.7% of non-obese
					subjects.
	*Healthy = Participants free of chronic disease at recruitment; BMI = Body mass index; DXA = Dual-energy x-ray absorptiometry; Hb = Haemoglobin; TIBC = Total Iron Binding Capacity	of chronic disease at recruitr orptiometry; acity	nent;		

2.4.5 Vitamin E

Several studies have investigated vitamin E status in relation to adiposity levels with conflicting results. The literature surrounding the relationship between vitamin E status and BF% was explored. Studies that included women between 16 and 45 years and those that investigated fasting blood samples were reviewed. Few studies were relevant to this thesis and those that were are presented in table 2.13. Chai *et al* (2010) found significantly higher levels of serum γ -tocopherol in the obese subjects and no significant difference between the groups when comparing serum α -tocopherol levels. Y-tocopherol has anti-inflammatory properties and thus its levels would rise in response to inflammatory signals in the body. The elevated serum γ -tocopherol levels in these obese individuals may therefore reflect systemic inflammation which is present with excess adiposity (Chai, 2010). Garcia *et al* (2009) also found a positive relationship between vitamin E levels and BMI, waist circumference and waist-to-hip ratio. Overweight and obese women had significantly higher vitamin E levels compared to the women of normal body weight. This significance however disappeared after adjusting for total cholesterol and triglycerides, as did the association between vitamin E levels and anthropometrical measurements.

In contrast to the above, several studies have reported low vitamin E levels in obese women. Vitamin E is important for the protection of LDL against oxidation, thus playing a preventative part in CVD development. Low vitamin E levels therefore have been associated with increased levels of oxidative stress, insulin resistance, impaired glucose metabolism and some cancers (Kimmons, 2006). Both Kimmons *et al* (2006) and Mehmetoglu *et al* (2011) reported significantly lower vitamin E levels among obese women compared to those of normal weight. A study by Viroonudomphol *et al* (2003) found 10.7% of overweight/obese women showed significantly lower serum concentrations than those of normal weight. This was in combination with the presence of dyslipidaemia in these subjects where 39.3% had total cholesterol levels above or equal to 6.48 mmol/L and 51.8% had LDL cholesterol levels at or above 3.89 mmol/L.

2.4.6 Zinc

Few studies have investigated the effect of obesity on zinc; however those that have, reported negative correlations. The literature surrounding the relationship between zinc status and BMI was explored. Studies that included women between 16 and 45 years were reviewed. The studies that were relevant to this thesis are presented in table 2.13. Zinc plays an important function in regulating the metabolism of protein, carbohydrates, lipids and nucleic acids in the body (Ennes Dourado Ferro, 2011). Zinc levels may be lacking in many obese individuals and

this is thought to be linked to insulin resistance, hyperglycaemia and impaired glucose tolerance (Ennes Dourado Ferro, 2011). Ennes Dourado Ferro *et al* (2011) found the mean intake of zinc in obese women was greater than women of normal weight, however there was no significant difference in the zinc status between the two groups. This may be due to the study having only looked at zinc levels in plasma. Around 80% of zinc is found in erythrocytes and since erythrocytes have a half-life of approximately 120 days, this is a more sensitive indicator of zinc status in individuals. Further to this Marreiro *et al* (2006) found that despite the good zinc intakes through dietary sources, obese subjects still presented with low serum levels.

2.4.7 Vitamin C

A positive relationship between Vitamin C levels and BMI has been documented (Kimmons, 2006). The literature surrounding the relationship between vitamin C status and BMI was explored. Studies that included women between 16 and 45 years and those that investigated were reviewed. Few studies were relevant to this thesis however those that were are presented in table 2.13. Both Garcia *et al* (2012) and Johnston *et al* (2007) observed an inverse relationship between vitamin C levels and BMI and waist circumference in women. This relationship is thought to be due to the effect vitamin C has on leptin expression. A low vitamin C level was seen with higher leptin concentrations in all women and was still present after adjusting for BMI (table 2.13).

Mehmetoglu <i>et al.,</i> 2011	oruuy uesigii	Purpose	Participants	Methods	Findings and conclusions
<i>et al.,</i> 2011	Cross-sectional	To investigate the	21 male and 77	• FFQ	Vitamin E levels of the obese females were significantly
	study	relationship	females, BMI > 30	 Weight, height, BMI 	lower compared to the normal weight female subjects.
		between plasma	kg/m^2 ; 20 male and 58	(kg/m ²).	 Adjusted vitamin E levels had a significant negative
		vitamin A, E, serum	females, BMI < 25	 Fasting blood sample 	correlation with BMI, waist circumference and waist-to-hip
		coenzyme \mathbf{Q}_{10}	kg/m ² , 18-65 years.	(serum CoQ ₁₀ , vitamin A	ratio in the control group.
Turkey		levels and insulin		and E, TC, triglycerides,	 Adjusted vitamin E levels were negatively correlated with
		resistance in obese		HDL-c, LDL-c, blood	waist circumference and waist-to-hip ratio in the obese
		and normal weight		glucose).	group.
		individuals.			Vitamin E was positively correlated with age in obese
					group.
Garcia, 2012a	Cross-sectional	To determine the	580 women, 37 ± 7.5	 Weight, height, WC, HC 	High leptin concentrations were associated with lower zinc
	study	association of zinc,	years	 3x 24-hour recalls 	and vitamin C levels when stratifying by BMI and BF%.
		vitamin A, C and E		 Body composition 	 Vitamin E concentrations were positively associated with
		concentrations		(DXA)	adiposity, BMI, WC and W:H ratio.
		with BMI, central		 Fasting blood sample 	 Vitamin E concentrations were significantly lower in
Mexico		adiposity, body fat		(glucose, TC, HDL, LDL,	women with lower weight, however after adjusting for
		and leptin levels.		TGs, leptin, Vitamin A,	lipid concentrations this was no longer significant.
				C, E and zinc.	 Overweight and obese women consumed significantly

Table 2.13 Studies investigating vitamin E, zinc, vitamin C and their relationships with body composition.

Author	Study design	Purpose	Participants	Methods	Findings and conclusions
					more energy and total fat but similar micronutrient intakes
					as normal weight women.
Ennes	Case-control	To determine the	36 normal weight (BMI	 Height, weight, BMI 	Erythrocyte zinc concentrations are inversely associated
Dourado	study	relationship	$18.5 - 24.9 \text{ kg/m}^2$) and	(kg/m²)	with WC and BMI.
Ferro, 2011		between	37 obese (BMI > 30	 3-day food record 	No significant differences were seen between the two
		biomarkers of the	kg/m^2) healthy*	 Fasting blood samples 	groups and their plasma zinc concentrations.
		metabolic	premenopausal	(plasma and	• Mean zinc intake in the obese women was 10.8 \pm 4.7 mg/d
Brazil		syndrome and zinc	women, 20-50 years.	erythrocyte zinc levels,	and 7.6 \pm 2.9 mg/d in normal weight women.
		status in obese		lipids, fasting glucose)	
		women			
Johnston <i>et</i>	Cross-sectional	To examine the	35 men and 83	 Height, weight, BMI 	An inverse correlation was seen between plasma vitamin C
<i>al.</i> , 2007	study	relationship	women, 38.7 ± 1.0	(kg/m²)	levels and BMI, BF% and WC in both men and women.
		between vitamin C,	years, BMI 18.7 – 47.9	 Body composition (BIA) 	This significant inverse relationship remained in women
		adiponectin, fasting	kg/m ²	 Fasting blood samples 	after controlling for body mass.
		insulin and weight		(serum vitamin C)	
America		in healthy adults.			
Marreiro <i>et</i>	Prospective	To assess the effect	56 obese normal	Group 1 received 30 mg	Low plasma zinc concentrations were seen in obese
<i>al.</i> , 2006	double-blind,	of zinc	glucose tolerant	zinc aminochelate daily	individuals.
	randomised,	supplementation	women, 25-45 years,	for 4 weeks. Group 2	Dietary intake of zinc was adequate and obtained through

Author	Study design	Purpose	Participants	Methods	Findings and conclusions
	clinical,	on serum leptin	BMI 36.2 ± 2.3 kg/m ²	received a placebo.	mainly meat in the obese individuals.
	placebo-	levels in insulin		 Body composition (BIA) 	A non-significant increase in plasma zinc concentrations
Brazil	controlled	resistance.		 Fasting blood samples 	was seen in supplemented group.
	study.			(zinc status, leptin and	
				insulin levels) pre and	
				post intervention	
				period	
Viroonudomp	Cross-sectional	To examine the	Intervention group: 16	 Weight, height, BMI 	- Overweight and obese women had lower α -tocopherol
hol <i>et al.</i> , 2003	study	association	males and 56 females,	(kg/m ²), WC, HC, W:H	levels compared to the normal weight control group.
		between	BMI ≥ 25.0 kg/m ² , 16-	ratio, MAMC	• 10.7% of overweight and obese females had decreased $\alpha\text{-}$
		anthropometric	60 years.	 Fasted blood sample 	tocopherol levels.
		measurements,	Control group: 14	(serum retinol, α-	• A negative correlation was found between weight, BMI,
Thailand		antioxidant status	males and 58 females,	tocopherol, lipid	hip circumference and $lpha$ -tocopherol in both overweight
		and lipid profiles.	BMI 18.5-24.9 kg/m ² ,	profiles)	and obese females.
			16-60 years.		
*Healt [†] BMI = E WC = W	*Healthy = Participants free o BMI = Body mass index; WC = Waist circumference;	*Healthy = Participants free of chronic disease at recruitment; BMI = Body mass index; WC = Waist circumference;	nent;	W:H ratio = Waist to hip ratio; MAMC = Mid arm muscle circumference; BIA = Bioelectrical impedance analysis; Tr. Total cholecterol	DXA = Dual-energy x-ray absorptiometry; ;

2.5 Assessing dietary intake

There are generally considered to be five main techniques used to assess dietary intake (table 2.14). The 24-hour recall involves a trained interviewer asking the participant to describe in detail their intake in the last 24 hours. A food frequency questionnaire requires the participant to report on the frequency that they consume an extensive list of foods in the past number of days, months or year. A food record can be estimated intake or weighed intake by the participant and reports the time, type and the amount of each food eaten each day, usually ranging from one to seven days. Finally, a diet history involves the participant to report on their usual intake over an extended period of time, for example the past month or year. The strengths and weaknesses of each method are shown in table 2.14 (Lee, 2010).

Table 2.14. Strengths and weaknesses of the five main dietary assessment techniques (Lee,2010)

Dietary assessment technique	Strengths	Weaknesses
24-hour recall	 Fast and easy to administer Inexpensive Low respondent burden Can provide detailed information on types of food consumed Can be used for groups 	 Relies on memory Misreporting occurs One recall is not representative of an individual's usual intake
Food frequency questionnaire	 Data represents usual intakes Small participant burden Can be self-administered Inexpensive for large sample sizes Can identify dietary patterns Can be used for groups 	 Relies on the ability of respondent to describe their diet May not represent participant portion sizes
Weighed food record	 Does not depend on memory Can be very accurate and provide detailed intake data Can provide data on eating habits Multiple day data can be representative of usual intake 	 Requires a high degree of cooperation and compliance Act of recording may alter diet Large participant burden Good literacy skills required Takes more time to obtain data Data analysis labour intensive and expensive
Estimated food record	 Does not depend on memory 	Estimated not absolute therefore room for error

Dietary assessment technique	Strengths	Weaknesses
	 Can provide data on eating habits Multiple day data can be representative of usual intake 	 Act of recording may alter diet Good literacy skills required Data analysis labour intensive
Diet history	 Assesses usual intake Can detect seasonal changes Correlates well with biochemical measures 	 Long interview process Required trained interviewers High degree of cooperation required

2.5.1 Misreporting of energy intake

It is well documented that under-reporting of energy intake through self-reported dietary assessment is common (Bedard et al., 2004). This is a concern in any dietary assessment, especially when estimating habitual intake as it has the ability to introduce bias (Bedard et al., 2004). Factors which influence under-reporting include weight status, age, gender and lifestyle such as smoking and physical activity as well as psychological factors (Bedard et al., 2004, Hyun, 2007, Johansson et al., 1998, Mendez et al., 2004). Under-reporting could also be explained by under-recording, under eating during the assessment period, measurement error, lack of accuracy recorded by the measurement tools provided and participant motivation and comprehension of how to complete the questionnaire (Bedard et al., 2004, Neuhouser et al., 2008, Kipnis et al., 2003). Furthermore, specific food groups may be more likely to be influenced by misreporting depending on how they are perceived. Foods that are perceived as unhealthy (e.g. those rich in refined carbohydrate, fat and sugar such as cakes, confectionary and biscuits) and snack foods may be more likely to be under-reported compared to those thought of as healthy which may be over reported (e.g. fruit and vegetables) (Johansson et al., 1998, Svendsen and Tonstad, 2006). This is thought to be due to the association of these foods with a poor health image, individuals wanting to fit into the acknowledged ideal eating behaviours, and also wanting to please the researcher (Macdiarmid, 1998, Margetts, 2004). Women who under-report have been found to be more likely to report foods which are higher in carbohydrate and lower in fat compared to those who do not under-report their intakes, and obese women have been shown to under-report total energy intake compared to nonobese individuals, possibly leading to skewed predictors of chronic disease risk (Bedard et al., 2004, Gonelevu et al., 1997).

2.5.2 Recovery markers

Recovery markers are commonly used in the literature to measure the accuracy of selfreported energy and protein intakes from dietary assessment tools. The two main methods used are the doubly labelled water protocol (energy biomarker) and a 24-hour urine collection (protein biomarker). The doubly labelled water protocol is considered the gold standard to measure the energy expenditure of individuals under free-living conditions (Poslusna *et al.*, 2009). It uses the stable isotopes ²H and ¹⁸O to measure CO₂ production rate from which energy expenditure can be determined (Hill and Davies, 2001). Using the 24-hour urine collection method, nitrogen values are measured in the urine to determine the 24 hour protein intake of individuals. The nitrogen excretion of individuals can however vary widely and therefore this method needs to be performed several times to validate the protein intake of individuals (Poslusna *et al.*, 2009).

2.5.3 Goldberg cut-off

Another method to identify and assess the impact of reporting bias is the Goldberg equation. This has been routinely used to predict total energy expenditure and identify those weight stable individuals that have under-reported their energy intake. It is based on determining the ratio of reported energy intake (EI) to the basal metabolic rate (BMR) of the individual (EI: BMR). The BMR is determined using the age, weight and sex through the Scholfield equation. Under-reporting is identified by using the minimum energy requirements for survival in sedentary individuals (EI <1.35 x BMR). Over-reporting is based on the energy requirements for extremely physically active individuals (EI > 2.4 x BMR). Although the doubly-labelled water technique is the gold standard for assessing individual energy expenditure, it is impractical to be used in large studies due to the cost of materials and the time and expertise required analysing the data. The Goldberg equation, although estimated, is a fast and more cost-effective alternative which is therefore routinely used in the literature (Bedard *et al.*, 2004).

Obesity is a worldwide health problem which increases the risk of many chronic diseases. Specific dietary patterns and macronutrient distributions have been strongly linked to increased adiposity, as have the consumption of energy-dense foods such as sugar- sweetened beverages and fast-food. Further, individuals who have a poor diet quality tend to have a higher risk of obesity. Investigating the dietary intake and eating patterns of New Zealand women will provide greater understanding of the development of different body fat profiles and how we can tailor our dietary advice according.

Chapter 3 Methods

3.1 EXPLORE Study Design

This study was a sub-study of the wider cross-sectional comparative designed study 'Examining Predictors Linking Obesity Related Elements' (EXPLORE) in young adult women. The EXPLORE study investigates how bodyweight and body fat profiles are related to the risk for chronic disease in premenopausal women as well as to investigate how diet and physical activity patterns may impact these profiles and how they may affect miRNA associated with body fat usage and storage. It aimed to recruit225 post-menarcheal, premenopausal New Zealand European (NZE), Maori and Pacific Island women (defined by at least one parent being of the ethnicity) to make a total of 675 participants. The EXPLORE study aimed to recruit an equal number of women (75 women) in each of the three body composition profile (BCP) groups outlined in table 3.1.

Table 3.1 Bod	y composition	profile (BCP) groups	(Kruger, 2015).
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BCP group	Body mass index (kg/m ²)	Body fat percentage (BF %)
"Normal fat" group (NN)	Normal (18.5-24.9 kg/m ²)	Normal (≥22%, <30%)
"Hidden fat" group (NH)	Normal (18.5-24.9 kg/m ²)	High (≥30%)
"Apparent fat" group (HH)	High (≥25 kg/m²)	High (≥30%)

BCP group = Body composition profile group; NN = Normal BMI, normal BF%; NH = Normal BMI, high BF%;

HH = High BMI, high BF%

This sub-study involved the New Zealand European (NZE) women only and investigated the dietary data obtained from the New Zealand Women's FFQ (NZWFFQ) that was validated in another sub-study of the EXPLORE trial (add ref). At the screening stage, women were categorised into the three BCP groups as outlined in table 3.1 using bio-electrical impedance (BIA) in a non-rigorous community setting. When women participated in data collection for the study, their body composition assessment was conducted using a strict and rigorous ADP protocol, resulting in further profile groups being identified. In addition to the three main BCP groups outlined in table 3.1, this sub-study therefore also included women who had a high BMI with a normal BF% and a normal BMI with a low BF%. As a result the dietary data from women characterising five different body composition profiles were analysed:

- NN normal BMI (18.5-24.9 kg/m²); normal BF% (≥22%, <30%)
- NH normal BMI (18.5-24.9 kg/m²), high BF% (≥30%)
- HH high BMI (≥25 kg/m²), high BF% (≥30%)
- HN high BMI (≥25 kg/m²), high BF% (≥30%)
- NL normal BMI (18.5-24.9 kg/m²), low BF% (<22%).

These five body composition profiles were included in the dietary analysis describing dietary intakes and patterns of the NZ European cohort to provide a complete dietary assessment of all profiles.

3.2 Ethical Approval

This study protocol was reviewed and approved by the Massey University Ethics Committee: Southern A, Application 13/13. The participants were provided with information sheets and consent forms which were completed prior to data collection. These documents provided the participant with information about the protocols and procedures of the study as well as outlining the inclusion and exclusion criteria for participating in the study, as well as their duties and obligations as participants in the study.

3.3 Study Population

3.3.1 Participants

The wider EXPLORE study aimed to recruit two hundred and twenty five New Zealand European women (75 per profile group). This will provide 80% power at significance levels of p < 0.05 to detect a medium effect *f* size of 0.25. It was calculated that approximately 380 NZE women needed to be screened to recruit the 75 eligible women per profile group (Kruger, 2015). Oversampling was performed to ensure that sufficient participants were recruited.

3.3.2 Recruitment

Participant recruitment was conducted over several months (August 2013 - December 2014) in parallel with data collection. The NZE participants were approached through a variety of routes including media and advertising (e.g. local newspapers, websites and local radio stations) and posters and flyers distributed throughout a variety of venues including community groups and churches, medical centres, community events, schools, organisations, and businesses. Social media (e.g. Facebook), emailing lists (e.g. Massey University staff, students, previous research participant databases) and word of mouth were also used throughout the Auckland region. The women were directed to the study website for further information and to register their interest of participation. The aim was to recruit a convenience sample of NZE, Maori and Pacific Island women.

Pre-screening process

Participants who registered their interest were provided with an information sheet and completed an online health screening and demographics questionnaire through Survey Monkey (or a paper version depending on internet availability) to determine eligibility. This process took 15-30 minutes and included questions on age, ethnicity, parent ethnicity, current and previous health issues (e.g. breastfeeding, diabetes, and cancer) and the regularity and date of their last menstrual cycle (appendix A). If the inclusion criteria were met, participants were deemed eligible and were invited to participate in the screening phase (phase 1) at Massey University, Albany, Auckland. Participants who met the exclusion criteria were contacted via phone or email and their body composition results sent to them.

Inclusion criteria

- NZE women between 16-45 years old
- Post-menarche (defined as having at least one full year or regular menstrual cycle)
- Premenopausal (defined as having had a continuous menstrual cycle for the last complete year)

Exclusion criteria

- Pregnancy
- Lactation
- Any diagnosed chronic disease/illness
- Those actively losing or gaining weight
- Milk allergy

3.3.3 Study setting

Recruitment took place off-site in various areas of Auckland, such as Albany, Henderson and Mangere. The study took place in the Human Nutrition Research Unit (HNRU) at Massey University, Albany Campus, Auckland.

3.4 Procedures

The study entails two phases: screening (phase 1) and main data collection (phase 2).

3.4.1 Phase 1

Screening process

Screening took place in the HNRU at Massey University Albany Campus, North Shore Auckland or off-site at a community venue convenient and appropriate for the participants. Simple anthropometric data of height, weight and body fat percentage (BF%) measurements needed to be obtained to allow for the body composition screen. Height was measured using a portable stadiometer. Weight and BF% were obtained using bioelectrical impedance analysis (BIA) machine (Biospace, Inbody 230, Cerritos, CA). The BIA was used during screening because it provided a non-invasive accurate way to quickly determine the participants body composition. For this measure participants were not fasted, were fully dressed and the time of day was not standardised. The BIA results were not shown to the participants to ensure a blinded study.

Participant's BMI (calculated using height and weight) and BF% results were used for preliminary categorisation into one of the three BCP groups. Eligible participants were invited back to the HNRU at Massey University via phone or email for further testing in phase 2.

3.4.2 Phase 2

Data collection

Participants recruited in phase 2 visited Massey University Albany campus for approximately 2 hours arriving between 7 am and 9:45 am. For the purpose of this study, participants completed:

- Health questionnaire and consent forms
- Anthropometric measures including height, and hip and waist circumference
- Body composition assessment using air displacement plethysmography (ADP) via the BODPOD (2007A, Life Measurement Inc, Concord, Ca., using software V4.2+ as supplied by the manufacturer).
- A validated food frequency questionnaire (FFQ) to assess dietary intake
- A validated New Zealand habits questionnaire to assess dietary and eating habits

Further details of the wider study methodology are outlined elsewhere (Kruger, 2015).

3.5 Body Composition

A full body composition analysis was conducted via air displacement plethysmography (ADP) (2007A, Life Measurement Inc, Concord, Ca., using software V4.2+ as supplied by the manufacturer) using the thoracic gas volume method (Noreen, 2006, Wingfield *et al.*, 2014). This analysis was performed between 8 am and 9:30 am. Participants were fasted overnight (for at least 12 hours) and had not drunk any fluids prior to the ADP analysis. The participants wore tight fitting clothing and a cap to cover hair and minimise its interference with the results. The ADP analysis was conducted by a qualified ADP operator. The body fat results from ADP were used for final categorisation of participants into one of the five BCP groups (Wingfield *et al.*, 2014, Noreen, 2006).

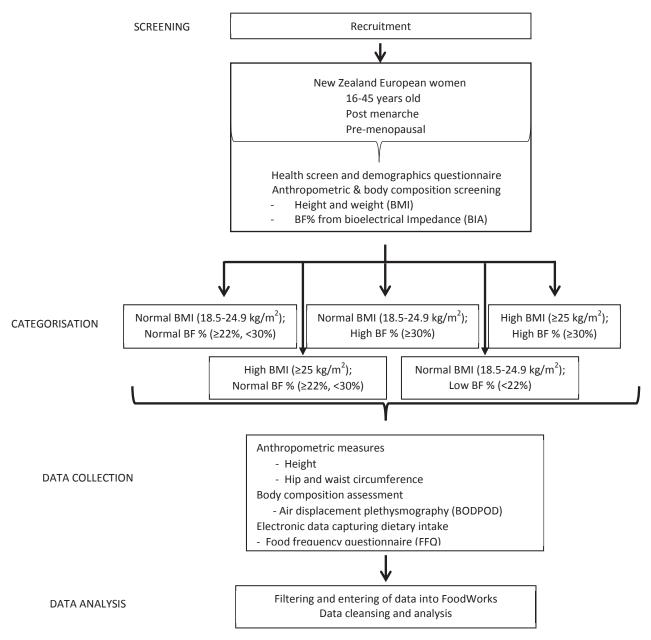


Figure 3.1 Methodological overview of the study.

3.6 Dietary Questionnaires

The participants completed a validated 220-item self-administrated semi-quantitative food frequency questionnaire (FFQ) and a validated eating habits questionnaire (EHQ). These were completed in the Human Nutrition Research Unit at Massey University Albany, Auckland campus. The questionnaires were designed and administered using online computer survey software (Survey Monkey). The questionnaires were completed online which minimises interpretation and coding errors, and also ensures a maximum response rate throughout the questionnaire by minimising missing data (Van Selm and Jankowski, 2006). Nevertheless, paper copies of the questionnaires were provided if required which were checked by the supervising research assistant to ensure all questions were answered.

3.6.1 Food frequency questionnaire

The validated, 220-item, self-administrated FFQ (appendix C) was used to assess energy, macronutrients and selected micronutrient intakes of participants over the previous month. It included food lists to encompass frequently consumed foods in each food category (e.g. dairy, breads and cereals, meat, fish and poultry, fats and oils, fruit and vegetables, drinks, dressings and sauces, miscellaneous (cakes, biscuits, spreads) and takeaways), using standard natural portions (e.g. 1 banana) or standard weight and volume measures (e.g. 45 g of oats, 1 pottle of yoghurt). Additional questions included information on usual food preparation methods (e.g. the removal of skin off chicken and cutting excess fat off meat), cooking practises (e.g. the use of oil or butter; frying or baking) and types of food items regularly consumed (e.g. white, wholemeal or wholegrain bread). The FFQ was validated as another sub-study of the EXPLORE study (thesis published; manuscript in preparation). Information on the development and validation of the FFQ is outlined in a Massey University Master's thesis by Houston (Houston, 2014).

Prior to the commencing the questionnaire, participants were provided with verbal instructions following the Standard Operating Procedures (SOP) (appendix B). The research assistant explained the first page of the questionnaire, emphasising the most important points e.g. base your answers on what you eat yourself and not others in your household, and explained two example questions to check participant comprehension (figure 3.2). The participant identification number was checked and entered by the research assistant who was also available for questions throughout completion of the questionnaire.

1. EXAMPLE: How often do you usually have sugar? (Please do not fill out) 1-3x/ 1x/week 2-3x/week 4-6x/week Once/day 2-3x/day 4+x/day <1x / month month Sugar - 1 tsp \bigcirc \bigcirc ()()If every day you have 2 cups of coffee with 1 tsp sugar, 4 cups of tea with 1 tsp sugar, one bowl of cereal with 1 tsp sugar and sugar on pancakes at dinner, you would choose four or more times per day = '4+ x / day'. Adjust your portion size and frequency of intake to suit your eating habits. 2. EXAMPLE: How often do you usually eat bread? (Please do not fill out) 1-3x / <1x / month 1x/week 2-3x/week 4-8x/week Once/day 2-3x/day 4+x/day Never Bread - 1 slice \bigcirc \bigcirc ()If every day you have two slices of toast for breakfast, and you have a sandwich for lunch three times per week, you would choose two - three

Adjust your portion size and frequency of intake to suit your eating habits.

times per day = '2-3x / day'.

Figure 3.2 Food Frequency Questionnaire (FFQ) example questions explained to participant prior to completion.

For each food item, participants reported their average frequency of consumption during the previous month, keeping in mind the standard serving size. This was done by choosing one of nine frequency categories as outlined in table 13. Data was collected from July 2013 to December 2014 therefore all four seasons were captured, thus reducing any discrepancy between seasonal food availability and eating patterns.

3.7 Data Analysis

Food frequency questionnaire data was downloaded from SurveyMonkey into an excel spreadsheet and rearranged into a comprehensive document. All frequencies were translated (table 3.2) and entered into Foodworks 7 (FoodWorks Professional, 2013) dietary analysis database by two MSc students and one research assistant using a template developed specifically for this purpose. Foodworks utilises the New Zealand Food Composition Database (NZ, FOODfiles 2010) developed by New Zealand Plant and Food Research to determine energy, macro- (carbohydrate, protein, total fat, saturated fat) and micronutrient intakes (vitamins A, D and E, C, B6, B12, folate, thiamin, riboflavin, and niacin, dietary fibre), and to assess diet quality. When specific food items were missing from the database, a substitute was entered (e.g. a diet energy drink was entered as diet soft drink). Where specific recipes were required (e.g. protein shake) a new recipe was entered manually e.g. 100g skim milk powder and 250 mL water. The data was checked twice by one of the MSc students to ensure data entry accuracy. Assumptions made during data entry are recorded in Appendix D.

Table 3.2 Frequency translations.

Frequencies	Translation
Never	0m
<1x/month	0.25m
1-3x/month	2m
1x/week	1w
2-3x/week	2.5w
4-6x/week	5w
Once/day	1d
2-3x/day	2.5d
4+x/day	4d

M = Month;

W = Week;

D = Day

Data obtained from the FFQ will be compared to the New Zealand Ministry of Health nutrient reference values (NRVs) for Australia and New Zealand. Specifically, the estimated average requirements (EARs) for women between 19 and 50 years will be used. These values are based on the nutrient requirements estimated to meet the needs of half the healthy population and are used to assess inadequate intake of nutrients within a group setting (National Health Medical Research Council, 2005). The recommended daily intake (RDI) values are set at a higher threshold (2 Standard deviations above the mean) to accommodate for variations in absorption and metabolism of nutrients (National Health Medical Research Council, 2005). They are sufficient to meet the daily needs of 97-98 per cent of healthy individuals, therefore are not appropriate to assess group dietary intakes (National Health Medical Research Council, 2005). Nevertheless, RDI values will still be included to compare nutrient intakes in relation to a higher threshold.

3.8 Food Groupings

All food items in the FFQ were categorised into 56 food groups based on similar nutritional composition and characteristics (table 3.3). Consideration was also given to those foods which are typically consumed together (e.g. crumbed and deep fried) and other individual foods and beverages which are typically or commonly consumed. Individual foods that solely create a food group had to be consumed by at least 10% of the study population. The amount of food groups created were based on the recommendation by Kass & Tinsley (1979) to allocate 5-10 participants per food grouping.

Table 3.3 Food groups

Food Group	Food items included	Justification
Full fat milk	Full fat milk (dark blue top)	High fat content compared to other milks
Low fat milk	Lite milk (light blue top), trim milk (green top)	Often chosen on low fat properties
Sweetened milk	Breakfast drinks, flavoured milk, evaporated	Sugar added
products	milk, hot chocolate	
Yoghurt	Yoghurt (plain, fruity, Greek, unsweetened)	Separate group to other dairy products
High fat cheese	Cheddar, processed cheese, cream cheese, blue	High fat content
	vein	
Low fat cheese	Edam, cottage cheese, brie, camembert, feta	Lower fat content
Apple, banana, orange	Apple, banana, orange	Common fruit available all year round in
		NZ
Other fruit	Fresh, canned, frozen, dried	Seasonal fruits
Tomatoes	Tomatoes	Good source of lycopene, eaten by >10%
		of people so can be a group on its own
Dark-yellow vegetables	Carrots, pumpkin	Good source of vitamin C
Green vegetables	Lettuce, spinach, cabbage, broccoli, watercress,	Good source of B vitamins
0	green beans, sprouts, courgette	
Other non-starchy	Capsicum, onion, mushrooms, frozen mixed	Other non-starchy vegetables
vegetables	vegetables, beetroot	, , ,
Potatoes (excluding	Potato (boiled, mashed, baked, stuffed,	Commonly eaten starchy vegetable,
chips and crisps)	scalloped)	eaten by >10% of people so can be a
	Sunopeay	group on its own
Starchy vegetables	Kumara, yam, parsnip, turnip, swedes (boiled,	Other starchy vegetables
Stareny vegetables	mashed, baked)	other starting vegetables
	Taro (flesh, roots, stalks), green banana, sweet	
	corn kernels	
White broads		Refined and processed staple bread
White breads	White bread, wraps, fruit bread, focaccia, bagel,	
Discustion on the set	pita, paraoa bread, rewena bread, doughboys	products, low fibre
Discretionary breads	Crumpets, scone, savoury muffin, croissant,	Refined and processed discretionary
<u> </u>	pancakes/waffles, iced bun	items, low fibre
Crackers	Cream, cruskit, corn, rice, vitawheat	Different nutrient composition to bread
		and grains
Whole grain breads	Bread (high fibre, wholemeal, wholegrain)	High fibre
Refined grains	White rice, pasta (penne, spaghetti, vermicelli),	White refined carbohydrates
	noodles (instant, egg, rice), canned spaghetti	
Wholegrains	Brown rice, quinoa, couscous, bulgur wheat	Wholegrains, high fibre
Oats	Porridge, rolled oats, oat bran, oat meal	High fibre, no added sugar
Sweetened cereals	Milo cereal, coco pops, nutrigrain, honey puffs,	Added sugar, highly processed
	fruit loops, special K, light and tasty, sultana bran	
Red meats	Beef (mince dishes, casserole, stew, stir-fry,	Red meats high in protein and saturated

Food Group	Food items included	Justification
	roast, steak)	fat
	Lamb (stew, casserole, stir-fry)	
	Venison, hogget (roast, chops, steak, casserole,	
	stew, stir-fry), offal (liver, kidney, pate)	
White meats	All chicken (breast, leg, wing, casserole, stir-fry)	White meats high in protein
	Turkey/quail, pork (roast, chop, steak), mutton	
	bird/duck, veal	
Processed meats	Sausages, frankfurters, saveloys, cherrioes,	All processed, high in saturated fat and
	bacon, ham, luncheon meats, salami, chorizo,	salt
	meatloaf, corned beef, patties	
Fish and seafood	Canned salmon, canned tuna, canned mackerel	All fish, high in protein
	Snapper/hoki, gurnard, shark, tuna, salmon	
	Shrimp/prawn, crab, mussels, pipi, whitebait,	
	kina, squid	
Egg and egg dishes	Eggs, egg mixed dishes (omelette, quiche,	All egg protein based
	frittata, other baked egg dishes)	
Legumes	Canned/dried (lentils, chickpeas, peas, beans,	All legumes
	baked beans), hummus, dahl	
Soy products	Soybeans, tofu	Soy higher protein than legumes
Peanut butter and	Peanut butter, peanuts	Different nutrient composition to plain
peanuts		nuts and seeds – sugar and salt added
Nuts and seeds	Nuts (peanut, brazil, walnut, almond, cashew,	All nuts and seeds
	pistachio)	
	Seeds (pumpkin, sunflower)	
Fats	Butter, lard, dripping, ghee	All solid fats
Coconut fats	Coconut milk, cream, oil	Topical
Oil and oil-based	Canola, sunflower, olive, vegetable oils, cooking	Avocado classed as a fat rather than a
dressings	sprays	vegetable
	Salad dressing (French, Italian), Avocado	
Margarine	Margarine – all types	Different to butter and other fat spreads
Creamy dressings	Mayonnaise, creamy dressings, white/cheese	Richer and higher fat content than the
	sauce, sour cream	sauces
Sauces	Tomato, barbeque, chilli, mint, soy, gravy,	Less saturated fat than the creamy
	mustard, chutney, instant soup	dressings
Sweet spreads	Jam, honey, marmalade	High sugar and energy content
Savoury spreads	Vegemite, marmite	Lower energy and sugar
Cakes and biscuits	Cakes, loaves, muffins, sweet pies, pastries, tarts,	All baked sweet discretionary items
	doughnuts, biscuits (plain, chocolate coated)	

Food Group	Food items included	Justification		
	instant)	a dessert item		
	Other non-dairy based puddings (pavlova, sticky			
	date pudding), jelly, ice blocks			
Sweet snack foods	Chocolate, lollies, muesli bars	Sweet snack foods		
Savoury snack foods	Potato chips, corn chips, twisties	Savoury snack foods		
Crumbed and deep	Crumbed chicken/fish, battered fish, potato fries,	These are typically eaten together in a		
fried	chicken nuggets	meal		
Fast-food	Meat pie, sausage roll, savouries, burgers, kebab,	Bought high fat fast-foods		
	Chinese, Indian, Thai, Pizza			
Fruit juice	Fruit and vegetable juice	Higher fruit content than fruit drinks		
Fruit drink and other	Fruit drink, sparkling grape juice, cordial, iced	Higher added sugar		
beverages	tea, energy drinks, sports drinks, flavoured			
	water, soft drinks			
Diet drinks	Diet energy drinks, diet soft drinks, diet cordial	Low calorie drink options		
Теа	Black tea, herbal tea	May drink tea not coffee		
Coffee	Instant coffee, brewed water-based coffee,	May drink coffee but not tea		
	espresso			
Beer	Standard, low alcohol	May drink beer but not wine		
Wine (red and white)	White wine, red wine	May drink wine but not beer		
Water	Water	Essential in the diet, no energy		
		component, different to other beverages		
Other alcoholic	Cider, spirits, sherry, port, ready-to-drink	All other miscellaneous alcoholic		
beverages	alcoholic sodas (RTD's), kava	beverages		
Sugar added to food	White sugar	Sugar specifically added to foods		
and drink				

3.9 Data Accuracy

3.9.1 Misreporting

The Goldberg equation was performed to highlight participants who misreported their energy intake in relation to their energy requirements in the FFQ. Schofield's equations were used to determine participant basal metabolic rates (BMR) (kilocalories per day) using individual age, gender, weight and height (table 3.4). A physical activity level (PAL) cut off point of 1.55 was used for all participants. This cut off point was determined to be a conservative value which represented seated work and very little strenuous leisure activity (Black, 2000). To derive the cut offs for evaluation of misreporting of energy intakes the SD_{min} and SD_{max} values of -2 and +2 were used for 95% lower and upper confidence intervals, respectively. The revised factors outlined in Black (2000) were also used in this analysis.

Literature reports large bias in energy reporting of overweight and obese individuals (Schoeller *et al.*, 1990). To ensure accuracy participants who were identified as misreporters were excluded in the nutrient analysis but were included in the pattern analysis.

Table 3.4 Scholfield equations.

Age	BMR equation (kcal/d)
10-17	8.4wt + 466ht + 200
18-29	13.6wt + 283ht + 98
30-59	8.1wt + 1.4ht + 844

BMR = Basal metabolic rate Wt = Weight Ht = Height Kcal/d = Kilocalories per day

3.10 Statistical Analysis

The NZWFFQ FoodWorks dietary data were analysed using SPSS 21.0 for windows (SPSS Inc, Chicago IL) computer software system. Data was checked for normality using histogram plots as well as skewness and kurtosis values. Values between -1.96 and 1.96 suggest that the data follow an approximately normal distribution and that no significant difference between the test distribution and a normal distribution is present. Due to the population sample size (n = 231) the Kolmogorov-Smirnov and Shapiro-Wilk tests were not used as they are limited to a sample size of 50 (Royston, 1982). The data was also checked for normality visually, using a superimposed bell curve shape on the histogram, the boxplot, de-trended plot and the Q-Q normality plot. Where improvements to the data were required to improve normality, a natural logarithmic transformation was performed. Descriptive statistics were reported as the mean and standard deviation (SD) for normally distributed data, geometric means and 95% confidence intervals (CI) for the log transformed data, and where normality was not able to be reached, median values and lower and upper quartiles were reported. Other variables were reported as numbers, frequencies and percentages.

3.10.1 Frequency of food item consumption

The 40 most frequently consumed food items were determined for the total population and the three main (NN, NH, HH) profile groups. This included the 26 women identified as under and over reporting their energy intake who subsequently were also included in the food pattern analysis. The HN and NL BCP groups were not included due to small sample sizes (3 and 18 participants, respectively) thus the results would not have been representative of all individuals with these body compositions. The top 40 foods were analysed to identify differences in food items most commonly consumed between the BCP groups extending out from the main food group items (e.g. bread, milk, oil). Frequencies of consumption for the 220 foods investigated in the FFQ were converted into daily equivalent frequencies and the average frequency of consumption was calculated. The mean daily intake (g per person per day) for each food item was determined by taking the average of the food item weights. Additionally, the number of participants who consumed each of the top 40 foods daily was determined.

3.10.2 Food pattern derivation

Dietary patterns were derived using exploratory factor analysis with factor loadings extracted using principal component analysis, varimax rotation and eigenvalues greater than 1. Orthogonal rotation was used as it is believed that the variables are not highly correlated with each other. The eigenvalue (greater than 1) indicates that the factor is loaded with a greater number of variables or explains more of the variation compared to when the eigenvalue is less than 1. All 231 NZE participants were included in the factor analysis including those who had a body composition outside of the three main BCP groups. The majority of pattern analysis studies include misreporters in the analysis (Devlin *et al.*, 2012). Although evidence is unclear, it is thought that including misreporters in the analysis of dietary patterns does not significantly bias the data outputs (Bailey *et al.*, 2007). Each of the food items were arranged into one of 56 food groups which were entered into the principal component analysis as daily equivalent frequencies.

The Kaiser-Meyer-Olkin (KMO) measure of sampling adequacy and the Bartlett's test of sphericity were used to assess the adequacy of correlation matrices within the data set. If the KMO value is greater than >5, and if the Bartlett's test of sphericity is statistically significant (<0.05) we can be confident in doing a factor analysis (Dziuban, 1974, Frohlich, 2001). To decide how many factors to derive both scree plots and the factors themselves were analysed to identify which factors explained the data best. Food and beverages that loaded highly (\geq 0.2) were positively associated with the factor and largely contributed to the identified dietary pattern (McNaughton *et al.*, 2011). A negative loading indicated that the item was inversely associated with the factor and thus did not contribute to the dietary pattern. The strength of the association between the food item and the factor is increased as the factor loading increases. Items which cross-loaded with the least loading on a factor were removed from the relevant dietary pattern. Factors were named based food items that loaded highly, the nutritional characteristics and food groupings in each of the identified dietary patterns.

Chronbach's α coefficients were used to assess inter-item reliability. Each factor was assessed individually using the food items that loaded highly because the variance in each of the factors and within all of the food groups is likely to be different (Nunnally, 1994). Food items that indicated that they were skewing the reliability and would increases the scale when taken out were removed from the pattern (for example, factor 4 increased from 0.252 to 0.431 when oil and oil-based dressings, fats and soy milk were removed from the pattern). The factor analysis was re-run without these items to ensure that the removal increased reliability without significantly influencing the dietary patterns (Field, 2009).

Simplified dietary pattern scores were calculated by summing the daily equivalent frequencies for each food item that loaded highly on each of the patterns. The simplified dietary pattern score was developed by Schulze et al (2003) and overcomes the non-comparable limitation of factor analysis by allowing the dietary patterns to relate to other population groups (Schulze *et al.*, 2003). Tertiles for each of the patterns were then determined. Percentage of total number and Chi-square analysis was used to determine associations between the categorical socio-demographic factors and dietary habits across tertiles of each dietary pattern. A one-way ANOVA was performed to compare between the means of nutrient intakes within each dietary pattern, and to analyse the mean age, BMI and BF% within tertiles of each dietary pattern. Post hoc tests of Tukey and Gabriel were used to determine which groups are significantly different from one another. Post hoc tests were used because there is no specific hypothesis that is trying to be proved and it is simply exploring the data to identify trends. Tukey's posthoc test was used because the sample sizes are the same in tertiles of each dietary pattern.

Chapter 4.0 Results

This chapter presents the findings from a subset of data obtained as part of the women's EXPLORE study; in particular the dietary data of the NZE women. The data presented include participant characteristics, followed by a nutrient analysis of both the total study population and within the BCP groups, the top 40 food items consumed, dietary pattern analysis and finally the diet quality of the NZE women.

4.1 Study Population

A total of 1193 NZ women (including NZE, Maori, and Pacific women) were recruited and 798 women screened for participation in the New Zealand Women's EXPLORE study. Of these, 473 were of NZE decent, and 233 of these women met the inclusion criteria and were invited to complete the study protocol. Oversampling (n = 6) was performed to ensure enough participants were recruited to achieve the objective of recruiting 75 women in each of the main BCP groups (NN, NH, HH). However participant body composition parameters changed following ADP analysis, thus resulting in the following BCP groups: 63 NN, 59 NH, 88 HH, 18 NL, and 3 HN. Two women were excluded from the study as they did not complete the FFQ (1 NN, 1 HH). Twenty four women were identified as mis-reporters (19 under-reporters, 5 overreporters) based on energy intakes relative to their energy requirements. Of these, 6 NN, 6 NH, and 7 HH women were identified as under reporters, and 1 NN, 1 NH, and 3 HH women were over reporters. These women were excluded in the nutrient analysis but included in the pattern analysis. The final study population used for data analyses therefore was 207 women in the nutrient analysis and 231 women in the pattern analysis (figure 4.1). Allocation of the women into each of the BCP groups is outlined in table 4.1. In the nutrient analysis, 27% of women were in the NN group. In the pattern analysis 30% were in the NN group, 28% in the NH group and 42% in the HH group.

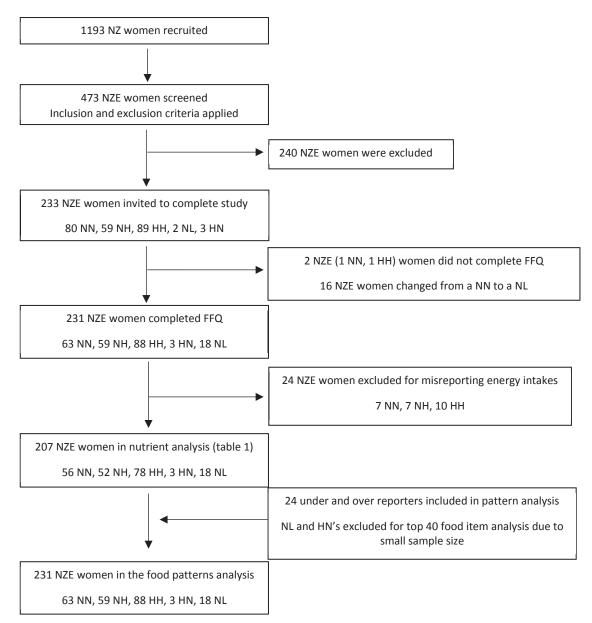


Figure 4.1 Outline of the recruitment process.

Table 4.1 Study population.

BCP groups	Participants included in nutrient analysis (n (%))	Participants included in pattern analysis (n (%))
NN	56 (27)	63 (27)
NH	52 (25)	59 (26)
НН	78 (38)	88 (38)
NL	18 (9)	18 (8)
HN	3 (1)	3 (1)
Total	207	231

NN = Normal BMI (18.5 – 24.99 kg/m²), normal BF% (22 -29.99 kg/m²);

NH = Normal BMI, high BF% (≥ 30%); *NL* = Normal BMI, low BF% (< 22%); HH = High BMI ($\geq 25 \text{ kg/m}^2$), high BF%; HN = High BMI, normal BF%

4.2 Participant Characteristics

The characteristics of the total study population are outlined in table 4.2. The women had an average age of 31.9 years and a median body mass index (BMI) of 24 kg/m². Their average waist to hip ratio, blood pressure and BF% were 0.7, 115/71.7 mmHg, and 32.6%, respectively.

Characteristics	N (%)	NZE women (N = 231)
Age at testing (years)		31.9 (25.2, 39.5) ^b
16-24	54 (23.4)	
25-35	88 (38.1)	
36-45	89 (38.5)	
Height (cm)		167 ± 6.6^{a}
Weight (kg)		70.1 ± 14.2 ^ª
Body Mass Index (BMI) (kg/m ²)		24 (21.7, 27.2) ^b
18.5 – 24.99	140 (60.6)	
25.0 – 29.99	60 (26)	
≥ 30.0	31 (13.4)	
Body fat percentage (%)		32.6 ± 8.1 ^a
< 22	18 (7.8)	
22 – 29.9	66 (28.6)	
≥ 30	147 (63.6)	
Waist Circumference (cm)(<80 ^c)		78.1 ± 11.0 [°]
Hip Circumference (cm)		104.2 ± 10.6^{a}
Waist: Hip ratio (< 0.85 [°])		0.7 ± 0.1^{a}
Systolic BP (mmHg) (120 ^c)		115 ± 9.5 ^a
Diastolic BP (mmHg) (80 ^c)		71.7 ± 7.4^{a}
^a = Mean ± Standard deviation;		

 Table 4.2 Characteristics of all New Zealand European study participants (n = 231).

^a = Mean ± Standard deviation; ^b = Median (25th, 75th percentile);

^c = Standard reference values (World Health Organisation, 2008);

BMI categories are defined as: Normal: 18.5-24.9 kg/m², Overweight: 25.0 - 29.9 kg/m², Obese: ≥ 30.0 kg/m²; Body fat percentage categories are defined as: Under: < 22% fat, Normal: <30% fat, High: ≥30% fat

Participant characteristics compared between the BCP groups are outlined in table 4.3. The NN group had the youngest median age (28.2 years (24, 36.4)) and the HH group the oldest (33.8 years (25.5, 40.5)). Body mass index and BF% increased as age increased when comparing the NN, NH and HH BCP groups respectively (BMI 21.4 kg/m² (19.9, 23.1), BF% 25.8% (23.5, 28.5); BMI 23.0 kg/m² (21.7, 24.4), BF% 33.4% (30.6, 36.4); BMI 28.6 kg/m² (24.4, 33.4), BF% 39.1% (33.3, 45.9)).

	BCP groups (n = 231)						
Characteristics	NN (n = 63)	NH (n = 59)	HH (n = 88)	NL (n = 18)	HN (n = 3)		
Age at testing ^a	28.2 (24, 36.4)	33.1 (25.7, 40.8)	33.8 (25.5, 40.5)	30.7 (23.4, 37.3)	39.0 (26.3, 40.9)		
(years)							
Height ^b (cm)	168.3 ± 6.7	167.6 ± 6.4	166.5 ± 6.9	166.5 ± 6.2	164.2 ± 1.9		
Weight ^b (kg)	60.9 ± 7.0*	64.8 ± 5.6*	82.2 ± 15.0*	59.9 ± 7.0*	73.9 ± 1.8		
BMI ^c (kg/m ²)	21.4 (19.9, 23.1)*	23.0 (21.7, 24.4)	28.6 (24.4, 33.4)*	21.5 (20.1, 23.0)	27.4 (25.7, 29.3)		
BF% ^c (%)	25.8 (23.5, 28.5)*	33.4 (30.6, 36.4)*	39.1 (33.3, 45.9)*	19.7 (18.1, 21.4)*	27.5 (22.9, 33.1)*		
WC ^b (cm)	70.1 ± 4.7*	74.5 ± 3.7*	$88.0 \pm 11.0^*$	69.0 ± 4.5*	80.3 ± 5.9		
HC ^b (cm)	97.5 ± 4.9*	101.4 ± 4.2*	112.4 ± 12.1*	96.7 ± 4.8*	105.6 ± 1.6		
W:H ratio ^b	$0.7 \pm 0.0^{*}$	$0.7 \pm 0.0^{*}$	$0.8 \pm 0.0^{*}$	$0.7 \pm 0.0^{*}$	0.8 ± 0.1		
Systolic BP ^b	112.2 ± 7.9*	114.3 ± 9.7	117.1 ± 9.9*	115.3 ± 11.0	120.0 ± 8.7		
(mmHg)							
Diastolic BP ^b	68.7 ± 6.3*	71.3 ± 7.8	73.9 ± 7.5*	71.3 ± 6.5	75.0 ± 2.6		
(mmHg)							
^{a =} Median (25 th , 2	75 th percentiles);	HC = Hi	p circumference;				
^b = Mean ± Stande	ard deviation (S);	W:H rat	io = Waist to hip rati	0;			
^c = Geometric me	an (lower, upper qua	rtiles); NN = No	rmal BMI (18.5 – 24.	9 kg/m²), normal BF୨	% (22 –29.9%);		
*statistically sign	ificant(ANOVA betw	een BCP groups);	NH = Normal BMI,	high BF% (≥ 30%);			
BMI = Body mass	,		5 1	HH = High BMI (≥ 25.0), high BF%;			
BF% = Body fat p	ercentage;		NL = Normal BMI, low BF% (< 22%);				

Table 4.3 Characteristics of participants compared between body composition profile (BCP)	
groups.	

BMI categories are defined as: Normal: 18.5-24.9 kg/m², Overweight: 25.0 - 29.9 kg/m², Obese: ≥ 30.0 kg/m²; Body fat percentage categories are defined as: Under: < 22% fat, Normal: <30% fat, High: $\geq 30\%$ fat

Participant waist circumference (WC), hip circumference (HC), systolic and diastolic blood pressures increased as BF% and BMI increased between the groups (NN 112.2 mmHg \pm 7.9 / 68.7 \pm 6.3 mmHg; NH 114.3 mmHg \pm 9.7 / 71.3 \pm 7.8; HH 117.1 mmHg \pm 9.9 / 73.9 \pm 7.5 mmHg).

HN = High BMI, normal BF%;

n = Number;

4.3 Dietary Analysis

WC = *Waist circumference;*

BP = *Blood pressure*;

4.3.1 Total NZE population

The average daily energy intake for the NZE women in this study was 9164 kJ (table 4.4). The women were meeting both the estimated average requirements (EARs) and the recommended daily intakes (RDIs) for all nutrients analysed, apart from Iron where only the EAR was met. Dietary fibre, vitamin E, and potassium intakes also met the adequate intake (AI) levels. Vitamin D intakes were below the AI. Sodium was consumed in excess of the upper limit.

Table 4.4 Mean daily dietary intakes from the NZWFFQ (n = 207) vs. New Zealand National recommendations and percentage of NZE women below the recommendations. (National Health Medical Research Council, 2005).

Nutritional analysis	Mean ± SD, Median (25 th , 75 th percentile) ^a	EAR ^b	Percent (%) below EAR	NRVs ^{cdef}	Proportion (%) under and over NRV	Cut off 77% of RDI	Percent (%) below cut off
Energy (kJ)	9164.5 ± 2278.8	na	na	7100-8700 ^h	23 U, 50 O ^h	-	-
Protein (g)	96.3 ± 25.7	37	0	46 ^c	0 ^c	-	-
(% of TE)	18.0 ± 3.3 ^e			15-25% of TE ^e	13 U, 3 O ^e		
Total fat (g)	86.7 ± 28.0	na	na			-	-
(% of TE)	34.8 ± 6.3 ^e			20-35% of TE ^e	0 U, 39 O ^e		
Saturated fat (g)	34.8 ± 13.9	na	na			-	-
(% of TE)	13.9 ± 3.5 ^e			<12% of TE ^e	66 O		
PUFA (g)	12.8 ± 4.6	na	na			-	-
(% of TE)	5.2 ± 4.0^{f}			6-10% of TE ^f	69 U, 2.4 O		
MUFA (g)	29.4 ± 9.5	na	na			-	-
(% of TE)	16.1 ± 3.1^{f}			10-20% of TE ^f	29 U, 0.5 O		
Cholesterol (mg)	289.3 ± 142.5	na	na	na	na	-	-
Carbohydrate (g)	233.9 ± 74.6	na	na			-	-
(% of TE)	41.9 ± 7.0 ^e			45-65% of TE ^e	64 U, 0 O ^e		
Sugars (g)	124.2 ± 46.6	na	na			-	-
(% of TE)	22.6 ^f			<15% of TE ^f	15 U, 84.5 O ^f		
Dietary fibre (g)	30.2 ± 9.6	na	na	25 ^d	31 ^d	-	-
Thiamin (mg)	1.7 ± 0.8	0.9	7.7	1.1 ^c	23 ^c	0.8	4
Riboflavin (mg)	2.6 ± 1.0	0.9	0.5	1.1 ^c	1 ^c	0.8	0
Niacin (mg)	22.4 ± 8.6	11	3	14 ^c	9 ^c	11	3
Vitamin C (mg)	158.7 ± 83.7	30	0.5	45 [°]	1.4 ^c	35	0.5
Vitamin D (µg)	4.6 ± 2.3	na	na	5 ^d	64 ^d	-	-
Vitamin E (mg)	13.5 ± 4.9	na	na	7 ^d	6 ^d	-	-
Vitamin B6 (mg)	2.5 ± 1.7	1.1	2.4	1.3 ^c	4 ^c	1.0	1
Vitamin B12 (µg)	4.8 ± 2.5	2.0	5	2.4 ^c	9 ^c	1.8	2
Total-folate (µg)	439.9 ± 155.6	320	25	400 ^c	46.3 ^c	308	23
Vitamin A (µg)	1564.7 ± 724.1	500	1	700 ^c	4.3 ^c	539	1.4
Sodium (mg)	2522.8 ± 861.0	na	na	2300 ^g	49 ^e	1771	-
Potassium (mg)	3815.5 ± 1080.1	na	na	2800 ^d	12.5 ^d	-	-
Magnesium (mg)	405.3 ± 112.3	255-	8.2	310-320 ^c	22.2 ^c	238	5.3
5 (0,	-	265					
Calcium (mg)	1233.1 ± 462.5	840	17	1000 ^c	38 ^c	770	11.5
Phosphorus (mg)	1742.8 ± 468.2	580	0	1000 ^c	4.3 ^c	770	0
Iron (mg)	12.6 ± 3.5	8	6	18 ^c	94 ^c	14	67
Zinc (mg)	11.9 ± 3.3	6.5	1.4	8 ^c	10 ^c	6	1.4
Selenium (µg)	83.4 ± 51.1	50	23	60 ^c	33 ^c	46	17
Alcohol (g)	4.3 (1.4, 10.2) ^a	na	na	na	na		_;

^a = Median (25th, 75th percentile);

^bNRV-EAR: Estimated Average Requirement;

^c NRV-RDI: Recommended Dietary Intake;

^d NRV-AI: Adequate Intake;

^eNRV-AMDR: Acceptable Macronutrient Distribution Range; SD = Standard Deviation;

^fNRV-SDT: Suggested Dietary Targets;

^gNRV-UL: Upper Limit;

^hNRV – EER: Estimated Energy Requirement;

All NRVs used were for women 19-30 and 31-50 years old (National Health Medical Research Council, 2005);

NRV = Nutrient Reference Values for New Zealand; PUFA = Polyunsaturated fat; O = Over;

MUFA = Monounsaturated fat; na =Not Available;

U=Under

TE = Total Energy;

Despite the nutrient recommendations being met by the population group, many women consumed less than the NRVs and / or the EARs for many nutrients. Dietary fibre and vitamin D were consumed less than the NRVs by 31% and 64% of women, respectively. Similarly for calcium, iron and selenium, 38%, 94% and 33% of participants consumed less than the NRV for each of these nutrients, respectively. For folate, 25% of participants consumed less than the EAR and 46.5% consumed less than the NRV.

All macronutrients except carbohydrate were within the recommended macronutrient distribution range (AMDR) with 41.9%, 18%, and 34.8% of energy intake coming from carbohydrate, protein and fat, respectively. Thirteen percent of participants consumed below the NRV for protein. Sixty four percent of participants consumed less than the recommended energy intake per day as carbohydrate. Nevertheless, 84.5% of participants consumed over the NRV for sugar per day. A total of 39% of participants consumed over the NRV for fat and 66% consumed over the NRV for saturated fat per day. Polyunsaturated fatty acid intakes were low with 69% of participants consuming less than the NRV per day.

4.3.2 Body composition profile groups

Energy and macronutrients

The energy and macronutrient intakes for each of the body composition profile (BCP) groups are outlined in figure 4.2 and table 4.6. The HH profile group had the largest energy intake per day (9296.2 kJ/d) compared to the HN profile group (7776.8 kJ/d). The NL group consumed the most protein (100.7 g/d), and the NN group consumed the least (93.4 g/d). The NN group consumed the most carbohydrate (242.2 g/d) and the most sugar per day (130.2 g/d), but only consumed 43.2% of energy from carbohydrate. The total and saturated fat content of the diets increased as BF% and BMI increased. The HH group also consumed the most total (89.4 g/d), saturated (36.5 g/d) and monounsaturated fat (30.4 g/d) compared to the other groups. The NL group had the highest intake of polyunsaturated fat (13.5 g/d).

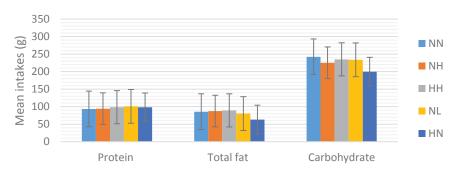


Figure 4.2 Mean (SD) macronutrient intakes compared between the body composition profile groups.

Table 4.5 Nutrient analysis comparison between body	analysis cor	nparison be	tween body composit	ion profile (BCP) group	s and with the estimat	composition profile (BCP) groups and with the estimated average nutrient requirements (EAR)	quirements (EAR).
	Recommendations	ndations			BCP groups*		
Nutrients	EAR	RDI	NN	HN	НН	NL	NH
Energy (kJ) ^c	7100-8700	na	9198.3 ± 2141.0	9057.8 ± 2114.9	9296.2 ± 2505.5	9028.8 ± 2206.2	7776.8 ± 2763.3
Protein (g) ^c	37	46	93.4 ± 25.4	94.3 ± 24.7	98.5 ± 26.5	100.7 ± 27.9	98.2 ± 20.5
Total fat (g) ^c	na	na	85.6 ± 24.3	87.5 ± 26.2	89.4 ± 31.5	80.4 ± 27.4	63.1 ± 24.3
Saturated fat (g) ^c	na	na	34.2 ± 13.0	35.0 ± 13.4	36.5 ± 15.4	31.2 ± 11.5	22.2 ± 9.5
PUFA (g) ^a	na	na	13.0 (9.6, 14.9)	12.2 (10.9, 13.5)	11.9 (10.9, 12.9)	13.5 (10.0, 15.1)	10.5 (5.1, 21.9)
MUFA (g) ^c	na	na	28.8±8.2	29.9 ± 8.8	30.4 ± 10.8	27.0 ± 10.0	22.2 ± 9.1
Cholesterol (mg) ^a	na	na	255.7 (182.7, 323.0)	247.0 (215.0, 283.7)	281.5 (256.3, 309.2)	250.7 (197.0, 344.3)	398.7 (160.3, 991.5)
Carbohydrate (g) ^c	na	na	242.2 ± 72.2	225.4 ± 62.1	235.0 ± 84.0	233.7 ± 71.5	199.6 ± 100.5
Sugars (g) ^c	na	na	130.2 ± 49.9	114.3 ± 33.6	126.2 ± 52.8	128.3 ± 34.6	108.7 ± 69.6
Dietary fibre (g) ^c	вп	25 (AI)	31.4 ± 10.6	28.8 ± 8.1	29.5 ± 9.9	32.9 ± 8.8	31.9 ± 15.0
Thiamin (mg) ^a	6.0	1.1	1.4 (1.2, 1.8)	1.6 (1.4, 1.8)	1.5 (1.4, 1.7)	1.65 (1.3, 1.9)	1.8 (1.1, 3.0)
Riboflavin (mg) ^c	6.0	1.1	2.6±1.0	2.7 ± 1.1	2.5 ± 1.1	2.7 ± 0.9	2.9 ± 0.7
Niacin (mg) ^c	11	14	21.4 ± 7.5	21.7 ± 9.1	23.6 ± 9.5	23.0 ± 6.3	20.4 ± 4.2
Vitamin C (mg) ^b	30	45	147.5 (103.1, 190.0)	144.0 (94.3, 160.0)	131.6 (104.9, 192.7)	175.4 (132.7, 210.9)	269.1 (49.7, 273.8)
Vitamin D (µg) ^c	na	5 (AI)	4.5 ± 2.2	4.6 ± 2.9	4.6 ± 2.2	4.6±1.8	4.4 ± 0.6
Vitamin E (mg) ^c	na	7 (AI)	13.4 ± 4.6	13.4 ± 4.8	13.4 ± 5.2	14.4 ± 5.2	13.6 ± 2.7
Vitamin B6 (mg) ^c	1.1	1.3	2.3 ± 0.7	2.5 ± 2.3	2.7 ± 1.8	2.6±0.8	2.6 ± 1.4
Vitamin B12 (µg) ^c	2.0	2.4	4.5±2.1	4.6 ± 2.3	5.1 ± 3.0	5.0±2.3	4.6 ± 1.3
Total-folate (μg) ^c	320	400	462.6 ± 141.7	453.4 ± 155.7	408.9 ± 156.7	467.4 ± 186.4	421.1 ± 136.4
Vitamin A (µg) ^c	500	200	1587.0 ± 767.7	1381.5 ± 496.3	1574.5 ± 756.4	1952.5 ± 903.9	1740.5 ± 507.3
Sodium (mg) ^c	na	2300 (NL)	2457.2 ± 713.3	2468.6 ± 771.6	2661.9 ± 1028.1	2358.2 ± 754	2058.8 ± 375.6
Potassium (mg) ^c	na	2800 (AI)	3823.3 ± 1104.8	3695.0 ± 911.6	3773.1 ± 1146.2	4261.1 ± 1027.5	4187.3 ± 1843.4
Magnesium (mg) ^c	255-265	310-320	412.1 ± 120.9	399.1 ± 95.6	395.2 ± 112.5	444.8 ± 122.8	408.2 ± 166.6
Calcium (mg) ^b	840	1000	1197.7 (979.0, 1494.7)	1257.7 (963.2, 1375.4)	1159.5 (855.4, 1398.7)	1409.4 (132.7, 1620.8)	1386.3 (1142.3, 1768.1)
Phosphorus (mg) ^c	580	1000	1749.5 ± 490.2	1735.0 ± 438.7	1703.9 ± 461.8	1884.7 ± 517.4	1911.0 ± 546.1
Iron (mg) ^a	8	18	11.9 (10.0, 15.5)	11.9 (10.9, 12.9)	12.2 (11.5, 12.9)	13.1 (10.9, 16.1)	12.3 (6.9, 21.9)
Zinc (mg) ^c	6.5	8	11.7 ± 3.1	11.8 ± 3.0	12.0 ± 3.5	12.8 ± 4.1	11.7 ± 3.4
Selenium (µg) ^a	50	60	70.7 (48.1, 92.4)	73.9 (65.1, 83.9)	73.9 (66.1, 82.5)	102.0 (46.1, 125.6)	90.2 (25.3, 321.4)
Alcohol (g) ^b	na	na	4.1 (1.2, 9.1)	5.65 (0.97, 13.0)	3.90 (0.42, 8.38)	4.4 (3.0, 7.1)	8.84 (1.82, 11.2)
^a = Geometric Mean (lower, upper quartiles);	wer, upper quo	ırtiles);	AI = Adequate intake	ke		PUFA = Polyunsaturated fatty acids	ids
^b = Median (lower, upper percentiles);	er percentiles);		NN = Normal BMI	NN = Normal BMI (18.5 – 24.9 kg/m ²), normal BF% (22 – 29.9%);		MUFA = Monounsaturated fatty acids	acids
^c = Mean ± Standard deviation;	viation;		NH = Normal BMI, high BF% (≥ 30%);	high BF% (≥ 30%);	∧ = *	* = Values not significant between BCP groups	en BCP groups
EAR = Estimated average nutrient requirement;	ige nutrient req	uirement;	HH = High BMI (≥ 25.0), high BF%;	25.0), high BF%;	= 7/1	UL = Upper limit;	
RDI = Recommended daily intake,	laily intake;		NL = Normal BMI, low BF% (< 22%);	low BF% (< 22%);	HN =	HN = High BMI, normal BF%	

hte (FAR) . 4 ć 70+0 1000 with the ł 'ofile (RCD) citic - Poor 5 10 0 analysis Table 4 5 Nutrient

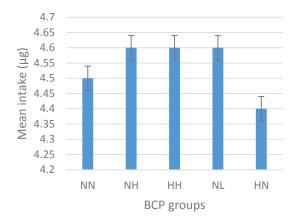
Micronutrients

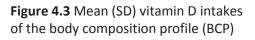
The micronutrient intake for each of the BCP groups is outlined in table 4.6. All of the EAR, RDI, and Al's were met for all of the nutrients analysed except for vitamin D and iron. Vitamin D intakes did not meet the AI in all BCP groups (figure 4.3). Estimated average requirements for iron was met, however the RDI was not (figure 4.3). Excess sodium was also present in all apart from the NL and HN group. Although meeting the nutrient recommendations, the HH profile group had the lowest intakes of riboflavin, vitamin C, folate, magnesium, calcium, and phosphorus compared to the other BCP groups (figure 4.4). They however had the highest intake of niacin, vitamin B6, vitamin B12, and zinc. These findings indicate that the HH group may consume a diet of poor nutrient density with less consumption of fruits, vegetables and dairy products and a higher consumption of meat compared to the other BCP groups.

Sodium intake was consumed in excess in all BCP groups apart from the HN group. The HH BCP group had the largest consumption per day (2661.9 mg/d) (figure 4.7).

The NH group had the lowest intakes of dietary fibre, vitamin E, vitamin A, potassium, and iron. They also had the highest intake of polyunsaturated fat along with the NN group. The NN group also had the highest intake of iron (figure 4.6), and the lowest intakes of thiamin, vitamin B6 and selenium.

The NL group had the highest intakes of thiamine, riboflavin, vitamin D, vitamin E, magnesium, calcium, iron and zinc indicating a nutrient dense diet (figures 4.3, 4.5, 4.6, 4.11). They also had the lowest intakes of cholesterol, and vitamin B12 (figure 4.4). The HN group had the highest intakes of cholesterol, dietary fibre, riboflavin, vitamin C, vitamin A, potassium, phosphorus, iron, and selenium. They had the lowest intake of niacin, vitamin D, sodium and zinc (figures 4.6, 4.8, 4.9, 4.10).





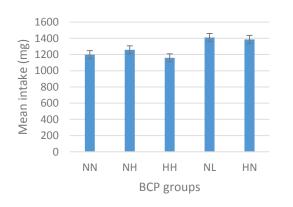


Figure 4.5 Mean (SD) calcium intakes of the body composition profile (BCP) groups.

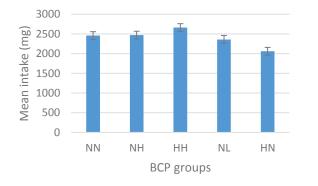


Figure 4.7 Mean (SD) sodium intakes compared between the body composition profile (BCP) groups.

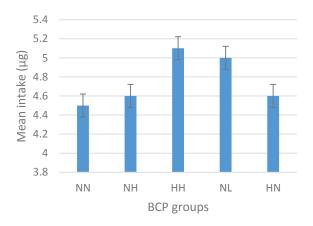


Figure 4.4 Mean (SD) vitamin B12 intakes of the body composition profile (BCP) groups.

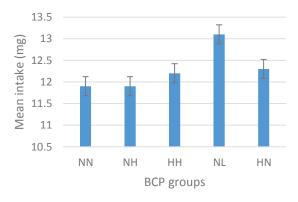


Figure 4.6 Mean (SD) iron intakes of the body composition profile (BCP) groups.

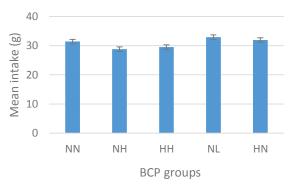
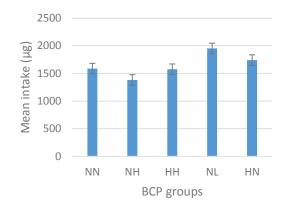


Figure 4.8 Mean (SD) intake of dietary fibre compared between the body composition profile (BCP) groups.



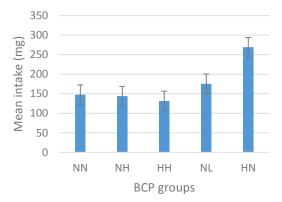


Figure 4.9 Mean (SD) vitamin A intakes of the body composition profile (BCP)

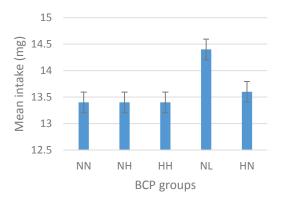


Figure 4.11 Mean (SD) vitamin E intakes of the body composition profile (BCP)

Figure 4.10 Mean (SD) vitamin C intakes of the body composition profile (BCP)

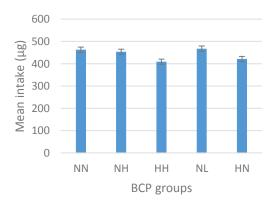


Figure 4.12 Mean (SD) total folate intakes of the body composition profile

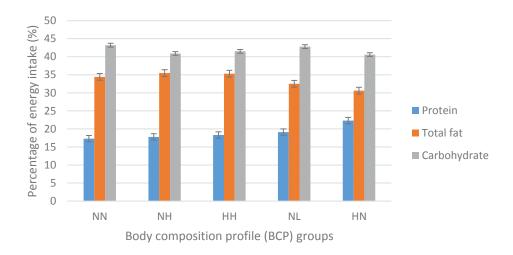
4.3.3 Macronutrient distribution

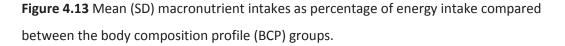
The average macronutrient distribution ranges for each of the BCP groups is outlined in table 4.6.

 Table 4.6 Mean macronutrient percentage intakes (%) compared between body composition

Macronutrient AMDR NN M				HH	NL	HN	
kJ from protein (%)*	15-25	17.3 ± 2.7	17.8 ± 3.2	18.3 ± 3.4	19.1 ± 3.6	22.3 ±3.7	
kJ from fat (%)*	20-35	34.4 ± 6.0	35.5 ± 5.6	35.3 ± 6.6	32.5 ± 6.9	30.6 ±7.8	
kJ from saturated fat (%)*	<12	13.7 ± 3.5	14.1 ± 3.2	14.3 ± 3.8	12.6 ± 3.2	10.7 ±3.1	
kJ from carbohydrate (%)*	45-65	43.2 ± 6.6	40.9 ± 5.6	41.5 ± 7.7	42.8 ± 8.3	40.6 ±9.9	
Values reported are mean ± st	andard dev	iation;	NN = Normal BMI (18.5 – 24.9 kg/m²), normal BF%				
kJ = Kilojoule;			(22 – 29.9	9%);			
AMDR = Acceptable macronut	NH = Nori	mal BMI, high	BF% (≥ 30%);				
* = Significantly different (ANC	HH = High	n BMI (≥ 25.0),	high BF%;				
			NL = Normal BMI, low BF% (< 22%);				
			HN = High BMI, normal BF%				
					1 77		

profile (BCP) groups.





None of the BCP groups met all of the acceptable macronutrient daily recommendations (AMDRs). All of the groups reported low carbohydrate intakes below the level of 45% of total energy intake. The NN group reported the highest intake of 43.2% with the NH and HH groups reporting intakes of 40.9% and 41.5%, respectively. Protein intakes were within the recommendations (NN 17.3%, NH 17.8%, and HH 18.3% of energy intake). Total fat intakes varied with only the NN and HN groups meeting the AMDR for total fat (34.4% and 30.6%, respectively). The other groups all consumed over the 35% of total energy cut off point for total fat intake. The HN group was the only group who consumed less than the recommended 12% of total energy intake as saturated fat per day (10.7%). The other groups consumed over the recommendation where 13.7%, 14.1%, 14.3%, and 12.6% of total energy intake was consumed as saturated fat in the NN, NH, HH and NL groups, respectively.

4.4 Top 40 Food Items

Total population

The 40 most frequently consumed foods reported in the NZWFFQ within the total study population are outlined in table 4.7. Twenty one of the 40 food items were from nutritious food groups, with 13 from the fruit and vegetable group, three from breads and cereals, three from low fat dairy products and two from eggs and nuts. No meat or poultry food items were present in the top 40 most frequently consumed foods.

Ranking	Food item	Frequency	Mean daily intake	# respondents with
		(per day)*	(g per person/d)	daily consumption
1	Water	783.2	850	224
2	Oil, composite used in cooking	341.1	3.3	51
3	Coffee, brewed	173.8	189.4	73
4	Tea, black, infused, weak	170.4	190.1	71
5	Butter	168.4	3.1	41
6	Multigrain bread	159.6	28.6	63
7	Herbal tea	156.7	175.3	73
8	Lettuce	106.9	18.2	45
9	Oil as a dressing/sauce	105.1	2.5	59
10	White sugar	95.5	1.8	23
11	Lite milk added to drinks	85.4	32.2	55
12	Trim milk added to drinks	85.4	19.2	36
13	Apple	84.7	58	42
14	Onion	82.3	21	35
15	Coffee, espresso (30 mL)	79.5	12.2	38
16	Carrot	79.3	36.9	31
17	Yoghurt	78.4	54	35
18	Banana	77.8	57.6	41
19	Margarine as a dressing/sauce	77.4	1.7	36
20	Egg, chicken	76	18.9	16
21	Tomatoes	70.2	55.3	23
22	Cream crackers	69	6.7	23
23	Broccoli	68.3	33.9	17
24	Almonds	67.9	4.3	28
25	Margarine, Mono, 55% fat	65.9	1.3	26
26	Lollies	64.4	4.6	21
27	Margarine, Mono, 70% fat	61.5	1.3	22
28	Chocolate bar	60.2	12.7	25
29	Courgette	60	21.5	18
30	Cheese - edam	57.9	13.1	24
31	Capsicum	56.1	11.7	14
32	Peanut butter	55.2	3.7	19
33	Orange	54.3	34.8	16
34	Wholemeal bread	53.4	7.1	19
35	Tomato sauce	50.6	4	14
36	Diet soft drink	40.0	46.4	17
37	Plain biscuit	46.9	4.4	14
38	Avocado	46.2	11.5	10
39	Spinach	44.8	22.9	13
40	Potato	43.3	18	5

 Table 4.7 Top 40 foods consumed for the total population.

*Frequency per day indicates the number of times the food item is eaten per day by the total (n = 231) NZE women

= Beverages

= Nutritional food items

= Discretionary food items

Body composition profile groups

The top 40 most frequently consumed foods within the three main BCP groups are outlined in table 4.8. The food items as previously mentioned for the NL and the HN BCP groups are not presented due to low sample sizes. Thirty of the 40 food items were present in all three BCP groups as being frequently consumed. Water and oil were the first and second items in all groups, similar to that of the total group. The HH group consumed water and oil for cooking more frequently than the other groups, although they were both ranked first and second, respectively in the BCP groups. They also used oil as a dressing more frequently than the other groups with this making the third top food in the HH group compared to the 8th and 9th food item in the NN and NH groups, respectively.

Fruit and vegetables were present in all BCP groups. The NN BCP group had 5 fruit and 6 vegetables in their top 40 foods, whereas the HH had 2 fruit and 8 vegetables and the NH 3 fruit and 7 vegetables. The HH BCP group was the only group to have an animal protein source in their top 40 food items, however all groups contained eggs as an alternative protein source. Additionally lite or trim milk and yoghurt were also present in all BCP groups. The highest ranking item from the breads and cereals food group was multigrain bread. This was ranked as the third most commonly consumed food in the NH group with this group having the highest frequency of consumption per day, consuming the most per day and having the most individuals consuming it per day.

All BCP groups contained discretionary items such as lollies and chocolate bars. Both these items ranked the highest in the HH BCP group, where they had the highest frequency of consumption, consumed the most per day and had the most individuals consuming on a daily basis. The HH group were also the only group to have diet soft drink in their top 40 food items.

					BCP groups				
		NN (n = 63)			NH (n = 59)			HH (n = 88)	
Food item	Freq/d	Mean intake	# subjects	Freq/d	Mean intake	# subjects	Freq/d	Mean intake	# subjects
	(ranking)	(g/b/d)	with DI	(ranking)	(b/d)	with DI	(ranking)	(b/d)	with DI
Water (250 mL)	211.2 (1)	842.4	59	196.4 (1)	828.2	58	297.1 (1)	844.2	87
Oil used in cooking (5 mL)	115.8 (2)	3.6	16	87.8 (2)	3.6	15	287.8 (2)	3.3	20
Butter (1 tsp)	62.3 (3)	3.9	21	34.3 (8)	3.1	10	66.1 (6)	3	14
Coffee, brewed (1 cup)	54.7 (4)	196.9	22	39.6 (6)	175.3	16	67.9 (5)	203.3	31
Multigrain bread (1 slice)	51.0 (5)	33.7	19	53.5 (3)	35.2	25	44.1 (10)	22.1	15
White sugar (1 tsp)	15.5 (30)	3.3	9	25.3 (10)	1.8	10	ı	ı	ı
Herbal tea (1 cup)	38.2 (6)	153.5	19	51.4 (4)	224.6	24	50.1 (8)	151.3	22
Black tea (1 cup)	33.5 (7)	141.2	15	51.1 (5)	223.7	22	71.5 (4)	207.6	28
Oil as a dressing/sauce (1 tsp)	32.3 (8)	2.8	20	31.9 (9)	2.9	20	193.1 (3)	2.2	17
Lettuce (0.5 cup)	27.0 (10)	16.8	12	21.2 (14)	16.2	8	49.2 (9)	20.3	23
Lite milk added to drinks (50 mL)	ı	ı	ı	39.01 (7)	36	15	63.7 (7)	39	24
Carrot (0.5 cup)	25.9 (12)	40.5	12	15.9 (27)	28.5	4	26.7 (22)	35.6	6
Coffee, espresso (30 mL)	26.2 (11)	14.8	12	21.8 (13)	13.5	11	22.5 (32)	8.1	6
Apple (1 fruit)	23.0 (15)	59.7	14	16.5 (26)	46.1	9	33.6 (11)	58.3	14
Banana (1 fruit)	25.1 (13)	60.6	17	16.5 (25)	53.5	7	28.8 (18)	53.8	12
Margarine as a dressing/sauce (1 tsp)	23.1 (14)	1.9	13	20.7 (15)	1.9	6	25.2 (26)	1.4	11
Onion (0.25 cup)	19.3 (19)	16.8	80	19.1 (18)	18.9	7	30.7 (15)	22.3	7
Yoghurt (0.5 cup)	22.6 (16)	58.0	13	17.9 (20)	47.5	7	29.5 (16)	52.7	11
Trim milk added to drinks (50 mL)	17.8 (24)	15.0	6	25 (11)	21.9	10	33.1 (12)	19.4	12

Table 4.8 Top 40 foods compared between the three main body composition profile (BCP) groups.

					BCP groups				
		NN (n = 63)			NH (n = 59)			HH (n = 88)	
Food items	Freq/d	Mean intake	# subjects	Freq/d	Mean intake	# subjects	Freq/d	Mean intake	# subjects
	(ranking)	(b/d/g)	with DI	(ranking)	(b/d/g)	with DI	(ranking)	(g/p/d)	with DI
Broccoli (0.5 cup)	19.1 (20)	35.5	4	15.7 (28)	27.1	4	25.5 (25)	34.9	9
Eggs (1 medium)	17.3 (27)	15.6	2	17.1 (23)	16.5	4	31.9 (13)	20.4	7
Margarine, Mono, 55% fat (5 mL)	22.0 (17)	1.6	10	13.7 (33)	1.1	ŝ	27 (21)	1.4	12
Cream crackers (2 crackers)	17.4 (26)	6.6	9	18.9 (19)	7.3	ß	24.4 (27)	9	∞
Tomato (1 medium)	18.0 (23)	51.0	5	14.5 (32)	52.9	2	30.9 (14)	57.5	13
Lollies (2 items)	19.5 (18)	4.7	9	12.6 (35)	3.4	4	28.8 (19)	5.3	6
Peanut butter (1 tbsp)	18.4 (22)	4.2	8	ı	I	ı	20.9 (34)	3.6	9
Almonds (10 nuts)	17.5 (25)	3.8	6	19.6 (17)	4.6	8	25.6 (23)	4.5	6
Chocolate bar (1 small)	16.2 (28)	11.0	8	11.4 (39)	9.3	ŝ	29.3 (17)	15.1	10
Margarine, Mono, 70% fat (5 mL)	19.0 (21)	1.4	9	ı	ı	ı	28 (20)	1.5	6
Orange (1 fruit)	15.6 (29)	34.8	5	17.2 (22)	39.9	7			
Courgette (0.5 cup)	14.3 (34)	19.9	3	15.7 (29)	20.9	5	23.1 (29)	22.1	8
Cheese - edam (3 tbsp)	14.1 (35)	12.5	7	20.7 (16)	17.1	ø	17.7 (39)	11	9
Honey (1 tsp)	13.2 (38)	1.6	ß			ı	ı	1	
Rolled oats (0.5 cup)	13.8 (36)	12.5	6	ı	ı	ı	ı	ı	ı
Wholemeal bread (1 slice)	14.9 (32)	7.1	4	16.6 (24)	8.9	7	19.4 (35)	9.9	7
Grapes (9 fruits)	14.6 (33)	10.6	ю	ı	ı	ı	ı		ı
Natural muesli (0.5 cup)	15.4 (31)	15.2	8			ı	ı		
Fruit juice	13.4 (37)	62.5	ъ		ı	ı	ı		
Potato	12.8 (39)	19.1	2	I	ı	ı	I	ı	ı

					BCP groups				
		NN (n = 63)			NH (n = 59)			HH (n = 88)	
Food items	Freq/d	Mean intake	# subjects	Freq/d	Mean intake	# subjects	Freq/d	Mean intake	# subjects
	(ranking)	(b/d)	with DI	(ranking)	(b/d)	with DI	(ranking)	(b/d)	with DI
Strawberries	12.6 (40)	18.2	5	1	1	1	I	1	
Spinach (0.5 cup)	ı	ı	I	ı	ı	I	18.5 (37)	22.1	7
Seeds (sunflower)	ı	ı	I	11.6 (37)	1.9	ю	I	ı	1
Toasted sandwich (1 sandwich)	ı	ı	ı	22.9 (12)	8.1	0	ı	ı	1
Lite milk added to cereal (125 mL)	ı		ı	17.8 (21)	44.9	12	ı		
Plain biscuit (2 biscuits)	ı	ı	ı	15.5 (30)	9	7	17.5 (40)	4.1	4
Capsicum (0.5 of a whole)	ı	ı	ı	13.5 (34)	10.5	2	25.5 (24)	13.2	8
Tomato sauce (1 tbsp)	ı	ı	ı	ı	ı	ı	24.3 (28)	5.3	8
Cheese – cheddar (3 tbsp)	ı	ı	ı	ı	ı	ı	23.0 (30)	13.2	7
Avocado (0.25 of a fruit)	ı	ı	I	14.6 (31)	13.8	4	I	ı	1
Chocolate biscuit (2 biscuits)	ı	ı	I	11.9 (36)	4.8	ŝ	I	ı	1
Margarine, Poly, 60% fat (5 mL)	ı	ı	I	11.5 (38)	0.9	Ъ	I	ı	1
Muesli bar (1 bar)	ı	ı	ı	11.2 (40)	7.2	4	ı	ı	1
Diet soft drink (1 cup)	ı	ı	I	ı	ı	ı	22.9 (31)	70.8	10
Frozen mixed vegetables (0.5 cup)	ı	ı	I	ı	ı	I	21.6 (33)	25.8	4
Marmite (1 tsp)	ı	ı	I	ı	ı	ı	19.1 (36)	1.5	ß
Chicken breast (125g)	ı			ı			18.4 (38)	31.3	4
= Beverages = Nutritional food items = Discretionary food items	< < T	NN = Normal BMI (18.5 – 24.9 kg/m ² , NH = Normal BMI, high BF% (≥ 30%); HH = High BMI (≥ 25.0), high BF%	(18.5 – 24.9 kg high BF% (≥ 3 25.0), high BF%	3/m²), normal 0%); 6	NN = Normal BMI (18.5 – 24.9 kg/m ²), normal BF% (22 – 29.9%); NH = Normal BMI, high BF% (≥ 30%); HH = High BMI (≥ 25.0), high BF%		Freq/d = Frequency per day; g/p/d = grams per person per day; DI = Daily intake	er day; rson per day;	n = number

Bold items = new food items not in the top 40 foods identified for the total NZE population

4.5 Dietary Pattern Analysis

Dietary patterns were identified using the FFQ data obtained from the total NZE population (n=231). Principal components factor analysis was used to analyse the data which identified four dietary patterns. Food items that loaded on each of the factors are outlined in table 4.9.

The four patterns that were identified were named as follows:

- Snacking pattern (pattern 1) This pattern was characterised by high loadings on 12 items which are commonly consumed as snacks, especially peanut butter, margarine, sweet spreads, cakes and biscuits and sweet and savoury snack foods e.g. chocolate and potato chips. It was also characterised by negative loadings on green vegetables, red meats and egg and egg dishes.
- Energy-dense meat pattern (pattern 2) This pattern was characterised by high loadings on 15 items which were either high in energy or meat products such as red, white and processed meats, pudding, and crumbed and deep fried food items e.g. crumbed and battered fish. It was also characterised by negative loadings on high calcium milk, yoghurt, soy products and water.
- Fruit and vegetable pattern (pattern 3) This pattern was characterised by high loadings on 7 items all of which were fruits and vegetables. It was also characterised by negative loadings on coffee.
- Healthy pattern (pattern 4) This pattern was characterised by high loadings on 9 items including water, whole grains, and meat alternatives such as fish and seafood, legumes, and nuts and seeds. It was also characterised by negative loadings on white and brown breads, sweetened cereals and fruit juice.

		Dietary p	atterns	
-	1	2	3	4
Full fat milk		-		
Low fat milk	_	_	_	
High calcium milk	.284	238	-	
Sweetened milk products	.204	238 .297	-	
	-	214	-	
Yoghurt High fat choose	-	214 .361	-	
High fat cheese Low fat cheese	.303	.501	-	
	.303	-	424	
Apple, Banana, Orange	-	-	.424	
Other fruit	-	-	.568	
Tomatoes	-	-	.476	
Dark-yellow vegetables	-	-	.628	
Green vegetables	249	-	.588	
Other non-starchy vegetables	-	-	.673	
Potatoes	-	.402	-	
Starchy vegetables	-	-	.593	
White breads	-	.480	-	
Discretionary breads		.422	-	
Crackers	.280	-	-	
Brown breads	.384	-	-	22
Refined grains	-	.390	-	
Wholegrain	-	-	-	.583
Non sweetened cereals	-	-	-	
Sweetened cereals	-	-	-	29
Red meats	231	.609	-	
White meats	-	.495	-	
Processed meats	-	.577	-	
Fish and seafood	-	-	-	.50
Egg and egg dishes	233	-	-	.29
Legumes	-	-	-	.54
Soy products	-	248	-	.39
Peanut butter and peanuts	.549	-	-	
Nuts and seeds	-	-	-	.41
Coconut fats	-	-	-	.46
Margarine	.538	-	-	
Creamy dressings	-	.297	-	
Sauces	-	.396	-	
Sweet spreads	.572	-	-	
Savoury spreads	.336	-	-	
Cakes and biscuits	.674	-	-	
Pudding	-	.565	-	
Sweet snack foods	.477	-	-	
Savoury snack foods	.531	-	-	
Crumbed and deep fried	-	.544	-	
Fast food	-	-	-	
Fruit juice	-	-	-	
Fruit and other drinks	-	.328	_	
Tea	.206	.320	_	
Coffee	.200	-	245	
Beer	-	-	245	.23
Wine	-	-	-	.234
Water	-	- 279	-	.43
Water Other alcoholic beverages	-	279 .312	-	.43

Table 4.9 Factor loading matrix for the four dietary patterns identified in NZE women (n=231).

Dietary patterns based on analysis of FFQ data from 231 NZE women; Pattern 4: 'Healthy' pattern Factors identified based on factor loadings > 0.2;

Cross-loadings removed from factors;

Food items with no loadings had loadings < 0.2

Pattern 1: 'Snacking' pattern;

Pattern 2: 'Energy-dense meat' pattern;

Pattern 3: 'Fruit and Vegetable' pattern;

Cronbach's α indicated that there was a moderate inter-item reliability (table 4.10) which was improved for each dietary pattern after the removal of white sugar, and drinks in pattern 1, oats in pattern 2, and oil-based dressings, fats and soy milk in pattern 4.

Pattern	Original Cronbach's α	New Cronbach's α	Items removed
1	0.619	0.639	White sugar, diet drinks
2	0.634	0.634	NA – highest value
3	0.641	0.648	Oats
4	0.252	0.431	Oil and oil-based dressings, fats, soy milk

Table 4.10 Cronbach's α score for each of the dietary patterns.

NA = Not applicable;

These patterns explained 6.9, 6.8, 5.6 and 4.8% of variation in food intakes, respectively

Each dietary pattern was then broken down into tertiles (T1, T2, T3) for further analysis. Those who scored highly on the pattern, reflected high consumption of those with positive factor loadings, and low consumption of those foods with negative factor loading, and were categorised as tertile 3.

In the 'snacking' pattern, there was a statistically significantly difference in age (F(2,228) = 3.36, P = 0.036) between tertiles 1 and 3. Women who followed the 'snacking' pattern more closely were older compared to those in lower 'snacking' pattern tertiles.

In the 'energy-dense meat' pattern, there was also a statistically significant difference between the age (32.8 years \pm 8.2, 32.9 years 8.1, 30.0 years \pm 8.3) and BMI (24.6 kg/m² \pm 4.2, 24.4 kg/m² \pm 4.0, 26.4 kg/m² \pm 6.7) between tertiles one, two and three, respectively.. Women who followed the 'energy-dense meat' pattern were younger compared to those in the lower 'energy-dense meat' pattern tertiles. Additionally women who followed the 'energy-dense meat' pattern had a lower BMI compared to those in the lower 'energy-dense meat' pattern tertiles. No other patterns showed a significant association with age, BMI or BF% (table 4.11).

	Age (years)	P-value	BMI (kg/m ²)	P-value	BF%
Pattern 1					
T1	31.9 ± 8.7*		24.2 ± 3.7		31.9 ± 7.3
T2	29.9 ± 8.0	0.026	25.2 ± 4.8		32.1 ± 8.4
Т3	33.9 ± 7.9*		25.9 ± 6.5		33.7 ± 8.5
Pattern 2					
T1	32.8 ± 8.2*	0.048	24.6 ± 4.2*	0.036	32.2 ± 7.4
T2	32.9 ± 8.1		24.4 ± 4.0		31.4 ± 7.2
Т3	30.0 ± 8.3*		26.4 ± 6.7*		34.3 ± 9.2
Pattern 3					
T1	31.7 ± 7.8		24.8 ± 4.4		33.2 ± 7.2
T2	31.1 ± 8.4		24.4 ± 4.3		31.5 ± 7.2
Т3	32.9 ± 8.6		26.2 ± 6.4		33.1 ± 9.5
Pattern 4					
T1	32.2 ± 8.6		25.2 ± 4.8		33.7 ± 7.5
T2	31.6 ± 8.6		25.5 ± 5.6		33.0 ± 8.1
Т3	31.9 ± 7.6		24.8 ± 5.1		31.1 ± 8.4

Table 4.11 Age, body mass index (BMI) and body fat percentage (BF%) characteristics of the tertiles in the dietary patterns for the NZE women (n = 231).

Pattern 1 = Snacking;

Values are means ± standard deviations;

T1, T2, T3 = tertiles of dietary pattern score

Pattern 3 = Fruit and vegetables; Pattern 4 = Healthy

Pattern 2 = Energy-dense meat;

* = statistically significant (P < 0.05) (ANOVA analysis within each pattern between tertiles);

When age, BMI and BF% were categorised into groups, age remained significantly associated with dietary pattern 2 only (χ^2 (4) = 10.132, P <0.038). A significantly higher proportion of women aged 16 to 24 years were in the highest tertile of the 'energy-dense meat' pattern reflecting higher consumption of those foods compared to women 36 to 45 years of age (table 4.12).

There was no association between 'snacking' pattern and BMI and BF% groups indicating that women consume foods from this pattern regardless of their body composition (table 4.12). In comparison, a higher proportion of women in the low BF% (<22%) group had higher consumption of food items from the 'fruit and vegetable' and 'healthy' patterns.

					Diet	tary pat	terns						
		Snacl	king pa	attern	Energ	gy-dens	e meat	I	Fruit an	d	Hea	Ithy pat	tern
						patteri	า	veget	ables p	attern			
Characteristics	n	T1	T2	Т3	T1	T2	Т3	T1	T2	Т3	T1	T2	Т3
Age (years)													
16-24	56	39.3	1.8	58.9	26.8	23.2	50.0*	26.8	44.6	28.6	33.9	33.9	32.1
25-35	89	33.7	3.4	62.9	32.6	37.1	30.3	39.3	30.3	30.3	32.6	33.7	33.7
36-45	86	29.1	0	70.9	38.4	36.0	25.6	31.4	29.1	39.5	33.7	33.7	32.6
BMI (kg/m ²)													
18.5-24.99	140	29.3	2.9	67.9	35.7	36.4	27.9	34.3	33.6	32.1	31.4	33.6	35.0
25.0-29.99	61	39.3	0	60.7	34.4	31.1	34.4	32.8	41.0	26.2	39.3	29.5	31.1
≥ 30.0	30	40.0	0	60.0	20.0	23.3	56.7	30.0	16.7	53.3	30.0	43.3	26.7
BF%													
< 22%	18	33.3	0	66.7	33.3	44.4	22.2	11.1	44.4	44.4	16.7	22.2	61.1
≥22-29.9%	66	30.3	4.5	65.2	34.8	30.3	34.8	37.9	28.8	33.3	30.3	39.4	30.0
≥ 30%	147	34.7	0.7	64.6	32.7	33.3	34.0	34.0	34.0	32.0	36.7	32.7	30.6
alues are percentage	s of the	total nu	mber ir	n that gro	oup;		N =	number	of partic	cipants;			

 Table 4.12 Socio-demographic characteristics of the NZE women in the tertiles of each dietary pattern.

Values are percentages of the total number in that group; T1, T2, T3 = tertiles of dietary pattern score;

* = proportions across tertiles within the dietary pattern differed significantly (P = 0.035; significant = P<0.05) (Chi-square analysis)

4.6 Diet Quality

The 'snacking' pattern had significantly different intakes of 18 out of 29 nutrients across the tertiles. This dietary pattern had moderate intakes of all nutrients compared to the other patterns (table 4.13).

The 'energy-dense meat' pattern had significantly different intakes of all nutrients except vitamin A and selenium across tertiles. Women with the highest and lowest dietary scores in this pattern had the highest and lowest intakes of energy, protein, carbohydrate, total fat, saturated fat, sugar, iron and zinc respectively (table 4.13). The nutrient and energy density of this pattern is changed considerably depending on the extent to which it is followed. Considering the food items associated with this pattern these findings were expected and thus confirm the naming of this pattern.

Across tertiles of scores on the 'fruit and vegetable' pattern, there were significantly different intakes for 18 out of 29 nutrients. Women with the highest and lowest dietary scores in this pattern had the highest and lowest intakes of vitamin C and vitamin A, respectively (table 4.13).

The 'healthy' pattern had significantly different intakes between the tertiles in 17 out of 29 nutrients. Women with the highest and lowest dietary scores in this pattern had the highest and lowest intakes of vitamin E, and those who had moderate scores in this pattern had the lowest intake of vitamin D, whereas those in tertile 3 had the highest intakes of vitamin D (table 4.13).

	EAR	S	Snacking pattern	5	Energ	Energy-dense meat pattern	oattern	Fruit	Fruit and vegetable pattern	ittern	T	Healthy pattern	
		T1	Т2	T3	11	12	ET	T1	T2	L3	T1	12	T3
Energy (kJ)	Na	8515.5	6.000	9420.4	7058.7 ±	8773.9 ±	11142.8 ±	8469.5 ±	8832.7 ±	9673.1±	9158.7±	8393.3±	9437.0 ±
		±2792.8	±2862.1	±3204.1	2106.6	1973.6	3057.6	3033.1	2617.3	3062.8	3368.9	2882.7	2432.6
P-value			0.159			0.000*			0.033*			0.073	
Protein (g)	Na	91.6	96.8	96.8	78.5 ±	91.4 ±	116.5 ±	87.8 ±	94.5±32.7	104.0±	93.8±	± 8.78	$104.9 \pm$
		±33.2	±31.0	±34.3	25.8	22.1	37.3	31.5		33.3	35.7	29.5	31.8
P-value			0.532			0.000*			0.009*			0.005*	
Total fat (g)	Na	83.7	84.7	88.2	64.6±	83.8±	107.9 ±	85.0±	82.4 ± 30.2	89.0 ± 32.5	86.5±	76.7 ±	93.3 ±
		±36.3	±28.5	±34.9	23.6	25.0	34.6	36.6			37.5	28.0	31.8
P-value			0.674			0.000*			0.469			0.007*	
SF (g)	Na	34.8	33.9	34.8	24.2 ±	33.5 ±	45.4 ± 18.2	35.0±	33.2 ± 13.3	34.8 ± 17.2	36.6±	30.6±	35.8±
		±18.9	± 14.1	±16.2	10.4	11.6		18.1	_		19.1	12.7	16.0
P-value			0.937			0.000*			0.753			0.044*	
PUFA (g)	Na	11.5	12.5	13.7 ±5.5	10.7 ±	12.4 ± 4.3	14.8 ± 4.7	12.1 ± 4.7	12.0 ± 4.7	13.8 ± 5.0	$11.6 \pm$	$11.3 \pm$	15.0±
	Ĩ	±4.2	±4.4		4.6					_	4.5	4.1	5.0
P-value			0.013*			0.000*			0.038*			0.000*	
MUFA (g)	Na	28.2	28.7	30.0	22.4 ±	28.6 ± 9.0	36.1 ± 11.1	28.9±	28.0 ± 10.7	30.3 ± 10.3	28.8±	26.2±	32.2 ±
		±11.6	±9.6	±11.9	8.6			12.3			12.0	9.6	10.9
P-value			0.580			0.000*			0.439			0.003*	
Cholesterol	Na	278.5	300.2	284.1	233.6±	282.9 ±	355.1±	272.9 ±	281.3 ±	317.3±	262.8±	246.1±	364.2 ±
(mg)		±164.2	±139.7	±147.1	127.8	141.0	189.4	171.3	155.4	158.2	118.0	100.0	219.0
P-value			0.651			0.000*			0.197			0.000*	
CHO (g)	Na	207.2	234.7	245.7	177.7 ±	221.4 ±	284.1 ±	207.2 ±	225.0 ± 69.1	250.9±	239.4 ±	219.3±	224.5 ±
	_	±75.4	±89.3	±96.7	59.7	71.6	95.4	87.8		100.6	93.0	91.0	80.0
P-value			0.021*			0.000*			0.008*			0.344	
Sugars (g)	Na	109.6	129.4	126.9	± 7.39	117.2 ±	148.3 ±	105.9 ±	115.8 ± 40.5	140.4 ±	125.8±	114.5±	121.9 ±
		±45.0	±59.6	±55.5	40.8	44.9	60.8	52.8		60.7	60.6	52.8	47.1
P-value			0.047*			0.000*	_		0.000*			0.420	
Alcohol (g)	Na	7.4 ±8.7	6.3 ±7.5	6.2 ±7.4	5.9 ± 8.3	7.5 ± 7.9	6.9±7.5	7.4 ± 8.8	7.4 ± 8.3	5.5 ± 6.4	6.1 ± 8.3	6.7 ± 7.9	7.5 ± 7.6
P-value			0.425			0.459			0.227			0.567	
Dietary fibre	Na	26.4	30.5	31.8	27.3 ±	29.1±9.2	32.2 ± 10.6	22.8 ± 7.0	28.9±7.4	36.8±9.9	26.6±	28.7±	33.2 ±
(g)		±8.5	±9.2	±11.4	9.6					_	9.6	9.9	9.4
P-value			0.002*			0.008*			0.000*			0.000*	
Thiamin (mg)	0.9	1.5 ±0.9	1.6 ±0.8	1.8 ±0.8	1.3 ± 0.8	1.6 ± 0.7	2.0 ± 0.9	1.5 ± 0.9	1.6 ± 0.8	1.9 ± 0.8	1.6 ± 0.7	1.7 ± 1.0	1.6 ± 0.6
P-value			0.164			0.000*			0.008*			0.366	
Riboflavin	0.9	2.5 ±1.1	2.7 ±1.4	2.5 ± 1.2	2.2 ± 1.1	2.5 ± 1.0	3.0 ± 1.5	2.4 ± 1.2	2.4 ± 0.9	2.8 ± 1.5	2.7 ± 1.5	2.4 ± 1.2	2.7 ± 0.9
(mg)													

Table 4.13 Nutrient intakes within the dietary pattern tertiles identified from the NZWFFQ among NZE women.

	EAR	SI	Snacking pattern	Ę	Ener	Energy-dense meat pattern	oattern	Fruit	Fruit and vegetable pattern	attern	T	Healthy pattern	
		T1	Т2	Т3	T1	Т2	T3	T1	Т2	Т3	Τ1	Т2	Т3
P-value			0.476			0.000*			0.050			0.226	
Niacin (mg)	11	21.9 ±9.8	22.7 ±11.4	22.7 ±9.9	18.7± 7.3	22.3 ± 8.1	26.7 ± 13.2	20.6 ± 9.8	21.2 ±8.0	25.8 ±12.2	22.3 ±12.6	21.2 ±9.5	24.3 ±8.4
		T1	T2	Т3	T1	T2	T3	T1	12	T3	T1	T2	T3
P-value			0.856			0.000*			0.003*	13*		0.173	
Vitamin C	30	142.5	168.2	165.3	140.6±	154.5 ±	175.0 ±	103.6 ±	153.4 ± 61.9	213.2±	143.2±	153.4 ±	173.8±
(mg)		±82.9	±84.5	±86.3	77.5	64.3	102.6	46.9	_	95.6	84.8	85.0	79.7
P-value			0.120			0.037*			0.000*			0.070	
Vitamin D (IJE)	Na	4.2 ±2.5	4.8 ±2.6	4.5 ±2.4	4.0 ± 2.4	4.3 ± 2.5	5.2 ± 2.6	4.5 ± 2.9	4.3 ± 2.3	4.8 ± 2.3	4.4 ± 2.7	3.9 ± 2.1	5.3 ± 2.5
P-value			0.329			0.006*			0.520			0.003*	
Vitamin E	Na	12.2	13.5	14.2 ±	12.1 ±	12.9 ± 4.4	14.9±4.8	11.7 ± 4.5	12.5 ± 4.5	15.6 ± 5.1	11.6±	11.8±	16.5 ±
(BIII)	T	14.0	14:0	7°C	C.C	÷,000			+000 000			0.0	7.0
P-value	,	1	0.035*			0.001*			0.000*			0.000*	
Vitamin B6 (mg)	1.1	2.5 ±2.0	2.7 ±2.2	2.6 ±1.9	2.1±0.9	2.5 ± 2.0	3.1±2.7	2.3 ± 2.1	2.4 ± 1.0	3.1 ± 2.6	2.7 ± 2.9	2.3 ± 1.7	2.8 ± 1.0
P-value			0.859			*600.0			0.014*			0.373	
Vitamin B12 (ug)	2.0	4.7 ± 2.9	4.9 ±2.7	4.5 ±2.4	3.7 ± 2.0	4.5 ± 1.9	6.2 ± 4.0	4.5 ± 3.2	4.6±2.7	5.3 ± 3.0	4.5 ± 2.7	4.3 ± 2.3	5.6±3.6
P-value			0.569			0.000*			0.242			0.009*	
Total-folate	320	410.6	424.7	451.4	381.8±	440.7 ±	463.7 ±	371.1±	405.6±	509.5±	416.3±	428.8±	441.3 ±
(hg)		±172.1	±134.3	±174.5	142.8	162.9	172.3	140.2	151.1	165.1	167.9	175.5	144.2
P-value			0.282	_		0.005*			0.000*				0.640
Vitamin A	500	1351.9 +444 0	1637.8 +777 1	1672.8 +004.0	1407.3 ± 605 2	1568.8 ± 777 E	1667.5 ± 761 0	1171.6± Ene 2	1504.9 ± 172 6	1967.0±	1384.6± 6106	1570.0 ±	1690.5 ±
P-value		0.44	0.011*	0.400-	0.000	0.081	0.10/	2.000	0.000*	7:700	0.010	0.031*	7.02/
Sodium (mg)	Na	2406.2	2458.3	2608.4	1832.9 ±	2386.9±	3280.0 ±	2469.2 ±	2438.9 ±	2591.8±	2631.7 ±	2339.6±	2531.0±
j		±959.9	±988.7	±1132.2	691.8	619.5	1124.1	1127.4	871.9	1077.9	1249.9	914.7	872.6
P-value			0.450			0.000*			0.623			0.200	
Potassium	Na	3465.3	3932.0	3903.7	3420.3 ±	3686.7 ±	4173.7 ±	3182.2 ±	3630.3 ±	4468.2 ±	3570.2 ±	3517.8±	4201.5 ±
(mg) D vichio	T	±10/0.4	±12/3.3	0.CUE11	C.CELL	948.2	1398.4	9/3./	7.000 0	L294.5	C.6/21	4.8011	1139./
		0 1.7 0	7007	0.077		000	- 1	. 0 0 7 0	0,000			100.0	
Magnesium (2014)	4 42	305.8 0 C 1 1 1	408.6 ±125 2	413.9 ±125 7	362.4 ±	388./ ± 106.2	436./±	348.U ± 107 0	391.6± ⊐ <<<	448.2 ±	367.5± 1 0.01	361.1±	460.5 ± 710 7
ח עבונים	T	0.2111	7.021T	/.CCLT	0.4CL	C-00T	0.621	6./UL	2.221	T-07T	T-77T	*000 0	/.011
P-Value	010	0,00,00	1.033 "	1 1 7	1000	. 100.0			0.000	1 200 1		112 1	
(mg)	040	±493.3	±532.2	±1/3.2 ±512.3	536.4	422.7	531.7 531.7	545.8	419.6	1203.0 ± 553.2	1211.4 - 589.1	489.2	443.1
P-value			0.399			0.001*			0.126			0.202	
Phosphorus	580	1616.3 ± 5 12 1	1760.9 4565 1	1748.3 4500 2	1482.3 ± EE2 2	1643.5 ±	2006.8 ±	1561.8± 526.7	1676.7 ± 522 2	1894.0± 502 7	1663.2 ±	1567.6± בסע 7	1906.2 ±
(1118)		4.040	T.COCT	C.UECI	C.2CC			1.000	7.000	1.000	COTO	1.400	7.00C
						1(105						

	EAR	S	Snacking pattern	rn	Ener	Energy-dense meat pattern	oattern	Fruit	Fruit and vegetable pattern	attern	-	Healthy pattern	
		T1	Т2	T3	T1	12	T3	11	7 7	T3	T1	T2	T3
P-value			0.217			0.000*				0.001*		0.001*	
Iron (mg)	8	11.5	12.6	13.1 ±4.3	10.5 ±	12.1 ± 3.2	14.6 ± 4.4	10.9 ± 3.7	12.1 ± 3.9	14.2 ± 3.7	$11.6 \pm$	$11.8 \pm$	14.0 ±
		±3.8	±3.6	_	3.3						4.0	3.9	3.8
P-value			0.035*			0.000*			0.000*			0.000*	
Zinc (mg)	6.5	11.3	11.9	12.0 ±4.3	9.4 ± 3.0	11.3 ± 2.8	14.6 ± 4.7	10.7 ± 4.1	11.7 ± 4.0	12.9 ± 4.2	$11.8 \pm$	$10.8 \pm$	12.8 ±
		±4.2	±3.8								4.5	3.6	4.2
		T1	T2	Т3	11	T2	T3	11	T2	T3	11	T2	Т3
P-value			0.541			0.000*			0.006*			0.011*	
Selenium	50	77.1	87.9	82.9	79.1 ±	76.0 ±	93.0 ± 44.9	75.1 ±	82.2 ± 45.3	90.8 ± 40.5	62.6±	69.5±	$116.5 \pm$
(bd)		±40.4	±68.0	±42.7	69.2	36.3		66.8			28.1	30.2	70.0
P-value			0.433			0.099			0.174			0.000*	
Values are mean intakes ± standard deviation;	ntakes ± si	es ± standard deviati			:		WFFQ = New Zea	aland Women's F	NZWFFQ = New Zealand Women's Food Frequency Questionnaire	uestionnaire			

* = statistically significant (P < 0.05) (ANOVA analysis within each pattern between tertiles); EAR = Estimated average requirements (National Health Medical Research Council, 2005) T1, T2, T3 = tertiles of dietary pattern score Va

MUFA = *Monounsaturated fat;*

PUFA = Polyunsaturated fat;

SF = Saturated fat; CHO = Carbohydrate

There were some significant differences between the macronutrient distributions in each of the dietary patterns (table 4.13). The macronutrient distributions were not significantly different across the tertiles of the 'snacking' pattern scores. Women who were in tertile 3 of the 'energy-dense meat' pattern had significantly higher protein and saturated fat intakes compared to those in tertiles 1 and 2. This suggests that women who closely follow this pattern may consume higher amounts of food items such as red meat or high fat cheese compared to those who follow it less closely. The 'fruit and vegetable' pattern had significantly higher intakes compared to those who follow it less closely. The 'fruit and vegetable' pattern had significantly higher intakes compared to women in tertiles 1 and 2. It is likely that these women have a higher fruit or starchy vegetable consumption compared to the women who follow the pattern less closely. The 'healthy' pattern had significantly higher intakes of protein and total fat in tertile 3 compared to tertiles 1 and 2. This suggests that women in tertile 3 consume more meat alternatives as protein sources and use more coconut oil, nuts and seeds in their cooking.

None of the dietary patterns met all AMDR's (table 4.14). All of the dietary patterns had mean protein intakes within the recommended range of 15 -25% of total energy intake. The carbohydrate intakes in all of the dietary patterns fell below the recommended range of 45 – 65% of total energy intake. The intake of total fat varied with half of the tertiles meeting the AMDR (20 – 35% of total energy intake), although these tertile intakes were at the higher end of the recommendations. All tertiles exceeded the AMDR of less than 12% of total energy intake for saturated fat. Overall, when analysing tertile 3 as the highest users of the dietary pattern, the 'fruit and vegetable' pattern showed the highest protein consumption and the 'energy-dense meat' pattern had the highest saturated fat intakes. Moderate intakes of carbohydrate and total fat were seen in tertile 3 of all the patterns as the highest intakes were present in tertile 1 of the 'healthy' pattern and tertile 1 of the 'fruit and vegetable'.

Macronutrient	•)	Snacking pattern	2	Energy	Energy-dense meat pattern	pattern	Fruit aı	Fruit and vegetable pattern	attern	-	Healthy pattern	c
(% of El)	T1	T2	T3	T1	T2	T3	T1	72	T3	T1	T2	T3
Protein	18.5 ± 3.0	18.5 ± 3.0 18.3 ± 3.3	17.8 ± 3.8	$19.1 \pm 3.8^{*}$	$17.9 \pm 3.4^{*}$	17.8 ± 2.9*	17.9 ± 3.7	18.3 ± 2.9	18.6±3.7	$17.6 \pm 2.9^{*}$	$18.1 \pm 3.3^{*}$	19.2 ± 3.9*
Carbohydrate	40.2 ± 7.8	40.2 ± 7.8 42.1 ± 6.7	42.6 ± 7.3	41.5 ± 7.1	41.2 ± 8.5	41.9 ± 7.0	40.1 ± 8.1	42.1±6.2	42.3 ± 8.1	43.2 ± 5.8*	42.5±6.4*	38.7 ± 9.3*
Fat	35.9 ± 7.6	34.5 ± 5.9	34.5 ± 5.6	33.8 ± 6.3	35.3 ± 6.9	35.9 ± 6.3	36.8±6.5*	34.2±5.3*	34.1 ± 7.3*	34.5±5.6*	33.8 ± 5.4*	36.7 ± 7.9*
Saturated fat	14.6 ± 4.6	14.6 ± 4.6 13.6 ± 2.9	13.4 ± 3.0		$14.1 \pm 3.4^{*}$	12.5 ± 3.0* 14.1 ± 3.4* 15.0 ± 3.9 *	$14.8 \pm 3.7^{*}$	$13.7 \pm 2.8^*$	14.8 \pm 3.7* 13.7 \pm 2.8* 13.1 \pm 4.1 * 14.3 \pm 3.6	14.3 ± 3.6	13.3 ± 2.7	13.9 ± 4.4
Values are means ± standard deviation; * = Statistically significant (P<0.05) (ANOVA analysis within each pattern between tertiles); T1, T2, T3 = tertiles of dietary pattern score; EI = Energy intake	± standard devia nificant (P<0.05) · of dietary patte	tion; (ANOVA analysi: rn score;	s within each pa	tern between tei	tiles);							

Table 4.14 Macronutrient distribution between dietary pattern tertiles.

Chapter 5 Discussion

Analysing dietary patterns within a population is increasing in popularity to assess diet-disease relationships. In contrast with a single nutrient approach, it analyses the whole diet giving insight to a population's eating patterns. From this, dietary advice can be given to improve nutrition and health outcomes. This study sought to explore the dietary intake and associated eating patterns of New Zealand European (NZE) women participating in the wider EXPLORE study. The objectives were to explore the dietary intakes of NZE women, identify eating patterns which are the strongest predictors of increased body fat, and explore differences in macronutrient profiles and micronutrient intakes between BCP groups. The findings demonstrate that several dietary patterns can be identified within this population, which explain some of the differences seen between individual body compositions. Other factors such as portion size, socio-economic status and food availability may play a role.

5.1 Participant Characteristics

The 231 NZE women involved in this study had on average a normal (18.5 – 24.9 kg/m²) BMI (24 kg/m²) and a high (\geq 30%) BF% (32.6%) body composition (World Health Organisation, 2014a, Oliveros, 2014). The 2013/14 New Zealand National Health Survey showed the mean BMI of NZE women to be 27.5 kg/m², indicating that the majority of women are overweight (Ministry of Health, 2014b). The 2014/15 National Health Survey also showed that 30.6% of NZE women have a BMI \geq 30 kg/m² indicating obesity (Ministry of Health, 2015b). This percentage has progressively increased since the 2006/07 National Health survey where 24.6% of NZE women were obese (Ministry of Health, 2015b). This cohort of women may therefore not be a representative sample of the New Zealand population (Ministry of Health, 2014a). The study attracted many women who were interested in health and nutrition and also those who were a part of sports clubs and teams. It is possible that these women eat a healthier diet and participate in physical activity more regularly than others in the population thus leading to a lower BMI for this cohort of women.

When comparing the NN, NH and HH BCP groups, a positive relationship was present between age, BMI and BF%. The NN group contained on average the youngest women by 5.6 years with a mean age of 28.2 years compared to the NH group (33.1 years) and the HH group (33.8 years). The NN BCP group also had women with on average the lowest BMI (21.4 kg/m²) and BF% (25.8%) compared to the NH BCP group (23.0 kg/m², 33.4%) and the HH BCP group (28.6 kg/m², 39.1%). This shows that there is a positive trend between increasing age and adiposity.

The 2014/15 National Health Survey also showed similar results where the percentage of NZE women classed as obese increased from 20.1% in women 15 to 24 years, 27% in women 25 to 34 years and 32.6% in women 25 to 34 years (Ministry of Health, 2015b). The difference in BMI between the NN and HH BCP groups was significant, as was BF% between all BCP groups apart from the HN's. These differences were expected as the participants were recruited and categorised based on their individual body compositions and allocated into the relevant groups. Age was not significantly different between any of the BCP groups which was also expected because the women were recruited in approximately equal numbers within age categories (16-24, 25-35, 36-45). This indicates that there is a significant difference between body compositions in women with similar BMIs. Although the NL and HN BCP groups were not originally body compositions recruited for, several participant body compositions changed category after the ADP analysis due to the increased sensitivity by this machine and strict testing protocol, compared to the BIA machine used for screening without any of the testing requirements. These women were therefore included in the analysis.

5.2 Dietary Intake Analysis

5.2.1 Total population

On average, the women's dietary intake met the Ministry of Health's EAR's for all micronutrients. This was also true for the RDI's and the AI's apart from vitamin D and iron. The EAR is set at a level which is sufficient to meet the nutrient needs of at least half of healthy individuals and are the lowest level of which is required to maintain a defined level of nutriture in an individual (Yates, 1998). These values are used to determine the adequacy of nutrient intakes in population groups, whereas the RDI is set at two standard deviations above the EAR and is set an intake level that exceeds the requirements of 97-98% of healthy individuals (Gibson, 2005). It is therefore not usually used for group analysis and is more appropriate for assessing an individual's nutrient intake and if, met indicates that that individual has a low risk of nutrient deficiency (National Health Medical Research Council, 2005, Gibson, 2005). For some nutrients the RDI as well as the EAR were compared to the intakes of these women. Some nutrients were consumed in high amounts and it was therefore interesting to analyse these intakes using both the EAR and RDI values.

It was expected that the average nutrient intakes of the NZE women recruited in this study would meet the EAR guidelines because the women included in this study were well nourished with no signs of nutrient deficiency, and were free of chronic disease. It was also expected that a proportion of women would fall below the RDI values for several nutrients because the RDI

values exceed the actual nutrient requirements of nearly all healthy individuals and therefore it will always result in an overestimation of the proportion of the group with inadequate nutrient intakes (National Health Medical Research Council, 2005, Gibson, 2005). It is important to identify a point at which nutrient intakes below the EAR and RDI are a concern. Gibson (2005) states that if the intake is below the EAR there is a 50% probability of inadequate intake in that group. This indicates that any intake below the EAR reference values would be concerning and needs to be addressed. Some nutrients do not have an EAR value and therefore their intake needs to be compared against the RDI. As the proportion of the group below the RDI is likely to be exaggerated as stated above, a cut-off value has been outlined by Gibson (2005) which allows approximation of inadequate intakes. It states that 77% of the RDI should be met by individuals to ensure nutrient adequacy and anything below indicates insufficient nutrient intakes (Gibson, 2005).

Iron

The dietary iron intakes in the NZE women met the EAR but not the RDI. This indicates that on a group level at least half of the women are meeting their iron requirements per day; with only 3.4% of NZE women below the EAR however on an individual level iron intakes are poor. The majority of women (91.7%) were consuming iron intakes (12.6 mg/d) below the RDI (18 mg/d). These findings are similar to that of the 2008/09 National Nutrition Survey with an estimated 34.2% of women aged 15 to 18 years and 15.4% aged 31 to 50 years having inadequate iron intakes (University of Otago and Ministry of Health, 2011). Similar results have also been seen in premenopausal women elsewhere, with inadequate iron intakes prevalent in the USA and Canada (Gadowsky *et al.*, 1995, Stang *et al.*, 2000, Cooper *et al.*, 2006). Iron deficiency is the most common nutrient deficiency worldwide and although many dietary factors such as vitamin C, phytates, calcium and polyphenols modify iron absorption, inadequate dietary iron intake is the main cause of iron deficiency (Swanson, 2003, Beck, 2014, Coad and Pedley, 2014). Conversely, Kennedy and Meyers (2005) found American women to have adequate intakes above the EAR. The premenopausal women in a study by Mazess (1991) also had iron intakes above the EAR showing that intakes are very variable between different populations.

Although the majority of women in this study did not meet the RDI this was expected as the value tends to overestimate the proportion of women with poor intakes. Kennedy and Meyers (2005) found that if the RDI value was used for comparison the rate of insufficient iron intake would be a lot higher compared to when using the EAR value. More importantly is that all women meet the EAR as well as at least 77% of the RDI to ensure minimal risk of deficiency. The majority of the women (94%) met the EAR indicating that NZE women on average are

consuming iron intakes which are likely sufficient to meet their nutrient requirements. Nevertheless, it is well known that premenopausal women in New Zealand consume low dietary iron (University of Otago and Ministry of Health, 2011, Gibson *et al.*, 2002). It is therefore likely that there may be a few women who in fact do have excellent iron intakes and this may be giving a false impression of the actual iron intakes of the other NZE women. This is indicated by the large proportion (91.7%) of women who fall under the RDI value. It is also shown by the RDI cut-off point where 67% of women were found to have iron intakes below 77% of the iron RDI. In addition, the 6% of women who did not meet the EAR reference value are not consuming sufficient dietary iron, showing that a total of 82% of women have inadequate iron intakes. This indicates over three quarters of the NZE women are consuming inadequate dietary iron.

Premenopausal women need higher amounts of dietary iron due to menstruation, growth in adolescence and during pregnancy (National Health Medical Research Council, 2005). Fergusson *et al* (2001) found seven to 13% of New Zealand women aged 15 to 49 years to be iron depleted. This indicates therefore that many women in this study may in fact be iron depleted and at a high risk of iron deficiency anaemia. This is concerning given the essential role of iron in several metabolic functions and the adverse effects associated with low iron status such as impaired cognitive and immune function, decreased work performance, low energy levels, and poor pregnancy related outcomes (Brownlie *et al.*, 2004, Murray-Kolb, 2011, Scholl, 2005, Allen, 2000). Given the low iron intakes among these women interventions to increase dietary iron consumption and to increase iron absorption enhancers such as ascorbic acid should be implemented to increase iron status in premenopausal women (Heath *et al.*, 2001).

Vitamin D

The average intake of vitamin D in this study was 4.6 µg per day, and it was found that 64% of women consumed below the adequate daily intake for vitamin D. Vitamin D's primary role in the body is to assist with calcium absorption thereby maintaining bone health (National Health Medical Research Council, 2005). Vitamin D is obtained through both dietary sources and cutaneous synthesis from sunlight, making its status in individuals difficult to measure (National Health Medical Research Council, 2005). Given the current food supply, sufficient vitamin D status is nearly impossible to obtain through diet alone (Fuller, 2001). Calvo *et al* (2005) concluded that without fortification of foods dietary intake of vitamin D is insufficient to obtain adequate vitamin D status. Rockell (2006) conducted a study investigating vitamin D status in New Zealanders over 15 years of age and found the mean serum 25-hydroxyvitamin D

concentrations in New Zealanders to be 50 nmol/L indicating poor vitamin D status. Only 3% of individuals were found to be vitamin D deficient (≤17.5 nmol/L), however insufficient vitamin D levels were found in 84% of participants (≤80 nmol/L) (Rockell, 2006). Although dietary vitamin D consumption was not determined, these results indicate that dietary vitamin D was inadequate to maintain adequate vitamin D status. New Zealanders tend to consume low levels of foods rich in vitamin D such as fatty fish and organ meats and this is reflected in 55% of the women consuming inadequate vitamin D. The low intakes observed in this study are concerning as the adequate intake value is estimated to be adequate for a group of healthy individuals (Gibson, 2005). Intakes below this level therefore suggest inadequate intakes, especially during the winter months in New Zealand as UV exposure is poor due to less time spent outside and people tend to wear more clothing (Rockell, 2006).

Poor vitamin D intakes can be supplemented with adequate sun exposure throughout the summer months in New Zealand. Sunlight exposure is crucial to obtaining adequate vitamin D status for good health, and seasonal changes, especially in New Zealand have a significant impact on the cutaneous synthesis of vitamin D through the skin (National Health Medical Research Council, 2005, Rockell, 2006). It is likely that seasonality plays a large role in vitamin D synthesis among New Zealanders with large differences seen in serum vitamin D (Rockell, 2006). Concentrations were found to be lowest during spring and the highest during the summer months with the estimated vitamin D synthesis during the winter months in Christchurch to be only 60 IU per day (Rockell, 2006). This indicates that dietary vitamin D intake is essential during the months with minimal UV light exposure. Similarly, Pasco (2001) also confirmed that during the winter months dietary vitamin D influences vitamin D status. As blood samples were not analysed as part of this study we speculate that the vitamin D status in these NZE women is poor, especially throughout the months with minimal UV exposure.

Obtaining adequate dietary vitamin D is also challenging due to no mandatory fortification of food vehicles in New Zealand. In America the fortification of milk and some breakfast cereals is mandatory, whereas in New Zealand fortification of food items is voluntary (Rockell, 2006, Calvo *et al.*, 2005, Calvo and Whiting, 2003). The American population have a higher vitamin D status compared to those in New Zealand which is likely due to their mandatory fortification of ways to improve vitamin D consumption are warranted. Mandatory fortification of food vehicles such as margarine and milk may help NZE women obtain sufficient dietary vitamin D intakes, especially throughout the winter months.

Calcium

On average, the NZE women were meeting the EAR for calcium, however 17% still had intakes below the EAR, and 38% were not meeting the RDI. Low calcium intakes are common throughout dietary assessment with the 2008/09 National Nutrition Survey showing that 71.2% of NZE women in the NZ population were consuming inadequate calcium intakes (University of Otago and Ministry of Health, 2011). Calcium is stored in teeth and bone and is essential to maintain structure and strength of the skeleton (National Health Medical Research Council, 2005, Flynn, 2003). Calcium in addition to vitamin D therefore also plays a critical role in bone health as shown by Ransdale et al (1994) who found a significant inverse relationship between dietary calcium and bone density independent of body mass. The proportion of women consuming less than 77% of the RDI was 11.5% and this is concerning because combined with the 17% below the EAR means that 28.5% of NZE women have insufficient calcium intakes. Adequate calcium consumption is essential, especially during young adulthood to attain peak bone mass, and also during adulthood to slow the decline of bone density (University of Otago and Ministry of Health, 2011). The low calcium intakes seen in New Zealand also extend to other countries such as America. Fleming (1994) showed that a large proportion of individuals do not meet the RDA and that women between 12 and 29 years consume less than 60% of the recommended daily calcium intake. It is essential that premenopausal women attain adequate calcium intake to decrease the risk of fractures and osteoporosis during later adulthood (Flynn, 2003). Strategies therefore should be implemented to improve the calcium status of NZE women.

Folate

Dietary folate intakes were low with 25% of NZE women not meeting the EAR and 46.3% of women not meeting the RDI. This is reflected in the 2008/09 National Nutrition Survey, where 4% of premenopausal women had red cell folate levels indicating a high risk of a neural tube defect (NTD) pregnancy, and only 27% of women had levels indicating a low risk (University of Otago and Ministry of Health, 2011). Folate is vital for healthy cell growth and development and is an essential nutrient in pregnancy to prevent neural tube defects (National Health Medical Research Council, 2005). In 2003, the prevalence of NTD rates was the highest in early pregnancy terminations in New Zealand (5.7 cases per 10,000 births), followed by live births (3.4 per 10,000) (Ministry of Health, 2015a). This prevalence has increased since 1999 where NTDs accounted for five per 10,000 births (live births and stillbirths) (Ministry of Health, 2003). Lewis *et al* (1999) estimated that 68-87% of women premenopausal women had inadequate serum folate levels and a review of folate intakes in Europe also showed that only a small

proportion of women meet the recommended dietary intake of folate per day (>350 µg/d) (De Bree *et al.*, 1997). This aligns with the intakes in this study where a quarter of the women are not consuming adequate folate intakes. Further, the proportion of women consuming below 77% of the RDI is 23%. This indicates that 48% of women are consuming inadequate folate intakes. As these women are of child-bearing age this is concerning as folate is essential for proper brain function, aids in the production of DNA and RNA, especially during periods of rapid growth such as during pregnancy, therefore deficiencies may give rise to neural tube defects (Mrc Vitamin Study Research, 1991, Smithells *et al.*, 1976, Wagner, 1996). Public health strategies to increase the folate intakes of these women should be considered such as the mandatory fortification of folic acid in bread and baked items.

Currently folic acid fortification of bread is voluntary in New Zealand. USA, Chile, Canada and Australia all have mandatory folic acid fortification of flour (Ministry of Health, 2015a). This fortification has been proven to be effective in Canada, reducing the rate of NTDs by 46% (De Wals *et al.*, 2007) and in 2012 Australian women of all ages were consuming folate intakes above the EAR (Australian Bureau of Statistics, 2014). Although many bread companies in New Zealand currently fortify their products with folic acid, mandatory fortification may be beneficial in increasing folate status in NZE women. Nevertheless, as this strategy has already been considered in New Zealand, alternative interventions should be developed.

Dietary fibre

Dietary fibre intakes for the majority of the NZE women were adequate (30 g/d), however 28% were still consuming less than the adequate intake (AI) per day (25 g/d). This contradicts the findings in the 2008/09 National Nutrition Survey which found that women were consuming on average 18 g of fibre per day (University of Otago and Ministry of Health, 2011). The women in this study therefore may have much higher fibre intakes compared to the rest of the New Zealand population. One reason for this may be the type of women this study attracted. Some of the women were interested in health and were part of sports teams therefore they may have already been eating a balanced healthy diet, including foods from the four main food groups. Alternatively, given that 28% of women had intakes below the AI, it is likely that there are a few women who have excellent dietary fibre intakes have also been shown in the United States of America (USA) where on average less than half the recommended intakes (15 g) are consumed (Slavin, 2005, Howarth *et al.*, 2001). This has also been seen in the United Kingdom (UK) where the average dietary fibre intake is 20 g per day. (Bingham, 1979). Additionally, the women in this study may also have over-reported their intakes of high fibre foods in the

NZWFFQ. Nevertheless, the proportion of individuals who have low fibre intakes is concerning as inadequate fibre intake has been clearly linked with body weight regulation and bowel health (Burton-Freeman, 2000, Howarth *et al.*, 2001, Marlett *et al.*, 2002). A study by Du *et al* (2010) found an inverse relationship between dietary fibre intake and BMI and waist circumference as did Murakami *et al* (2007) who showed an independent negative association between body weight and dietary fibre intakes. Interventions to increase the dietary fibre intakes of NZE women should be implemented and targeted at those who currently do not meet the recommended guidelines.

5.2.2 BCP groups

There were notable differences in the nutrient intakes between some of the BCP groups; however these were not significantly different. Presented below are findings from the HH, NN and NL BCP groups which had the highest and lowest intakes of the nutrients.

HH group

The HH group had the largest intake of energy (9296.2 \pm 2505.5 kJ/d), total fat (89.4 \pm 31.5 g/d), saturated fat (36.5 \pm 15.4 g/d), monounsaturated fat (30.4 \pm 10.8 g/d), and sodium (2661.9 ± 1028.1 mg/d). The high energy intake may be due to the accompanying high fat consumption in this group of women. Fat is energy-dense providing 37.7 kJ/g compared to that of carbohydrate which provides 16.7 kJ/g, therefore providing more energy in smaller amounts of food (National Health Medical Research Council, 2005). Fat is also highly palatable and therefore may lead to overconsumption and increase body fatness (Bray et al., 2004). This is evident in these women who had both a total fat and saturated fat intake above that of the NN BCP group. Studies have shown that energy-dense food items high in fat tend to be less satiating leading to weight gain and subsequently overweight body compositions (Blundell and Macdiarmid, 1997). Tremblay (1989) found that a diet high in fat leads to increased adiposity which is likely due to the hyperphagic effects of a high fat intake. This suggests that individuals in the HH BCP group may not only consume more energy-dense foods than the other BCP groups, but they may also consume them in larger portion sizes leading to their high BMI's and BF%. Interestingly, the energy intake of the HH BCP group ($9296.2 \pm 2505.5 \text{ kJ/d}$) was similar to that of the NN BCP group (9198.3 \pm 2141 kJ/d). This may be due to the HH BCP group underreporting their intake. Murakami and Livingstone (2015) showed overweight and obese individuals to be associated with under reporting of energy intakes which may explain the similarities seen between the HH and NN BCP group energy intakes.

The HH BCP group also had the highest sodium intakes. Although the NN (2457.5 \pm 713.3 mg/d) and NH (2468.6 ± 771.6 mg/d) BCP groups also consumed sodium intakes above the recommended upper limit of 2300 mg per day, the HH BCP group had the highest intakes consuming 2661.9 mg per day (table 4.5). The high sodium intakes in these women are similar to the results of from Ministry of Primary Industries investigation (Skeaff, 2013). They found that almost 93% of adults had urinary sodium excretion values indicating sodium intakes above the suggested dietary target (SDT) of 1600 mg per day for sodium (Skeaff, 2013). Additionally, they found 76% of individuals had sodium excretion values indicating sodium intakes above the recommended upper level of 2300 mg per day (Skeaff, 2013). Energy dense food items high in both fat and salt lead to over consumption and increased adiposity (Birch, 1999). The positive relationship that can be seen between these two nutrients can have a profound effect on health outcomes such as a high salt intake leading to hypertension which when coupled with a high fat intake and high BF% is a major risk factor for coronary heart disease and stroke (Nishida et al., 2004). The high sodium intakes in the HH BCP group indicates this group may not only consume foods that are high in fat and saturated fat, but also those which are associated with a high salt content such as bread, breakfast cereals, cheese, processed meats, sauces and spreads (Heart Foundation, 2013).

The HH BCP group also had the lowest intakes of vitamin C (131.6 mg), folate (408.9 µg) and calcium (1159.5 mg) compared to the other BCP groups, although this difference was not significant. Interestingly, these mean values also met the EAR and RDI recommendations for each nutrient indicating sufficient intakes. This is similar to Chai (2010) who found no significant differences in the dietary intakes of lipid-soluble micronutrients between different BMI's. Kimmons (2006) also did not see any differences in the vitamin C intakes between individuals of different BMI's which is similar to the results from this study. Although the calcium intake (1159 mg) of these individuals met the RDI (1000 mg), the intake was still lower than the other BCP groups. Literature agrees with this finding with studies showing low calcium intakes to be associated with a higher BF% (Zemel, 2000). Tidwell and Valliant (2011) also showed that individuals with lower calcium and vitamin D intakes had higher levels of adiposity. The lower intakes of nutrients in the HH BCP groups, and strategies to increase the nutrient density of their diets would be beneficial.

NN group

The NN group had the lowest protein (93.4 g), B12 (4.5 μ g), iron (11.9 mg) and zinc (11.7 mg) consumption per day, however had the highest carbohydrate (242.2 g), and sugar (130.2 g)

intakes. Despite having the lowest protein, B12 and zinc intakes these values are still above the EAR and therefore are not a concern. Iron intakes in this group; although above the EAR, are still significantly lower than the RDI. Given the proportion of women in the total cohort who are consuming insufficient iron intakes it is likely that this also extends to this group of women (Looker *et al.*, 1997). The low intakes of nutrients seen in these women are all associated with animal protein. Red meat is a great source of protein, B12, iron and zinc (Williams, 2007). It is therefore possible that these women consume red meat less frequently than women of other BCP groups and therefore the low levels of these nutrients are seen. Schulze (2006) and Maskarinec *et al* (2000) both found an inverse relationship between a dietary pattern consisting of a high intake of red meat and weight gain among women which further confirms this association.

The higher carbohydrate intakes seen in the NN BCP group indicate that individuals who have a lower BMI and BF% tend to consume a greater proportion of carbohydrate. This is reflected in the body compositions associated with each of the BCP groups. The median BMI in the NN group was 21.4 kg/m² and BF% 25.8% compared to 28.6 kg/m² and 39.1%, respectively in the HH BCP group. Similar findings have also been demonstrated in the literature. Miller *et al* (1990) found obese individuals to consume 46% of energy intake from carbohydrate whereas lean individuals consumed 53% of total energy as carbohydrate. Merchant *et al* (2009) also found that consuming a diet with less than 47% of total energy from carbohydrate was associated with an increased risk of being overweight or obese among healthy adults. Similarly, Bowman's (2002) results also support the higher consumption of carbohydrate in individuals with lower BF%'s.

These findings of higher carbohydrate consumption among women with a NN body composition indicate that these individuals consume less meat and high fat items but more fruit, breads and whole grains. These food items are more energy restrictive and nutrient dense which likely contribute to these women's body composition (Bowman and Spence, 2002). Literature also reflects this trend and shows that individuals who follow a 'healthy' dietary pattern consume a diet with a higher percentage of total energy from carbohydrate and are more likely to have lower BF%'s (Newby *et al.*, 2003). Controversy however is present in the literature, where Chai (2010) showed that individuals with higher carbohydrate and fat intakes are more likely to be overweight or obese. This may be due to the type of carbohydrates consumed, where items that are both high in refined carbohydrate and fat tend to be nutrient poor and energy-dense leading to passive overconsumption and subsequent weight gain (Drewnowski, 2007).

Although the intakes in this group are referred to as the highest intake, they are in no way excessive and are no cause for concern as discussed in the following section on macronutrient distribution. Overall, these women consume a nutrient dense diet but still follow the trend of inadequate iron intakes.

NL group

The NL group had the highest PUFA (13.5 g), dietary fibre (32.9 g), folate (467.4 μ g), calcium (1409.4 mg), iron (13.1 mg), zinc (12.8 mg), and selenium (102.0 mg) intakes per day. These findings indicate that the NL group have a nutrient dense diet and this is reflected in their body composition of a normal BMI with a low BF%. It is likely that these individuals consume a low energy but nutrient dense diet. As far as we are aware, no other studies have been conducted looking at the dietary patterns of premenopausal women with normal BMI and low BF%, therefore no similarities to other studies can be drawn. Nevertheless, these individuals are still consuming iron intakes below the RDI continuing the trend of low iron consumption regardless of body composition.

5.2.3 Macronutrient distribution

The macronutrient distribution profile of the NZE women did not meet all of the AMDRs. The women on average consumed a low carbohydrate and high saturated fat diet with moderate protein intakes. This trend was consistent when comparing the sample population as a whole, within each BCP group and in each of the dietary patterns.

Protein

Dietary analysis revealed the NZE women in this study meet the AMDRs for protein, consuming on average 18% of total energy intake. The Ministry of Health's AMDRs advise an intake of 15 to 25% of energy from protein to maintain a balanced diet (National Health Medical Research Council, 2005). These results are similar to that found in the New Zealand National Nutrition Survey (Ministry of Health, 2014b) where the NZ population consumed 16.4% of total energy as protein.

Results between the BCP groups were also similar to the NZ population where all groups met the AMDR. There was however a significant difference between the protein consumption of the three main BCP groups (NN, NH, HH) where the NN group consumed the least amount of protein (17.3±2.7%) and the HH group the most (18.3±3.4%). Although the HN BCP group consumed 22.3% of energy from protein, however because there were only three participants in this group this is not a good representative of individuals with this body composition. The New Zealand National Nutrition Survey showed that the average protein intake was 73 g, whereas the intakes of the BCP groups were all over 90 g of protein per day. This indicates that the women in this study consume larger amounts of protein compared to the New Zealand population.

Many studies have shown beneficial effects of increased percentage intake of protein relative to carbohydrate intakes for weight reduction (Layman, 2003, Noakes *et al.*, 2005, Skov, 1999). The protein intakes in this study were the highest among the HH BCP group and high BF% and these women also had high energy intakes. The high protein intakes may be associated with larger portion sizes and overall energy intakes (Kral and Rolls, 2004). The correlation may also be due to the type of protein consumed. Studies have found positive relationships between meat consumption and weight gain (Wang and Beydoun, 2009, Vergnaud *et al.*, 2010). This relationship is thought to be due to meat containing higher amounts of energy, total and saturated fat leading to weight gain (Vergnaud *et al.*, 2010). Conversely, the consumption of dairy products have been shown to be associated with weight regulation (Mirmiran *et al.*, 2004, Zemel, 2004). Zemel (2003) showed that increasing dietary calcium results in reduced adiposity in the absence of energy restriction in obese individuals. As the HH BCP group had the largest protein intakes and the highest BF%, it is likely that they consume more meat and less dairy products compared to the other BCP groups which contribute to their body composition.

Total fat

The NZE women consumed on average 34.8% ($86.7 \pm 28.0 \text{ g/d}$) of energy from total fat, meeting the AMDR of 25 to 35% of total energy intake. This is higher than the NZ population who consume 33.6% (69 g/d) of energy as total fat (University of Otago and Ministry of Health, 2011). This trend has also been seen in the United States where dietary fat provides 31.9% to 36.9% of total energy intake per day in adults (National Health and Medical Research Council III, 1988-1991). Intakes of total fat over the AMDR were found in 39% of women. These high total fat intakes were also present within some of the BCP groups where significant differences were found. The NN, NL and HN groups, although in the upper end of the recommendations, met the AMDR; however the NH and HH groups consumed above the AMDR with 35.5% (87.5 g) and 35.3% (89.4 g) of energy coming from total fat, respectively. This indicates that the higher the BF% the more total fat is consumed. Miller *et al* (1990) showed that obese individuals consumed 35% of energy from fat, compared to lean individuals who consumed only 29% from fat. The individuals in the HH BCP group consumed the least amount of carbohydrate and the same amount of protein compared to the other groups, indicating that they consumed the rest of their energy as fat. They also had the highest amount of total

energy consumption (9296.2 ± 25055 kJ/d) out of all the BCP groups which likely plays a role in their high BMI and BF%. Previous studies have found that women had a higher risk of obesity when a greater percentage of energy was obtained from fat, protein or animal protein indicating that excessive total fat intakes in the NZE women may contribute to excess adiposity and an adverse body composition (Murtaugh *et al.*, 2007). High fat intakes in these women are therefore likely to be contributing to their excess adiposity which may lead to adverse health outcomes such as cardiovascular disease, insulin resistance and type 2 diabetes (Nicklas, 2001).

Saturated fat

The total NZE women in this study did not meet the AMDR for saturated fat (<12% total energy intake), consuming on average $13.9 \pm 3.9\%$ of total energy intake. These findings align with those from the New Zealand National Nutrition Survey where the NZ population consumed 13% of total energy intake as saturated fat (University of Otago and Ministry of Health, 2011). Each of the BCP groups and all four dietary patterns also had saturated fat intakes above the AMDR indicating that no BCP group follows the AMDR guidelines. This is concerning as excess saturated fat intake has been associated with increased adiposity (Hariri et al., 2010, Field et al., 2007). Miller (1990) found that obese women derive a greater proportion of total energy intake from fat $(36.3 \pm 1.5\%)$ compared to lean women $(28.6 \pm 1.5\%)$. This has also been shown in a study by Alfieri (1997) who showed that overweight and obese individuals consume a diet significantly higher in dietary saturated fat compared to their lean counterparts. This has been reflected in the study participants where women in the HH BCP group had the highest saturated fat intakes compared to those who were in the HN BCP group who had the lowest intakes. The women in the NN and NH BCP groups also had saturated fat intakes above that of the recommended level indicating that their saturated fat intakes may result in progressive weight gain over subsequent years.

A further concern is the link between excess saturated fat intake and adverse health outcomes. It is well established that excess adiposity is associated with the development of chronic diseases such as CVD, type 2 diabetes, hypertension, insulin resistance, dyslipidaemia, osteoarthritis, sleep apnoea, metabolic syndrome, and some cancers (Kannel *et al.*, 1996, Nicklas, 2001, Oliveros, 2014, James *et al.*, 2004, Manson, 1990). These women were screened prior to recruitment for the presence of any chronic diseases, however on-going consumption of high saturated fat intakes may lead to an increased risk of these morbidities. Decreasing saturated fat intakes have been shown to improve health outcomes and decrease the risk of chronic disease (Hu, 2001).

Carbohydrate

On average the NZE women consumed 41.9% (233.9 ± 74.6 g/d) of total energy intake from carbohydrate (table 4.4) which is below the AMDR guidelines (45-65% of total energy intake) (National Health Medical Research Council, 2005). This intake is also below the mean carbohydrate intake for the NZ population who consume 47.1% (211 g/d) of energy intake as carbohydrate (University of Otago and Ministry of Health, 2011). These findings indicate that these women consume a higher proportion of protein and fat to make up their total energy intake each day. Further analysis revealed 64% of the NZE women were consuming less than the AMDR guidelines. Merchant et al (2009) suggested that consuming less than 47% of energy from carbohydrate is associated with an increased risk of obesity. This has also been shown by other studies where Lluch (2000) found carbohydrate intakes to be 44% of total energy intake and total fat intakes to be 37.5% of total energy intake showing that this pattern of macronutrient intake is common among overweight women. Nevertheless, the low carbohydrate intakes in these women also trended into the BCP groups where the NN group consumed significantly more carbohydrate (43.2 \pm 6.6%) than the HH BCP groups (41.5 \pm 7.7%). As these intakes are below 47% of total energy intake it may be that the women in the NN BCP group are at an increased risk of weight gain. Conversely, many studies show the role of a higher carbohydrate diet in weight gain moderation (Miller et al., 1990, Bowman and Spence, 2002). Nevertheless, Miller et al (1990) and Bowman et al (2002) both referred to carbohydrate intakes of 53% and 55% of total energy intake, respectively to aid in weight management. These intakes are therefore higher than what the women in this study are consuming which further confirms that they may be at an increased risk of weight gain.

Several other factors may also explain the low carbohydrate intakes seen in these women. Firstly, a low carbohydrate diet may be eaten to compensate for the consumption of foods with undesirable nutrient profiles. For example, consuming fewer carbohydrates means this energy needs to be replaced with protein or fat to maintain satiety and body weight (Miller *et al.*, 1990). The dietary data suggest that these women compensate the low carbohydrate intakes for a higher consumption of fat, where excess total fat intakes were seen in 39% of the women and 66% were consuming over the recommendations for saturated fat.

Secondly, a low carbohydrate diet has been highly publicised as a healthy diet to maintain or lose weight. The presence of this pattern in the majority of the women is not surprising as dietary trends in the media currently promote the 'high fat low carbohydrate' and 'paleo' fad diets which typically consist of low carbohydrate and high protein and fat (Cordain *et al.*, 2005). Carbohydrates, especially sugar have been portrayed in the media to be detrimental to

health and the cause of major weight gain and the current obesity epidemic (Swinburn, 2004). Although this claim may be true for many energy-dense high added sugar foods such as cakes, biscuits, refined breads and SSBs; whole grain breads and cereals provide essential nutrients to a balanced diet (Gil *et al.*, 2011). Many individuals, especially women succumb to current trends in the media and thus this may be why the identified dietary patterns reflect this way of eating. Low carbohydrate intakes however have also been reported as having the potential to be detrimental to health when followed over several years (Bilsborough, 2003). Complications include health arrhythmias, cardiac contractile function impairment, osteoporosis, kidney damage, lipid abnormalities and increased cancer risk (Bilsborough, 2003). As the emergence of this dietary pattern is relatively new, further research is required to determine the safety and efficacy of following this pattern long term.

Thirdly, underreporting may play a role. Dietary reporting from the participants may not have been quantifiably precise. Retrospective dietary intake is often misreported as it is cognitively challenging for many participants to complete an extremely accurate FFQ (Thompson, 2008). An individual's perception of what they eat may be completely different to what they actually consume each day, therefore leading to discrepancies such as under reporting in the final data.

Finally, social desirability and social approval bias have the potential to severely skew participant reporting (Hebert, 1995, Gibson, 2005). Social desirability is the participant wanting to display an image keeping within the social norm, and social approval is where participants report their intake to avoid criticism and achieve what the researchers are looking for (Hebert, 1995, Gibson, 2005). Scagliusi *et al* (2003) found that women were more likely to report their energy intake in a socially desirable way and food items which are considered 'fattening' or 'unhealthy' such as lollies, fried foods and refined breads were more likely to be under reported. It is therefore possible that social desirability in regards to their carbohydrate intake beliefs may have played a role in their reporting.

5.3 Top 40 Food Items

The top 40 food items consumed by the NZE women were determined for the total population and the three main BCP groups (NN, NH, HH). These were determined using daily equivalent frequencies of consumption reported in the NZWFFQ. Mean daily weights and the number of participants consuming the items daily were also determined.

Whole population

The most frequently consumed foods for the total NZE women included a range of food and beverages outside of the four food groups such as water, oil, coffee, tea and butter. Fruit and vegetables comprised the majority of the other items, however foods from the breads and cereals and milk and milk product food groups were also present (e.g. milk, yoghurt, wholemeal bread, and crackers) (table 23). Interestingly given the adequate protein intakes of this group, the only item from the lean meat and legume food group was eggs with no animal protein in the top 40 food items, suggesting the reason for the increased frequency of consumption seen for fruit and vegetables per day. For example, most individuals would be consumed several times per day (University of Otago and Ministry of Health, 2011).

Butter was the fifth most commonly consumed food in the NZE women which aligns with the National Nutrition Survey, where butter was the largest contributor to saturated fat intake (University of Otago and Ministry of Health, 2011). Wholemeal and multigrain breads were also commonly consumed by the NZE women aligning with the national nutrition survey where 65.9% of women ate a light or heavy grain bread (Ministry of Health, 2014b). In association with the macronutrient distributions, only four items from the breads and cereals food group were present in the top 40 items. This may contribute to the low carbohydrate intakes seen within this population as both the portion size and the number of people consuming these daily, were small. Although apples, bananas and oranges were also present in the top 40 foods these were also consumed in small quantities with a mean 84.7g per day of apple consumed equalling approximately one small apple. In countries around the world such as the USA, few individuals have been shown to meet the fruit and vegetable guidelines (Kimmons et al., 2009, Casagrande et al., 2007). Further discrepancies have been seen in the literature between the recommended daily intake of actual fruit consumption by individuals. Gonelevu et al (1997) found that both normal and overweight women did not meet the recommended five plus servings per day of fruit and vegetables and on average reached three and a half serves. In New Zealand, 65.8% and 72.7% of women ate the recommended fruit and vegetable intakes per day, indicating that over a quarter of women still have inadequate intakes (University of Otago and Ministry of Health, 2011).

Discretionary items which were commonly consumed include lollies, chocolate bars and diet soft drink. This is similar to those identified by Quatromani *et al* (2002) who found a dietary pattern characterised by high intakes of lollies, and high fat foods, and low intakes of fruits, vegetables and lean meats. A similar pattern was also identified by McNaughton *et al* (2008)

which was characterised by sweet and savoury snack foods, potato chips, confectionary, cakes, soft drinks, and flavoured milks.

BCP groups

The dietary trends present in the total NZE women were also evident within the BCP groups. Water and oil used for cooking were ranked the top two items most commonly consumed in all BCP groups. The HH group consumed water and oil for cooking more frequently than the other groups, although they were both ranked first and second, respectively in the BCP groups. They also used oil as a dressing more frequently than the other groups with this making the third top food in the HH group, compared to the eighth and ninth food item in the NN and NH groups, respectively. Oil is very energy dense, therefore it can significantly increase the caloric content of a meal (Bes-Rastrollo, 2008). This result shows that women with a high BMI consume oil more frequently compared to those with a normal BMI, therefore it is possible that the habitual intake of oil contributes to their excess body fat. Available evidence however indicates that olive oil does not contribute to weight gain. Bes-Rastrollo *et al* (2008) and (2006) found that high fat items such as oil and nuts were not associated with weight gain. Benitez-Arciniega *et al* (2012) also showed that oil consumption was not associated with BMI when adjusted for energy intake. Other factors such as the consumption of other discretionary foods may therefore contribute to a high BMI and BF% body composition.

Fruit and vegetables were commonly consumed by all BCP groups; however the NN and HH group had the highest rankings, mean daily intakes and the most individuals consuming them each day for the majority of the foods. The most commonly eaten fruits were apples and bananas. This is similar to Russell's (1999) findings who identified bananas to be the most commonly consumed fruit by New Zealanders followed by apples, oranges and stone fruit when in season. The most commonly consumed vegetables were lettuce, carrot, onion and broccoli. This is also similar to Russell's (1999) findings that carrots, tomatoes, lettuce, onions and peas were the most commonly consumed vegetables by New Zealanders per week. Seasonality differences are unlikely to explain the variances seen in consumption because participants were recruited throughout all four seasons; therefore seasonality differences were accounted for. Increasing fruit and vegetable intake may reduce the risk of overweight and obesity (He *et al.*, 2004). The comparable consumption of fruit and vegetables seen in the HH BCP group to those in the NN BCP group therefore indicates that the HH BCP group may consume other items which are energy-dense that may contribute to their increased adiposity.

The highest ranking item from the breads and cereals food group was multigrain bread, which was ranked as one of the top 10 food items consumed by all three BCP groups. This supports data indicating that in New Zealand, bread is the main contributor to carbohydrate intakes (Russell, 1999). Bread is a household staple in New Zealand and therefore is likely to contribute to the carbohydrate intakes of these women (Ni Mhurchu and Ogra, 2007).

Discretionary foods such as chocolate bars and diet soft drink were ranked the highest in the HH BCP group. These individuals consumed the largest amounts and had the most people consuming these items each day(Swinburn, 2004). Literature clearly identifies the link between SSB consumption and weight gain (Nicklas, 2001, Nikpartow *et al.*, 2012, Schulze, 2004, DiMeglio, 2000, Johnson *et al.*, 2009). A causal relationship has also been shown between diet soft drinks and a high BMI (Liebman *et al.*, 2006). Liebman *et al* (2006) found that individuals who consumed diet soft drinks were more likely to also consume fast-food and larger portion sizes and also engage in less physical activity to help facilitate a healthy body weight. As the HH BCP group were the only group to have diet soft drinks identified in their top 40 food items consumed, it is likely that these play a role in increasing energy and fat intakes thus contributing to their body composition.

Overall, the consumption of the core staples in the New Zealand diet such as bread, tea, coffee and butter were common among all of the BCP groups. The higher consumption of oil, diet soft drink and the lower consumption of fruit and vegetables may distinguish individuals in the HH BCP groups from those with lower levels of adiposity.

5.4 Dietary Patterns

Dietary patterns were identified to investigate how dietary intake affects body composition profiles. Using factor analysis, four dietary patterns were derived: 1. 'Snacking' pattern, 2. 'Energy-dense meat' pattern, 3. 'Fruit and vegetable' pattern, 4. 'Healthy' pattern. Participants were categorised into tertiles according to their dietary pattern scores. Participants in the top tertile had a diet considered to be following this pattern. Individuals with low scores on this pattern (tertile 1) were considered to be loosely associated with the pattern and not following it directly.

Dietary pattern characteristics

The four dietary patterns identified are distinctly different from each other and three of the four are similar to those found in previous studies. The 'snacking' pattern (pattern 1) is similar to the 'junk food' pattern identified by Kourlaba *et al* (2009) and the 'snacky' pattern identified by Aranceta *et al* (2003). The pattern in this study is characterised by cheese, crackers, peanut

butter, cakes, biscuits, and sweet and savoury snack foods. It is not associated with green vegetables, red meat or eggs. The other patterns identified are similar to those outlined by other studies. The 'energy-dense meat' pattern (pattern 2) is similar to the 'Western' pattern identified by Schulze (2006) or the 'empty calorie' pattern identified by Quatromani *et al* (2002). Participants following this pattern frequently consume potatoes, white and discretionary breads, all types of meat, puddings and crumbed and fried foods. They do not consume high calcium milk, yoghurt, soy products or water. The 'fruit and vegetable' pattern (pattern 3) is similar to the 'healthy' pattern identified by Suliga (2015) or Newby *et al* (2006), however both of these patterns included other items such as dairy and fruit juice, whereas the pattern in this study only loaded on fruit and vegetables. This pattern was not associated with the consumption of coffee. The 'healthy' pattern (pattern 4) was similar to the 'bean' pattern identified by Maskarinec *et al* (2000). Those following the pattern consume whole grains, fish and seafood, legumes, soy products, nuts, seeds and coconut fats. They do not consume brown breads or sweetened cereals.

Age, BMI and BF%

The 'snacking' pattern had a significant difference in the age of the NZE women between tertiles 1 and 3. Women who followed this pattern (tertile 3) were older (33.9 ± 7.9 years) compared to those who followed it less closely (tertile 1) (31.9 ± 8.7 years) (table 4.11). Women following this pattern, although not significant had a higher BMI (25.9 \pm 6.5 kg/m²) and BF% (33.7 ± 8.5%). This suggests that women who are older snack more often than younger women and this may lead to a high BMI, high BF% body composition. Conversely, Basdevant et al (1993) found those who snacked throughout the day were younger (40 ± 13) years) compared to those who did not snack (43 \pm 12 years), however they also found a positive correlation between BMI and snacking. When age, BMI and BF% were broken down into their groups, the majority of women of all ages, BMI's and BF%'s loaded in tertile 3 (table 4.12). This indicates that all NZE women consume equal amount of the foods associated with the 'snacking' pattern. In agreement with this is Piernas and Popkin (2010) who found that women from 19 to 39 years derive a larger amount of energy from snacks which was also associated with increased energy intake compared to those older or younger. This age range contains the majority of the NZE women, therefore suggests that snacking is prevalent at any age and may significantly contribute to women's energy intakes eventually leading to an increased BMI.

The 'energy-dense meat' pattern also had a significant difference in both age and BMI across all tertiles (table 4.12). This pattern had a positive association with BMI and an inverse

association with age, where younger women aged 15 to 24 years, followed this pattern more closely compared to those who were older (36 to 45 years). It also suggests that those who follow this pattern tend to be overweight compared to those who only consume a few foods associated with this pattern. No other dietary patterns showed a significant relationship between age, BMI or BF% which is consistent with other studies where no significant differences between BMI and dietary patterns at baseline were present (Newby *et al.*, 2003). Adolescents have been shown to have an increased consumption of soft drinks and fast-food, both of which similar to this dietary pattern (St-Onge *et al.*, 2003). The similar 'western' pattern has been associated with an increase in weight gain and a higher risk of being overweight as the loadings in the pattern increased (Schulze, 2006, Quatromani *et al.*, 2002). This indicates that if the young women (15 to 24 years) in our study continue to follow this pattern into adulthood, they have an increased risk of becoming overweight or obese. Interventions can be developed and targeted at these young women to improve their nutrition and decrease their risk of excess adiposity.

The lack of significant association between the other dietary patterns and age, BMI and BF% indicates that the women eat a varied diet and do not solely follow one dietary pattern. Body composition may therefore have stronger associations with other factors such as portion sizes, physical activity and food availability rather than the overall specific dietary pattern. Much literature has been reported on the relationship between portion size and increased adiposity. Rolls *et al* (2002) found larger portion sizes regardless of individual characteristics and serving methods to lead to increased energy intake. Additionally, Kral and Rolls (2004) and Berg *et al* (2009) also found increased portion sizes to contribute to weight gain.

In the 'fruit and vegetable' and the 'healthy' patterns participant reporting accuracy and social desirability bias may have influenced the lack of significant associations seen between age, BMI, BF% (table 4.11, 4.12). Reporting dietary intake retrospectively is difficult and may lead to inaccuracies between actual and reported intakes and therefore is not quantifiably precise (Thompson, 2008). When completing the NZWFFQ they may have reported what they think they should be eating, how they would like to eat, or what they think the researcher wants to see, which would contribute to the lack of association between the dietary patterns and BCP groups (Hebert *et al.*, 1997). These biases in the reported dietary data may contribute to biased results and therefore significant associations were not seen.

Micronutrient intakes

All micronutrient intakes were significantly different between the dietary patterns (tertile 3) apart from vitamin B6 and alcohol. These differences demonstrate that each of the dietary patterns has different nutrient profiles, and that individuals following each of the patterns will also have different levels of nutrient intake and adequacy.

Women following the 'snacking' pattern had moderate intakes of all nutrients compared to the other dietary patterns (table 4.13). Many food items in this pattern are nutrient poor and energy-dense, nevertheless, the more nutritious items associated with this pattern such as high calcium milk, cheese, and brown bread provide valuable nutrition to the dietary pattern to such a degree that dietary recommendations are met.

Individuals following the 'energy-dense meat' pattern are most likely to have the highest energy, protein, total and saturated fat, sugar, carbohydrate, vitamin B12, iron, calcium and zinc intakes (table 4.13). Food groups that loaded on this pattern include high fat cheese, and red, white and processed meats which are likely to contribute to the high protein and saturated fat content as well as the vitamin B12, iron and zinc intakes. It also includes white and discretionary breads, sauces, creamy dressings, crumbed and deep fried foods, sweet snack foods and fruit drinks which all contribute to the high energy, total and saturated fat and carbohydrate intakes (Swinburn, 2004). As the energy intake of this pattern is significantly larger than the other dietary patterns, it is reasonable to assume that this pattern contributes higher amounts of many other micronutrients. Having the highest energy and saturated fat intakes, it can be suggested that individuals from the HH BCP group follow this pattern most closely as they have been found to also consume a high energy and high fat diet. Kong et al (Kong et al., 2009) found a diet high in fat and carbohydrate to lead to weight gain. Although the HH BCP group had lower carbohydrate intakes, these women may consume more meat, creamy dressings and fried foods contributing to higher fat consumption. This is also supported by Murtaugh et al (2007) who found that individuals consuming a diet higher in saturated fat were associated with a higher risk of obesity.

Individuals who consumed few foods from the 'energy-dense meat' pattern (tertile one) had the lowest intakes of protein, polyunsaturated fatty acids, vitamin B12, iron and zinc. As this pattern is the only one which contains meat it is logical that this pattern contains both the highest and lowest protein, iron and zinc intakes depending on the extent to which the pattern is followed. Although this pattern does have some benefits such as the high calcium, zinc and B12 intakes, it also has some aspects such as the high saturated fat and sugar content which

are detrimental to health and can lead to increased adiposity (Swinburn, 2004). This pattern is similar to the 'empty calorie' diet identified by Quatromani *et al* (2002) which was also regarded as an energy-dense diet and associated with a 40% increase risk of being overweight when following this pattern. Individuals who loaded highly in the 'energy-dense meat' pattern in this study had a mean BMI of $26.4 \pm 6.7 \text{ kg/m}^2$ and a BF% of $34.3 \pm 9.2\%$ indicating that women following this pattern were also overweight and had a high BF%. This further confirms the patterns association with these women. Interventions targeting this dietary pattern may be beneficial to improve the weight regulation and health of these women.

The 'fruit and vegetable' pattern had the highest intakes of vitamin A, C and dietary fibre (table 4.13). This pattern may be associated with the HN BCP group who also had the highest intakes of these nutrients. Women who were in tertile 1 of this pattern had the lowest intakes of fibre, vitamin C and vitamin A. Fruit and vegetables are valuable sources of these nutrients therefore individuals consuming few foods from this pattern are likely to find it challenging to obtain sufficient amounts of these nutrients (Van Duyn and Pivonka, 2000). It is likely that women in the NH or HH BCP groups do not follow this pattern. The NH BCP group had the lowest fibre and vitamin A intakes which is also shown by Garcia (2012) who found an inverse relationship between measures of obesity and vitamin A intakes. Additionally, the HH BCP group had the lowest vitamin C intakes which also correlated with previous research showing an inverse relationship between BMI, BF% and vitamin C intakes (Johnston et al., 2007). This finding contradicts that of the top 40 food items where fruit and vegetable intakes in the HH BCP group were shown to be comparable to that of the NN BCP group. The differences seen may be due to smaller quantities of these foods being consumed by individuals in the HH BCP group. Although the overall energy intake of these individuals is high, it is likely that they consume energy-dense, nutrient-poor food items which contribute to their low levels of micronutrient intakes (Garcia, 2012). This is confirmed through their relationship with the 'energy-dense meat' pattern.

The women following the 'healthy' pattern had the highest vitamin D intakes, lowest calcium intakes and moderate intakes of all other nutrients (table 4.12). This dietary pattern loaded highly on fish and seafood as well as egg and egg dishes, which are all good vitamin D sources (Chen *et al.*, 2007). Nevertheless, this pattern contains no dairy products contributing to the lowest calcium intakes of all the identified dietary patterns. The nutrient characteristics of this pattern align with the HH, NH and NL BCP groups; therefore one group cannot be linked to this pattern. Women in New Zealand and around the world on average have low vitamin D and calcium intakes as mentioned previously and this is therefore concerning for bone health in

later life (Rockell, 2006, Fleming, 1994, Ramsdale *et al.*, 1994, University of Otago and Ministry of Health, 2011).

The 'energy-dense meat' pattern can be clearly associated with the HHBCP group while looser correlations can be made between the other patterns. It is likely that women of all ages and body compositions consume foods from the 'snacking' pattern, whereas younger overweight women were more likely to consume foods from the 'energy-dense meat' pattern and not from the 'fruit and vegetable' pattern. Given the clear relationship between younger overweight individuals with the 'energy-dense meat' pattern, dietary interventions can be developed to target women with a high BF% in an attempt to reduce body weight and improve their health outcomes.

Macronutrient intakes

Protein

The AMDR (15-25% of total energy) for protein was met within each of the dietary patterns Table 4.14). The 'healthy' pattern had the highest percentage of energy from protein (19.2 \pm 3.9%) compared to the other dietary patterns, and significant differences were also seen between the pattern tertiles. The 'healthy' pattern is associated with high protein items such as fish and seafood, legumes, eggs and soy products. This is similar to the 'bean' pattern identified by Maskarinec (2000) which has been classed as a health conscious pattern due to the inverse relationship with BMI. Increases in protein intake, especially meat alternatives have been associated with increased satiety and weight regulation (Layman, 2003, Noakes *et al.*, 2005, Te Morenga and Mann, 2012). This suggests that individuals following this pattern consume less animal products and also have a normal BMI. Women in tertile three of this pattern have a normal BMI (24.8 kg/m²) indicating that women in the NL BCP group may be associated with this pattern as they also had the highest protein intakes.

The 'energy-dense meat' pattern also had significantly different protein intakes between the tertiles. This pattern was associated with red, white and processed meat consumption. Increased red meat consumption has been associated with increased BMI due to its high energy and saturated fat content (Wang and Beydoun, 2009, Vergnaud *et al.*, 2010).

These results show that higher protein intakes from meat alternatives may attenuate weight gain and lead to a healthy body weight. The NL BCP group may follow the 'healthy' pattern as associations can be made due to their normal BMI and low BF%, showing this pattern may aid in body weight regulation.

Total and saturated fat

Total fat intakes varied between the dietary patterns. The 'snacking' and 'fruit and vegetable' patterns both met the AMDR but the 'energy-dense meat' and the 'healthy' pattern both had excessive intakes ($35.9 \pm 6.3\%$ and $36.7 \pm 7.9\%$, respectively).

Individuals following the 'fruit and vegetable' pattern had total fat intakes within the AMDR, however those who consumed only few items from this pattern had intakes exceeding the recommendations ($36.8 \pm 6.5\%$). Fruit and vegetables were the only food items which loaded highly on the pattern thus indicating that individuals who follow this pattern also consume other foods which are higher in saturated fat. Confirming this is Schulze (2006) who found individuals following a 'western' dietary pattern characterised by low intakes of fruit, vegetables and whole grains was associated with a diet of higher energy and total fat intake.

The 'energy-dense meat' pattern also had a total fat intake above the AMDR in tertile three however the difference between the tertiles was not significant. Nevertheless, individuals who loaded in tertile three of this pattern had significantly higher saturated fat intakes compared to those in tertile one. Overweight and obese individuals are more likely to consume a greater amount of saturated fat compared to those of normal weight, therefore this further confirms the association between the 'energy-dense meat' pattern and the HH BCP group (Alfieri, 1997).This is likely due to the high animal protein consumption associated with this pattern (Wang and Beydoun, 2009, Vergnaud *et al.*, 2010). Additionally, the high saturated fat items such as snack foods, fried foods, and high fat cheese associated with this pattern are likely contribute to the increased adiposity seen among these women (Swinburn, 2004).

Carbohydrate

Significant differences in carbohydrate intakes were identified between each of the tertiles in the 'healthy' pattern. Individuals who loaded in tertile one of the 'healthy' pattern had the highest carbohydrate intakes ($43.2\% \pm 5.8$). Studies have shown that whole grain consumption may contribute to a healthy body weight (Good *et al.*, 2008). It could be speculated that the NN group consume only few foods from this pattern as this group had the highest carbohydrate intakes. Decreasing the amount of energy from fat and increasing the consumption of complex carbohydrate has been shown to have a beneficial effect on body weight and body fatness (Saris *et al.*, 2000, Poppitt *et al.*, 2002). Bowman and Spence (2002) showed that adults eating greater than 55% of energy from carbohydrate were more likely to have a BMI below 25 kg/m². Shay *et al* (2012) showed that a diet higher in complex carbohydrates and lower in saturated fat and protein is associated with lower energy intakes, a

more favourable micronutrient profile and a lower BMI. This further confirms the loose association of the NN BCP group with the 'healthy' pattern. Although carbohydrate intakes for this pattern are still well below this threshold, it indicates that the higher carbohydrate intakes of the NN BCP group may be dictating their body composition through decreasing the overall consumption of total fat and energy.

5.5 Diet Quality

Women in the HH BCP group did not consume significantly different nutrient intakes compared to other groups. They met all of the EAR's except for iron and vitamin D; however this was consistent within all BCP groups. Individuals who are overweight or obese consume adequate energy intakes. Once the EAR's are met through the consumption of nutrient dense foods, the discretionary foods consumed provide excess energy which will contribute to weight gain (Garcia, 2012). Women who have poor micronutrient status may therefore make poor food choices and / or have alterations in micronutrient metabolism (Kimmons, 2006). Women with higher BF%'s may metabolise specific micronutrients via different pathways and therefore may have higher micronutrient requirements compared to those with lower adiposity (Chai, 2010).

Vitamin A

Women in the HH BCP group had vitamin A intakes which met both the EAR and RDI and were not significantly different to the other BCP groups (table 4.5) (National Health Medical Research Council, 2005). Previous studies have shown the inverse association between BMI and carotenoids to be independent from dietary intake (Chai, 2010). As the vitamin A intakes in these women appear to be sufficient from a dietary intake perspective, further research is warranted to investigate the nutritional status of these ladies from a biological perspective.

Vitamin D

Individuals with a HH body composition had one of the highest vitamin D intakes, although this still did not meet the adequate level of 5 µg per day (table 4.5) (National Health Medical Research Council, 2005). This was consistent however throughout the whole study population. Not only is vitamin D obtained through dietary means, but sunlight in the New Zealand summer months is also a main source (Rockell, 2006, National Health Medical Research Council, 2005). Vitamin D status has been strongly associated with different levels of adiposity and it was found that vitamin D concentrations in obese New Zealanders were 6 nmol/L lower than that of individuals who were of normal weight (Rockell, 2006, Cheng *et al.*, 2010). Previous studies have reported the fat soluble vitamin D to be sequestered and stored in fat cells in those with high BF%'s leading to a lower vitamin D status compared to individuals with

less body fat (Cheng *et al.*, 2010, Martini and Wood, 2006). These results suggest that another mechanism may be present as individuals of a HH body composition had vitamin D levels comparable to the rest of the NZE women. Additionally, other factors may also influence their status such as sunlight exposure, where individuals with a leaner body composition may participate in a higher amount of physical activity outdoors and therefore a greater absorption of vitamin D through the skin (Cheng *et al.*, 2010).

Iron

Women in the HH BCP group had the highest intakes of iron; however, they still did not meet the RDI (table 4.5). This BCP group was most associated with the 'energy-dense meat' pattern which contained red, white and processed meat. No other dietary pattern contained animal protein, thus explaining the highest iron intakes in this pattern. New Zealand women and those with a high BF% have routinely been shown to have low iron levels, with studies showing a higher prevalence of iron deficiency in overweight and obese women compared to their normal weight counterparts (Cepeda-Lopez *et al.*, 2011, Yanoff *et al.*, 2007). As the HH BCP group in this study consumed the highest iron intakes, it is likely that iron is metabolised by a different mechanism compared to those with a lower BF% (Cepeda-Lopez *et al.*, 2011, Yanoff *et al.*, 2007, Zimmermann *et al.*, 2008).

Vitamin E

Women in the HH BCP group had the same vitamin E intakes as the NN and NH BCP groups (table 4.5). Literature reports conflicting views on vitamin E metabolism and its status in individuals with a high BF% (Mehmetoglu *et al.*, 2011, García *et al.*, 2009). These data suggest there are no significant differences in vitamin E intakes between individuals of different body compositions, therefore different mechanisms for metabolism may be present in those with a higher BF%.

Zinc

Women in the HH BCP group had zinc intakes comparable to that of the other BCP groups. Previous research has shown those who are overweight or obese have high dietary zinc intakes, but the same zinc status as individuals with lower adiposity (Ennes Dourado Ferro, 2011). This indicates that those with excess adiposity may require a higher zinc intake compared to those with a lower BF%.

Vitamin C

Women in the HH BCP group had the lowest vitamin C intakes compared to the other groups (table 4.5). Lower dietary intakes may be responsible for the lower levels of vitamin C routinely

seen in this population with higher body fat. Garcia (2012) found a negative relationship between vitamin C status and BMI. Similar results were also seen by Johnston *et al* (2007) where vitamin C was negatively associated with BMI, BF% and waist circumference. As previously mentioned, the HH BCP group was largely associated with the 'energy-dense meat' pattern. This pattern does not load on any fruit or vegetables which are high in vitamin C and this may be a cause of the low intakes observed (Garcia, 2012). Although it does not rule out any differences in vitamin C metabolism between those of different body compositions, it does suggest that differences in food choices lead to different intakes of vitamin C and this may be a contributor to the low levels observed in these individuals. Increased vitamin C intakes in individuals with a HH body composition may be beneficial to increase their serum levels.

These results show that the intakes of micronutrients in overweight and obese individuals are not significantly different to that of normal weight women; yet previous research indicates that the concentrations seen in these women are low. Further research is warranted into the mechanisms behind the lower concentrations of these micronutrients and whether higher nutrient intakes would be beneficial for women with a higher BF%.

Chapter 6 Conclusions

6.1 Aim of the research

Body composition determines an individual's risk of many chronic diseases. Specifically, excess body fatness has been associated with a number of adverse health outcomes including hypertension, dyslipidaemia, insulin resistance, type 2 diabetes and cardiovascular disease (University of Otago and Ministry of Health, 2011, World Health Organisation, 2015, Stein, 2004, Oliveros, 2014, Poirier *et al.*, 2006, Pi-Sunyer, 2002). Body mass index is a commonly used measure to identify an individual with excess body fat (Flegal *et al.*, 2009). Many individuals however are misclassified using this measure and therefore those with excess body fat may not be identified as having increased health risks (Oliveros, 2014, Gallagher *et al.*, 2000). Habitual dietary intake is a factor which determines body composition and thus identifying eating patterns that are predictive of excess body fat in women may be of value in assessing chronic disease risk (Swinburn, 2004).

Few studies have investigated how the current dietary intakes and eating patterns of women relate to BF%, BMI and the tailoring of dietary advice according to these body fat profiles. The aim of this study therefore was to identify eating patterns of premenopausal NZE women participating in the wider women's EXPLORE study. The objectives of this sub-study were to explore the dietary intakes of the NZE women in different BCP groups, to identify eating patterns which are the strongest predictors of body fatness, and to explore the differences in the macronutrient distribution profiles and diet quality between the BCP groups and the identified eating patterns. Investigating these factors will contribute to our understanding of particular dietary intakes and eating patterns associated with different BCP groups. This will enable us to identify targeted interventions to help improve health outcomes in women with excess body fat.

This study was a sub-study of the wider cross-sectional comparative designed 'Examining Predictors Linking Obesity Related Elements' (EXPLORE) study in young adult women. The study took place at Massey University Albany Campus, North Shore Auckland. It investigated the dietary data obtained from a validated 220-item, self-administrated, semi-quantitative FFQ which was completed by 231 post-menarche, pre-menopausal NZE women. Height and weight were taken and BF% was measured using air displacement plethysmography (BodPod). Participants were categorised into one of three body composition profile (BCP) groups: normal BMI (18.5-24.9 kg/m²), normal BF% (\geq 22%, <30%) (NN); normal BMI, high BF% (\geq 30%) (NH); high BMI (\geq 25 kg/m²), high BF% (HH). Dietary intake, macronutrient profiles and diet quality,

along with dietary patterns identified using principal component factor analysis, were analysed for the total NZE population and BCP groups. Associations between dietary patterns and age, BMI and BF% were also investigated.

6.2 Main findings and Conclusions

The main findings and final conclusions of this study will be presented according to the research objectives as they were stated in chapter 1.

• The first objective was: "to explore the dietary intakes of women in the different body composition profile (BCP) groups".

The dietary intakes in the NZE women between each of the BCP groups were explored and clear differences micronutrient intakes have been found. A concerning number of NZE premenopausal women consumed inadequate intakes of iron (82%), vitamin D (55%), folate (48%), calcium (28%) and dietary fibre (28%) putting them at risk of nutrient deficiencies. Nevertheless, all nutrient intakes on average apart from iron and vitamin D within the BCP groups were adequate to meet the nutrient reference values.

The HH group had the largest intakes of energy (9296.2 kJ/d), total fat (89.4 g/d), saturated fat (36.5 g/d), monounsaturated fat (30.4 g/d), and sodium (2661.9 mg/d). This group also had the lowest intakes of vitamin C (131.6 mg), folate (408.9 μ g) and calcium (1159.5 mg) compared to the other BCP groups. These results indicate that these women may not consume as nutrient dense diets compared to women in the other BCP groups, and that their food choices may be significantly different to that of women with lower BF%'s. Strategies to reduce the energy and fat consumption in these women would be beneficial to aid in weight management.

The NN group had the lowest protein (93.4 g), B12 (4.5 μ g), iron (11.9 mg) and zinc (11.7 mg) intakes per day. These nutrients are clustered in animal protein-rich foods; therefore it is likely that these women consume less red meat compared to other BCP groups. Overall, these women consume a nutrient dense diet but still follow the trend of inadequate iron and vitamin D intakes.

The NL group had the highest PUFA (13.5 g), dietary fibre (32.9 g), folate (467.4 μ g), calcium (1409.4 mg), iron (13.1 mg), zinc (12.8 mg), and selenium (102.0 mg) intakes per day indicating that these women consume a nutrient dense diet which is reflected in their body composition of a normal BMI with a low BF%.

Common household staple items were identified as the most frequently consumed food items by the NZE women such as water, oil, coffee, tea and butter. Fruit and vegetables, bread, milk and yoghurt were also present. The HH group consumed diet soft drinks more frequently than the other groups which have been shown to increase the consumption of other food items high in energy and fat, such as chocolate bars. The HH group also consumed oil for cooking and chocolate bars more frequently than the other groups; likely increasing their energy intake and contributing to their excess adiposity.

• The second objective was: "to identify the eating patterns which are the strongest predictors of increased body fat".

The patterns which are the strongest predictors of body fat have been explored and four dietary patterns were identified: The 'snacking' pattern, 'energy-dense meat' pattern, 'fruit and vegetable' pattern, and the 'healthy' pattern.

The 'snacking' pattern was associated with women of all ages, BMI and BF%'s. This indicates that all women may consume food items from the 'snacking' pattern, such as peanut butter, sweet spreads, cakes, biscuits and sweet and savoury snack foods, regardless of their body composition. These women however are less likely to eat vegetables, red meat and egg and egg dishes.

The 'energy-dense meat' pattern was associated with younger (16-24 years) and overweight (26.4 kg/m²) women suggesting that this pattern leads to increased body fatness. This pattern was associated with the consumption of red, white, and processed meats, pudding, and crumbed and deep fried foods, but with a lower intake of high calcium milk, yoghurt, soy products and water. As a result, individuals following this pattern are also most likely to have the highest energy, protein, total and saturated fat, vitamin B12, iron, calcium and zinc intakes. Foods consumed from this pattern in conjunction with those from the 'snacking' pattern are likely to contribute to excess adiposity seen in the HH BCP group. Development of dietary strategies in the community targeting this specific dietary pattern will be beneficial to improve the health of individuals following this pattern.

The 'fruit and vegetable' pattern was associated with the HN BCP group as they had the highest vitamin A and C intakes showing that they consume plenty of fruits and vegetables but are less likely to consume coffee. This pattern was not however, associated with the NH BCP group as they had the lowest intakes of vitamin A and fibre, or the HH BCP group as they had

the lowest vitamin C levels. This indicates that women with a high BF% may have low fruit and vegetable intakes which likely contribute to their body composition.

Women following the 'healthy' pattern are likely to have the highest, although still insufficient, vitamin D intake, the highest protein intake, and the lowest calcium intake. These nutrients are obtained through whole grains, fish and seafood, legumes, nuts and seeds, but not white and brown breads, sweetened cereals or fruit juice. This pattern was linked to women in the NL BCP group who also had the highest protein intakes.

• The third objective was: "to explore the differences in the macronutrient profile intakes in women of different body compositions"

The differences in macronutrient intake between the BCP groups have been explored. NZE women of all body compositions following any dietary pattern consume protein intakes meeting the AMDR guidelines.

On average the NZE women met the total fat AMDR guideline but did not meet the guideline for saturated fat. A large proportion of NZE women (39%) consumed over the AMDR for total fat, as did the NH and HH BCP groups (35.5% and 35.3%, respectively). Additionally, a large proportion of NZE women (66%) and all of the BCP groups also consumed over the AMDR for saturated fat. High total fat intakes were also seen in the 'energy-dense meat' pattern and individuals who scored high on this pattern had significantly higher saturated fat intakes compared to those who were not. The saturated fat intake (13.1%) of women following the 'fruit and vegetable' pattern also exceeded the guidelines, indicating that women who follow this pattern may consume foods outside of this pattern that are high in saturated fat.

The average carbohydrate intake for the total population was low (41.9% of total energy intake), where 64% of the NZE women had intakes below the AMDR. The women associated with each of the BCP groups as well as the four dietary patterns also had carbohydrate intakes below the AMDR. The HH BCP group consumed significantly less carbohydrate (41.5%) compared to the NN BCP group who had the highest intakes (43.2%). As the HH BCP group exceeded the AMDR guidelines for both total fat and saturated fat, the high fat intakes seen in these women are therefore likely to be contributing to their excess adiposity, which may lead to adverse health outcomes. Individuals in the NH BCP group also consumed less carbohydrate and more total and saturated fat compared to the NN BCP group. This indicates that a higher carbohydrate and lower fat intake may be associated with a lower BMI and BF% body composition.

• The final objective was: "to explore the micronutrient intake in each BCP group and compare the differences in diet quality".

The micronutrient intakes of each BCP group were investigated in regards to diet quality. Women in the HH BCP group had vitamin A, E, and zinc intakes comparable to the other BCP groups. They also had one of the highest vitamin D and iron intakes, although not meeting the AI and RDI, respectively. The Iron and vitamin D intakes therefore indicate inadequate intakes among all NZE women and that the lower micronutrient concentrations seen in individuals with a high BF% may be independent of dietary intake, while biomarkers were not measured. Further research is therefore indicated into the metabolism of fat soluble nutrients, and whether they are sequestered in body fat leading to lower circulating plasma levels in women with a high BF%.

6.3 Study strengths

This study has several strengths which contribute to the final outcomes. They include the method of body composition analysis, using a validated FFQ, analysing seasonal eating patterns, and using dietary pattern analysis.

Body composition analysis

Using air displacement plethysmography (ADP) to determine body composition is a strength of this study. This method is emerging as the 'gold standard' of body composition analysis (Fields *et al.*, 2002, Collins and McCarthy, 2003). It provides a safe, fast and effective way to measure an individual's body composition accurately with minimal participant burden (Fields *et al.*, 2002, Lowry and Tomiyama, 2015, Ginde *et al.*, 2005, Collins and McCarthy, 2003). This highly accurate and precise assessment of body fat and lean mass was used together with dietary data to identify specific dietary patterns of the NZE women (Lowry and Tomiyama, 2015). We can therefore be confident that the BF% of these women is accurate and precise.

Validated FFQ

The FFQ has been validated and identified as a valid tool to determine the relationship between dietary intake, socio-demographic factors, and nutrition-related risk factors within a population (Houston, 2014). The FFQ included New Zealand specific foods and portion sizes making it relevant to the NZE women in assessing their dietary intakes. Misreporting of dietary intake was also accounted for in the nutrient analysis by using the Goldberg equation. This method highlights those individuals who over- or under-reported their energy intake in relation to their energy requirements in the FFQ. Individuals identified as mis-reporters were excluded from the nutrient analysis. The dietary data obtained from this study can therefore be considered a valid and reliable representation of the NZE women's dietary intakes and eating patterns.

Seasonal eating

Recruitment of participants in this study occurred from August 2013 to December 2014. This gave a variety of dietary intakes over a range of seasons and therefore reflects seasonal changes in nutrient intakes and eating patterns. This ensures the eating patterns identified are an average of the total NZE population throughout the year.

Dietary pattern analysis

Using pattern analysis to assess the dietary intakes of these women was a strength of the study. Assessing the collinearity of foods and nutrients can be effectively used to determine dietary patterns and thus habitual food intake (Sacks *et al.*, 1995). Patterns of food intake represent daily occurrences, where nutrients are consumed together rather than in isolation (Sacks *et al.*, 1995). These habitually consumed combinations of foods may determine body composition and subsequent health outcomes (McNaughton *et al.*, 2007). The dietary patterns identified in this study therefore can be used to predict relationships between dietary intake and health outcomes, allowing interventions specific to these individuals who follow the patterns to be created.

Factor analysis reliability

A way to determine the reliability of factor analysis is to calculate the Cronbach's α . Cronbach's α is a lower bound estimate of reliability and provides a measure of internal consistency (Tavakol, 2011). The reliability of factor analysis is closely related to its 'reproducibility', where the test is considered reliable if it is able to be reproduced (Margetts, 2004). Alpha is used to test if two items are correlated to one another; the higher the correlation, the higher the alpha value. Alpha values below 0.7 are considered a low reliability in the outcome (Tavakol, 2011). The Cronbach's α values in this study were low, however very close to 0.7 and this may be due to poor inter-relatedness between items or heterogeneous constructs (table 4.10). (Tavakol, 2011). Many dietary papers tend to use eigenvalues, scree plots and 'interpretability' to determine the reliability of their dietary patterns (Suliga, 2015, McNaughton *et al.*, 2007, Murtaugh *et al.*, 2007, Newby *et al.*, 2004). This study has considered all four approaches to determine reliability; therefore the dietary patterns identified were taken to be reliable estimates of dietary patterns followed by the NZE women.

6.4 Study limitations

There are a number of limitations associated with dietary pattern analysis which may have influenced the outcome of this study. These include the study population characteristics, social desirability bias, the physical activity level of the individuals, interpretability of the dietary patterns, and the lack of blood sample analysis.

Study population

A convenience sample of NZE women living in Auckland were recruited via email distribution, pamphlet advertisements and word of mouth. Women who responded were interested in health and wellbeing, therefore were motivated to participate in the study. Many of the participants were also a part of sports teams or attended the gym regularly therefore may not be representative of the general New Zealand population.

As previously mentioned, social desirability and social approval bias may influence the reporting of the NZE women. Reporting how they think they should be eating, how they would like to be eating or wanting to 'please the researcher' may bias the dietary data results. This is a factor that is present in all self-report dietary assessments. The extent to which it influences the dietary data results should be assessed and adjustments should be made accordingly (Hebert *et al.*, 1997). In an attempt to reduce this bias, the NZWFFQ has been validated and has been shown to give an accurate representation of an individual's eating habits in relation to a food diary over the last month of consumption (Houston, 2014). Additionally, as previously mentioned, participant perception of what they actually eat may provide a large bias in the results. Misreporting of energy intake was accounted for by using the Goldberg equation and individuals identified as under or over reporting their energy intakes were excluded from the nutrient analysis section of this study.

Physical activity level

A limitation the Goldberg equation is the estimation of individuals physical activity levels (PAL). The participant PAL was not known for each participant in this study. A general PAL level of 1.55 for a lightly active individual was used for all participants. This value was selected based on previous research showing that it is a conservative value (Black, 2000). If the average PAL level was higher than the 1.55 used then the proportion of under and over reporters would increase and decrease, respectively.

Principal components analysis method

A limitation of the principal component method stems from the subjective decision in determining the number of factors to extract (Newby *et al.*, 2004). Although, scree plots and eigenvalues, all provide some guidance in this decision, ultimately the final decision is open to interpretability by the researchers. The choice to include four factors in this study was based on all four elements above and was believed to be the best fit to represent the dietary patterns of the NZE women in this study.

Biomarkers of dietary intake

Blood samples were not analysed as part of this study. Although the dietary intake of individuals were determined, blood samples would have been valuable in identifying individuals of different BCP groups who were in fact deficient in certain nutrients.

6.5 Recommendations

- Dietary intakes of iron, vitamin D, folate and calcium are low in many NZE premenopausal women. Interventions and dietary advice should be given to increase the intakes of these micronutrients to prevent potential adverse health outcomes. For example, a blog could be designed outlining the importance of iron, vitamin D, folate and calcium in the daily diet. The blog could also include the current intake guidelines, food sources and recipes which provide high amounts of these nutrients. This blog could link to a free application for use during grocery shopping which provides the food sources for these nutrients and recipes from the blog.
- Women with a high BMI (≥25 kg/m²) and a high BF% (≥30%) body composition tend to consume a diet high in energy, total and saturated fat. Interventions to decrease the energy and fat consumption in these individuals will be beneficial to aid in weight management. For example, a policy could be introduced to reduce the advertisements and marketing stands of discretionary food items such as chocolate within supermarkets. These could be replaced with healthier alternatives such as fruit and vegetable stands. This will target all women who snack regularly; encouraging healthier alternatives, and also those with a HH body composition who tend to consume a diet high in energy, total and saturated fat.
- On average the NZE women do not follow the AMDR guidelines, consuming low carbohydrate and high saturated fat intakes. A visual aid, similar to the 'healthy plate

model' could be created which outlines the current AMDR guidelines and shows how individuals can easily meet this.

- All NZE women were found to frequently consume foods from the 'snacking' pattern. This
 pattern had a high saturated fat intake and contained nutrient-poor, energy-dense food
 items.
 - Dietary guidelines could be created targeting women who follow the 'snacking' pattern.
 - Focus on promoting the reduced consumption of cakes and biscuits, sweet and savoury snack foods and spreads to reduce energy, total and saturated fat intake.
 - Provide the rationale for reducing the intake of these items and include exciting healthy alternatives both for everyday consumption and social occasions.
- Overweight younger (16-45 years) women are more likely to follow the 'energy-dense meat' pattern. Dietary advice should be targeted at these women to decrease their energy intake, total fat and saturated fat intakes as well as limiting their added sugar intakes to decrease excess body fat.
 - Create dietary guidelines targeting overweight women aged 16 to 24 years who follow the 'energy-dense meat' pattern.
 - Encourage decreased consumption of white discretionary breads, processed meat, creamy dressings, pudding and fried foods.
 - Include ways to increase complex carbohydrates into their diet to replace white discretionary breads to aid in weight management.
- Individuals with a high BMI (≥25 kg/m²) and high BF% (≥30%) are likely to consume less fruit and vegetables compared to those with a NN body composition. Strategies to increase the consumption of these foods are warranted.
 - Create dietary guidelines for women with a high BMI and BF% who do not follow the 'fruit and vegetable' pattern.
 - Include information on the role of vitamin A, C and dietary fibre in the body which will provide rationale and motivation for increased consumption.
 - Emphasise the current Ministry of Health 'five plus a day' guidelines and provide clear pictures and examples of appropriate portion sizes.

- Include ways to limit total and saturated fat consumption given the high intakes seen in these women.
- Individuals with a normal BMI (18.5-24.9 kg/m²) and low BF% (<22%) body composition are more likely to be following the 'healthy' pattern. This pattern was not associated with the consumption of dairy products and had the lowest calcium intake out of all the dietary patterns.
 - Create dietary guidelines for these women following the 'healthy' pattern.
 - Focus on including calcium rich foods in their diet such as dairy products, and other sources such as tofu and bones of tinned fish.

Recommendations for further research

- Given the high fat and low carbohydrate intakes in these NZE women, further investigation should be undertaken to determine their cholesterol status and cardiovascular disease risk.
- Additional factors such as physical activity levels, socioeconomic status and food availability should be investigated to determine their relationship with body composition in these women.
- The consumption of snack foods is prevalent among all NZE women; therefore further investigation into the reasons and implications behind this snacking behaviour in regards to body composition is warranted. The newly developed S-EAT snacking tool can be used which provides information on individual's dietary habits (Philipsen, 2015). This tool could be used in conjunction with information on portion sizes and the dietary patterns obtained from the NZWFFQ, to gain a better understanding of the snack food choices of women with different body compositions.
- The dietary intakes of individuals in the HH BCP group indicate that they are consuming nutrient intakes at the same levels as their NN and NH counterparts. Further research is warranted to assess the prevalence of nutrient deficiencies in these individuals, such as vitamins A, D, E and zinc, and to explore the metabolic pathways of these micronutrients in individuals with excess adiposity.

6.6 Conclusion

Four main dietary patterns were identified among pre-menopausal women: The 'snacking' pattern, the 'energy-dense meat' pattern, the 'fruit and vegetable' pattern and the 'healthy' pattern. All NZE women frequently consume snack foods; however younger overweight women tend to follow the 'energy-dense meat' pattern more closely than other women, leading to high energy, total and saturated fat intakes. Women with a high BMI and high BF% were not associated with the 'fruit and vegetable' pattern, and women with a normal BMI and low BF% may follow the 'healthy' pattern. Additionally, New Zealand European women typically follow a low carbohydrate, high saturated fat diet, while those in the NH and HH BCP groups also had high total fat intakes. These are inconsistent with the AMDR guidelines and may contribute to the development of different body compositions. This clearly illustrates the dietary patterns that contribute to excess adiposity and provides rationale for the development of interventions targeted at women following these patterns, to reduce body fatness.

Inadequate intakes of iron, vitamin D, calcium and folate were prevalent among the NZE women and these results confirm the need for public health interventions to promote the importance and increase the consumption of these essential nutrients. Finally, vitamin A, D, E and zinc intakes among individuals with excess body fat appear to be comparable to that of women with normal body fatness. Further research should be conducted to investigate metabolic differences that explain the consistently low serum levels seen among these women.

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Appendices

Appendix A. Pre-screening health and demographic questionnaire

Women's EXPLORE study

Personal Information, Health and Demographics Questionnaire

First name:						
Family name:						
Medical Practitioner:						
Address:						
Phone:						
What is your first language?						
English						
Other						
If other, please state:						
l would like to receive a brief repor project:	t summarizing Yes □	the main findings of the				
I am willing to be contacted in future research projects within the Institute of Food, Nutrition and Human Health:						
	Yes □	No 🗆				
Health and Demographic information						
Health and De	emographic	Information				

- When was your youngest child born? __ / __ / ___ (DD/MM/YYYY)

When did your last period start? (Day / mo	nth / year)		
Are you pregnant?		Yes 🗆	No 🗆
Do you have any surgical or cosmetic in	nplants?	Yes 🗆	No 🗆
Are you currently in paid employmen	t?	Yes 🗆	No 🗆
lf yes,			
Full time		Yes 🗆	No 🗆
Part time		Yes 🗆	No 🗆
If yes, specify hours per	week:		
Describe your job or paid employment of	r work:		
TITLE / DESCRIBE WEEK		HC	URS PER
_			
Do you follow a specific diet for healt	h reasons?	Yes	□ No □
Please explain			
-			
Do you follow any diet for cultural or	religious reasons?	Yes 🗆	No 🗆
If yes, what type of diet do you follow?			
-			
Are you taking any form of medicatio medicine and contraception?	n, including traditio	onal or hor	neopathic
	Yes □	No 🗆	
Please specify the condition, the medica	ation and the dosage	in the table	e provided.

Condition	Medication	Dosage	Frequency

Are you taking any form of supplements, including tablets or drinks? Yes \hdots No \hdots

If yes, what are the name, brand and dosage of the supplements you are taking? _____

(Will send details by ema	ail Yes 🗆	No □)	
Supplement	Brand	Dosage	Frequency

Do you smoke cigarettes? Yes No No						
If yes, approximately how many ciga	arettes per day:					
Do you drink alcohol?	Yes □	No 🗆				
If yes, approximately how ma	any standard drin	ks per week:				
[1 standard drink = a glass of wine (120ml), 1 bottle/can of beer, I tot of spirits (45mL)]						
Do you have any allergies?	Yes □	No 🗆				
Please specify						

Please tell us how you found out about the Women's EXPLORE study. Did you found out from:

• A friend?

	0	If yes, what is him/her name?
• An	o en	nail list? If yes, what is the name of the email list?
• At		event? If yes, which event?
• Fly		on noticeboard? If yes, where was the noticeboard?
• Oti	her	···

Appendix B. Standard operating procedures for the food frequency questionnaire

Questionnaire Procedure

1. The questionnaires will be completed online using the computers in building 27. The two computers must first be turned on using the power button on the hard drive. You must then log in using the Explore network account, so that the computer is ready for the participants;

Username: XXXXXXXXXX Password: XXXXXXXXXX

- 2. Once logged in please ensure that there are three questionnaire links on the desktop i.e. Food Frequency Questionnaire, Eating Habits Questionnaire, and the Eating Behaviour Questionnaire. Open up each link, ensuring there is access. If there are any issues, please ask Zara, AJ, Sara, Kathryn Beck, or PC to help. If the issue cannot be resolved, there are hardcopies for each of the questionnaires which are kept in the pink Explore questionnaire folder which is will be beside the computers at the questionnaire station. However, the hardcopy version must be a last resort as this will have to be put into surveymonkey manually by an Explore staff member at a later date, which is time consuming.
- 3. The next component to check is the guidelines for the weighed food record. There is a pdf on the desktop beside the questionnaire links called 'Weighed Food Record Guidelines', open this up and make sure it plays as a video. Check the sound, if there is no sound you may have to set up the headphones from the pink Explore folder. These can be plugged into the computer hard drive in the hole marked with a headphone symbol, plug in, put the earphones in your ears and check the sound again. If there are any issues, please see one of the team members listed above. If you are going to use earphones for participants, please ensure there are small steriliser wipes at the computer station. The earphones must be cleaned by the staff member managing the questionnaire station with the wipes after each participant use. This is to maintain hygiene.
- 4. Participants will move to the questionnaire station after completing the BP station, this BP station is very quick (3-5minutes maximum) so ensure you are prepared by this stage. The participants are offered breakfast and a hot drink once BP has been recorded, if the participant declines please escort them straight to the questionnaire station. If the participant would like breakfast and/or a hot drink, please encourage them to consume this

177

at the computer stations where they will begin the questionnaires (as time may be short with multiple participants on the testing days). The questionnaire station is time consuming (25-40 minutes) so it is important to get this underway as soon as possible.

- 5. Once the participant is seated infront of the computer, inform them that there are three questionnaires to complete, and a video to watch afterwards.
- 6. The first questionnaire to complete is the Food Frequency Questionnaire; open the link for the participant. Then read the first page to the participant, taking your time through the question examples, asking at the end if they have any questions.
- 7. Click 'next' at the bottom of the page and this will take you to the first questionnaire question which asks for the participant ID number, please insert this full number yourself i.e. 250018 and click 'next' at the bottom of the page.
- 8. The questionnaire is now ready for the participant to begin. Let the participant know that you will be there to answer any questions they have whilst completing it. Leave the participant and sit at the big round table behind the computer station, so that you are close enough for any possible questions, but so that you still give the participant some privacy.
- 9. It is important that conversation is kept to a minimum whilst participants are completing the questionnaires, they need full focus.
- 10. If any questions arise during this time from the participant that you are unsure of, please ask one of the team members stated above. It is important that we obtain the most accurate information possible from the questionnaires.
- 11. If the links (questionnaires) stop working at any time, please see a team member stated above. You may need to provide a hardcopy for the participant to complete. If so, please ensure they complete this hardcopy version from the beginning, as their questionnaire data will not be saved unless they have 100% completed it.
- 12. Once the participant has completed the food frequency questionnaire, please open the 'Eating habits' questionnaire for them to complete next. Again, open up link and read

178

the initial statement(s) to the participant. Enter in their participant ID number at the top, and then remind the participant again that you are free to answer any questions they may have over this time.

- 13. If there are any complications/issues please contact one of the staff members stated, and provide a hardcopy version only as a last resort (to complete from question 1 to the end).
- 14. Once completed, open up the 'Dietary behaviours' questionnaire, entering in the participant ID again, with the instructions as above.
- 15. Upon completion of the third questionnaire, the participant must watch the food record video. Open this up for the participant, and provide cleaned earphones if required.
- 16. Once this is finished, close the video and escort the participant to the large round table for the take home package station. This package is already set up for the participant, you are required to take the participant through; the weighed food record diary, food record example booklet, allocate and record four days for them to complete this on, electronic scales are provided if required, and then through the dietary diversity questionnaire, before the participant is equipped with the accelerometer.
- 17. Firstly, show the participant the weighed food record diary and explain this must be completed over the next four days i.e. if today's date of testing is Thursday the 12th of September, then the participant will complete a weighed food diary the following four days: Friday 13th, Saturday 14th, Sunday 15th and Monday 16th from when they wake up in the morning until when they go to bed at night. They must record the time of each meal, and weigh all food items they eat providing as much detail as possible when recording foods. We need four consecutive days of recording, which include at least one weekend day. If they go out for a meal, get them to describe the food as best they can, using the palm size as a portion tool or plate/bowel size.
- 18. Provide the participant with the food record example booklet, explain they can refer the picture numbers i.e. spaghetti bolognaise picture b.3 in their food diary if this helps for meals out or those they could not weigh.

- 19. Next, record the four days and dates (as explained above) in the participant's food diary record booklet, there are four lines at the beginning of the booklet to record this. Also record this on the calendar in the Explore pink folder. Explain that there are contact details on the front of the food record dairy which the participants can use if any questions arise while they are weighing their food and recording it.
- 20. If the participant does not have a set of electronic scales at home, please provide them with some. If there are none in building 27, please see PC. Check that the scales have batteries in them, and that they work. Record the scale ID number on the back of the calendar with the participant name and date. The participant will be given a courier bag to return the scales in.
- 21. Finally, go through the Dietary Diversity Questionnaire with the participant. This is to be completed in 7 days' time when they take the accelerometer off. They only complete this questionnaire once, and emphasise how they must not look at their food record when completing this. The questionnaire is designed with a different purpose.
- 22. The weighed food record and the dietary diversity questionnaire need to be returned in the return envelope with the accelerometer after7 days. The courier bag with the scales can be sent back after the four days of recording.
- 23. Wendy will now complete the take home station setting the participant up with an accelerometer and petrol voucher.

Appendix C. Food frequency questionnaire

EXPLORE Food Frequency Questionnaire									
1. Please read carefully before you begin:									
Please make sure when	filling ou	t this quastic	oppoiro t	hat your					
Tell us what YOU usu	-			-	boldl				
Fill in the form YOURS Are correct, but don't s	SELF.		-		rioid:).				
Answer EVERY quest before moving onto the	ion; the a	sterisk symb			ning of each	n question	means th	at you mu	ist answer
This will help us to get t	This will help us to get the most accurate information about your usual food intake.								
	Please answer by ticking the box which best describes HOW OFTEN you ate or drank a particular food or drink in the LAST MONTH and HOW MUCH you would usually have.							or drink in	
For example:									
1. EXAMPLE: How	often d	lo you usu	ally h	ave sug	ar? (Plea	se do n	ot fill ou	ut)	
	Never	<1x / month	1-3x / month	1x / week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Sugar - 1 tsp	0	\bigcirc	0	0	0	0	0	0	0
If every day you have 2 cups of pancakes at dinner, you would					ugar, one bow	vi of cereal w	vith 1 tsp sug	ar and suga	ron
Adjust your portion size and fre	equency of	intake to sult yo	our eating I	nabits.					
2. EXAMPLE: How	often d	lo you usi	ually e	at bread	l? (Pleas	e do no	t fill out))	
	Never	<1x / month	1-3x / month	1x / week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Bread - 1 slice	\bigcirc	0	0	0	0	\bigcirc	0	\bigcirc	\bigcirc
If every day you have two slice times per day = '2-3x / day'.	s of toast fo	r breakfast, and	d you have	a sandwich	for lunch three	e times per w	veek, you wo	uld choose t	wo - three
Adjust your portion size and fre	umes per day = 2-5x / day. Adjust your portion size and frequency of intake to suit your eating habits.								
EXPLORE Food Frequency Questionnaire									
2. EXPLORE Study Food Frequency Questionnaire									
*1. Please enter your study ID (if you are unsure or don't know please ask the									
researcher)									
EXPLORE Food Frequency Questionnaire									
3. Eating Pattern									
*1. How would you describe your eating pattern? (Please choose one only)									
Eat a variety of all food		-	-	patteri	1: (11643	e eneo.	se one o	, , ,	
Eat eggs, dairy product				ats					
Eat eggs, dairy product									
Eat eggs and dairy proc									
Eat eggs, but avoid dai	ry products,	all meats and f	fish						
Eat dairy products, but	avold eggs,	all meats and f	fish						
Eat no animal products									

Other (please state)

Never <1x/ 1-3x/ 1x/ 2-3x/ 4-6x/ Once / 2-3x/ 4+x/ 4+ Cheddar (tasty, mild, colby) - 2 heaped Tbsp / day
250 mL/1 cup Image: control of the second secon
Milk on breakfast cereals or porridge - 125 mL / 1/2 cup Milk added to water-based hot drinks (coffee, tea) - 50 mL / 1/5 cup Milk-based hot drinks (Latte, Milo) - 250 mL / 1 cup *5. How often do you usually eat cheese? Never of the month week week day
Milk added to water-based hot drinks (coffee, tea) - 50 mL / Image: confee, tea) - 50 mL / Image: confee, tea) - 50 mL / Milkbased hot drinks (Latte, Milo) - 250 mL / 1 cup Image: confee, tea) - 50 mL / Image: confee, tea) - 50 mL / Milkbased hot drinks (Latte, Milo) - 250 mL / 1 cup Image: confee, tea) - 50 mL / Image: confee, tea) - 50 mL / Image: confee, tea) - 50 mL / Milkbased hot drinks (Latte, Milo) - 250 mL / Image: confee, tea) - 50 mL / Image: confee, tea) - 50 mL / Image: confee, tea) - 50 mL / Milkbased hot drinks (Latte, Milo) - 250 mL / Image: confee, tea) - 50 mL /
1/5 cup Milikbased hot drinks (Latte, Milo) - 250 mL / 1 cup
*5. How often do you usually eat cheese? Never <1x/
Never <1x/
Never math month week week week day day<
matchbox cube Imatchbox cube Imatch
Feta, Mozarella, Camembert - 1 heaped Tbsp / 1 med wedge Image: Commembert - 1 heaped Tbsp / 1 med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med med wedge Image: Commembert - 1 heaped Tbsp / 1 med weak Image: Commembert - 1 heaped Tbsp / 2 med weak Image: Commembert - 1 heaped Tbsp / 2 med weak Image: Commembert - 1 med weak Ima
wedge Brie, blue and other specialty cheese - 1 heaped Tbsp / (1 med wedge) Processed cheese silces - 1 silce Cream cheese - 2 heaped Tbsp Cottage or ricotta cheese - 2 heaped Tbsp Cottage or ricotta cheese - 2 heaped Tbsp Wever <1x/
1 med wedge Processed cheese silces - 1 silce Cream cheese - 2 heaped Tbsp Cottage or ricotta cheese or ricottage or ricottage or ricottage Cottage or ricottage or ricottage or ricottage or ricottag
Cream cheese - 2 heaped Tbsp Cottage or ricotta cheese - 2 heaped Tbsp * 6. How often do you usually eat these dairy based foods? Never s1x/ 1-3x/ 1x/ 2-3x/ 4-5x/ Once / 2-3x/ 4+ month month week week day
Cottage or ricotta cheese - 2 heaped Tbsp Image: Cottage or ricotta cheese - 2 heaped Tbsp Image: Cottage or ricotta cheese - 2 heaped Tbsp * 6. How often do you usually eat these dairy based foods? Never <1x/ 1-3x/ 1x/ 2-3x/ 4-6x/ Once / 2-3x/ 4-6x/ Never <1x/ 1-3x/ 1x/ 2-3x/ 4-6x/ Once / 2-3x/ 4-6x/ Ice cream - 2 scoops Image: Cottage or ricotta or dairy food - 1 pottie / ½ cup Image: Cottage or ricotta or dairy food - 1 pottie / ½ cup Image: Cottage or ricotta or dairy food - 1 pottie / ½ cup Image: Cottage or ricotta or dairy food - 1 pottie / ½ cup Image: Cottage or ricotta or dairy food - 1 pottie / ½ cup Image: Cottage or ricotta or dairy food - 1 pottie / ½ cup Image: Cottage or ricottage or ri
*6. How often do you usually eat these dairy based foods? Never <1x/
Custard or dairy food - 1 pottle / ½ cup O
Custard or dairy food - 1 pottle / ½ cup 0
Yoghurt, plain or flavour - 1 pottle / ½ cup O
Fermented or evaporated milk (buttermilk) - 15 cup

EXPLORE Food Frequency Questionnaire
5. Bread
*1. Do you eat bread?
O №
Ves
2. What type(s) of bread, rolls or toast do you eat most often? (You can choose up to 3 options, but please only choose the ones you usually have)
White
White - high fibre
Wholemeal or wheat meal
Wholegrain
Other (please state)
st 3. What type of bread slice do you usually have? (Please choose one only)
Not applicable
Sandwich siloe
O Toast slice
Mixture of both sandwich and toast slices
*4. On average, how many servings of bread do eat per day? (Please choose one
only) (A 'serving' = 1 slice of bread or 1 small roll)
Less than 1 serving
1-2 servings
3-4 servings
S-6 servings
7 or more servings
-

*5. How often do you usually eat the	ese bro	ead ba	ased f	oods	?				
	Never	<1x/ month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x day
Plain white bread - 1 slice	0	\bigcirc	\bigcirc	\bigcirc	0	0	Ó	Ó	Ó
High fibre white bread - 1 slice	0	0	Ο	0	Ο	0	Ο	\bigcirc	0
Wholemeal or wheat meal - 1 slice	0	0	0	0	0	0	0	0	0
Wholegrain bread - 1 slice	0	0	0	Ο	0	0	0	$^{\circ}$	0
Fruit bread or fruit bun - 1 slice	\odot	\odot	\odot	\odot	0	\odot	0	\odot	0
Wrap - 1 medium	0	0	0	Ο	0	0	0	0	0
Focaccia, bagel, pita, panini or other speciality breads - 1 medium	0	0	0	0	0	0	0	0	0
Paraoa Paral (fry bread) - 1 slice	\odot	\bigcirc	\bigcirc	\bigcirc	0	\bigcirc	\bigcirc	\bigcirc	0
Rewena bread - 1 silce	\odot	\odot	0	\bigcirc	\odot	\bigcirc	0	\bigcirc	0
Doughboys or Maori bread - 1 slice	\circ	$^{\circ}$	$^{\circ}$	0	$^{\circ}$	0	$^{\circ}$	\odot	0
*6. How often do you usually eat the	ese ot	her br	ead b	ased	foods	?			
	Never	<1x/ month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x day
Crumpet or muffin split - 1 crumpet / 1 whole muffin split	0	\bigcirc	0	\bigcirc	0	0	Ó	Ó	Ó
Scone - 1 medium	0	$^{\circ}$	0	\circ	0	0	0	$^{\circ}$	0
Bran muffin or savoury muffin - 1 medium	\odot	\bigcirc	0	\bigcirc	0	\bigcirc	\bigcirc	\bigcirc	0
Croissant - 1 medium	0	\bigcirc	0	0	0	0	0	$^{\circ}$	0
Waffle, pancakes or pikelets - 1 medium / 2 small	\bigcirc	\bigcirc	\bigcirc	\bigcirc	0	\bigcirc	\bigcirc	\bigcirc	0
loed buns - 1 medium	0	0	0	0	0	0	Ο	0	0
Crackers (cream crackers, cruskits, com / rice crackers, vitawheat) - 2 medium	0	0	0	0	0	0	0	0	0

st7. Do you have butter, margarine or spreads on bread or crackers?

Ο	No
\sim	

O Yes

7 or more servings

8. What type(s) do you have most often? (You can choose up to 3 options, but please
only choose the ones you usually have)
Not applicable
Butter (all varieties)
Monounsaturated fat margarine e.g. Olive, Rice Bran, Canola Oli Spreads
Polyunsaturated fat margarine e.g. Sunflower Oli Spreads
Light monounsaturated fat margarine e.g. Olivio Spread Light
Light polyunsaturated fat margarine e.g. Flora Spread Light
Plant sterol enriched margarine e.g. Pro Active, Logical Spreads
Light plant sterol enriched margarine e.g. Pro Active Spread Light
Butter and margarine blend e.g. Country Soft, Butter Lea
Parter and margame active c.g. County Con, Dater Cea
Other (please state)
Other (please state)
Other (please state) *9. On average, how many servings of butter, margarine or spreads do you have per
Other (please state) *9. On average, how many servings of butter, margarine or spreads do you have per day? (Please choose one only)
Other (please state) * 9. On average, how many servings of butter, margarine or spreads do you have per day? (Please choose one only) (A 'serving' = 1 level teaspoon or 5 mL)
Other (please state) * 9. On average, how many servings of butter, margarine or spreads do you have per day? (Please choose one only) (A 'serving' = 1 level teaspoon or 5 mL) e.g. 1 sandwich with butter thinly spread on two pieces of bread = 2 servings
Other (please state) * 9. On average, how many servings of butter, margarine or spreads do you have per day? (Please choose one only) (A 'serving' = 1 level teaspoon or 5 mL) e.g. 1 sandwich with butter thinly spread on two pieces of bread = 2 servings
Other (please state) * 9. On average, how many servings of butter, margarine or spreads do you have per day? (Please choose one only) (A 'serving' = 1 level teaspoon or 5 mL) e.g. 1 sandwich with butter thinly spread on two pieces of bread = 2 servings Not applicable Less than 1 serving

EXPLORE Food Frequency Que	stio	nnair	e						
6. Breakfast Cereals and Porridge									
 *1. Do you usually eat breakfast cere No Yes 	al and	l/or po	orridg	e?					
2. What breakfast cereal(s) do you eat	most	often	? (Yoı	u can	choos	se up i	to 3 o	ptions	s, but
please only choose the ones you usua	lly ha	ve)							
Not applicable									
Weetblx									
Refined cereals e.g. Cornflakes or Rice Bubbles									
Bran based cereals including fruity varieties e.g. Specia	l K, Mue	sil, Ali Br	an						
Sweetened e.g. Nutrigrain, Cocoa Pops									
Porridge									
Other (please state)									
week? (Please choose one only) (A 'serving' = ½ cup porridge, muesli, c e.g. ½ cup of porridge 3 times per week week Not applicable Less than 4 servings 4-6 servings 7-9 servings 10-12 servings 13-15 servings 16 or more servings	(+ 2 v	veetbi	ix 4 tii			(= 7 s	erving	gs per	
*4. How often do you usually eat por				real f	oods?	,			
	Never	<1x/	1-3x /	1x/	2-3x /	4-6x /	Once /	2-3x /	4+ x /
Porridge, rolled oats, oat bran, oat meal - ½ cup	\bigcirc	month	month	week	week	week	day	day	day
Muesil (all varieties) - ½ cup	ŏ	õ	õ	õ	õ	0	õ	0	õ
Weetbix (all varieties) - 2 weetbix	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ
Comflakes or rice bubbles - ½ cup	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ
Bran cereals (All Bran, Bran Flakes) - ½ cup	\bigcirc	0	0	\bigcirc	0	0	0	0	\bigcirc
Bran based cereais (Sultana Bran, Sultana Bran Extra) - ½ cup	Ο	Ο	Ο	Ο	Ο	Ο	Ο	Ο	0
Light and fruity cereals (Special K, Light and Tasty) - ½ cup	0	0	0	Ο	Ο	0	Ο	0	\bigcirc
Chocolate based cereals (Milo cereal, Coco Pops) - $\%$ cup	Ó	0	Ó	Ó	Ó	Ó	Ó	Ó	\bigcirc
Sweetened cereals (Nutrigrain, Fruit Loops, Honey Puffs, Frostles) - ½ cup	0	0	0	0	Ο	0	0	0	\bigcirc
Breakfast drinks (Up and Go) - Small carton / 250 mL	0	0	0	0	Ο	0	0	0	0

EXPLORE Food Frequency Qu	estio	nnai	re						
7. Starchy Foods									
*1. Do you eat any type of starchy fo No Yes	ods s	uch a	s rice	, past	a, noo	odles	and c	ousco	ous?
*2. On average, how many servings couscous do you eat per week? (Plea (A 'serving' = 1 cup cooked rice / pasta e.g. 1 cup of rice + ½ cup of pasta incl = 2.5 servings	ase ch a)	oose	one o	nly)					
Not applicable									
Less than 4 servings									
4-6 servings									
7-9 servings									
0 10-12 servings									
13–15 servings									
16 or more servings									
*3. How often do you usually eat the	se str	archy	foods	2					
· · · · · · · · · · · · · · · · · · ·	Never	<1x/	1-3x /	1x/	2-3x /	4-6x /	Once /	2-3x/	4+ x /
Rice, white - 1 cup	\bigcirc	month	month	week	week	week	day	day	day
Rice, brown or wild - 1 cup	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ
Pasta, white or wholegrain (spaghetti, vermicelli) - 1 cup	Ŏ	Ŏ	Õ	Õ	Õ	Õ	Õ	Ŏ	Ŏ
Canned spaghetti (Watties) - 1 cup	0	Ο	\bigcirc	0	\bigcirc	\bigcirc	0	0	0
Instant noodles (2 minute noodles) - 1 packet	000	0	0	-	Õ	Q	0	0	0
Egg and rice noodles (hokklen noodles, udon) - 1 cup	Õ	Õ	Õ	Õ	Õ	Q	Q	Õ	Õ
Other grain (quinoa, couscous, buigar wheat) - 1 cup	\circ	\circ	\bigcirc	\bigcirc	\circ	\circ	\circ	\bigcirc	\bigcirc

EXPLORE Food Frequency Que	stio	nnall	e						
8. Meat									
*1. Do you eat beef, mutton, hogget,	lamb,	or po	rk						
○ No									
Yes									
*2. Do you trim any excess fat (fat yo	u can	see)	off the	ese m	eats?	(Plea	se ch	oose	one
only)									
Not applicable Auror									
Always Offen									
Never cut the fat off meat									
* 3. On average, how many servings o you eat per week? (Please choose one (A 'serving' = palm size or ½ a cup of m e.g. ½ cup of savoury mince + 2 small Not applicable Less than 1 serving 1-3 servings 4-5 servings 7 or more servings	e only neat w lamb) vithou	t bon	e)		99cr,			K do
*4. How often do you usually eat mea	at?	<1X/	1-3x /	1x/	2-3x /	4-6x /	Once /	2-3x/	4+ x /
	Never	month	month	week	week	week	day	day	day
Beef mince dishes (rissoles, meatloaf, hamburger pattle) - 1 slice / patty / ½ cup	0	0	\bigcirc	0	Ο	\bigcirc	0	\bigcirc	0
Beef or veal mixed dishes (casserole, stir-fry) - $\%$ cup	0	0	0	0	0	0	0	0	0
Beef or veal (roast, chop, steak, schnitzel, corned beef) - paim size / % cup	Ο	Ο	Ο	0	Ο	0	Ο	Ο	\bigcirc
Lamb, hogget or mutton mixed dishes (stews, casserole, stirfty) - $\%$ cup	0	0	0	0	0	0	0	0	0
Lamb, hogget or mutton (roast, chops, steak) - paim size / $\%$ cup	Ο	Ο	0	0	Ο	Ο	0	Ο	\circ
Pork (roast, chop, steak) - paim size / ½ cup	0	0	0	0	0	0	0	0	0
Canned comed beef - 1 medium silce	0	0	0	0	0	0	0	0	0

*5. How often do you usually eat the	se oth	er me	eats?						
	Never	<1x/ month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Sausage, frankfurter or saveloy - 1 sausage / frankfurter/ 2 saveloys	0	\bigcirc	\bigcirc	\bigcirc	\bigcirc	\bigcirc	0	\bigcirc	0
Bacon - 2 rashers	Ο	0	0	Ο	Ο	Ο	\circ	Ο	0
Ham - 1 medium slice	0	0	0	\bigcirc	0	0	0	\bigcirc	\bigcirc
Luncheon meats or brawn - 1 slice	0	Ο	0	\bigcirc	Ο	0	0	0	0
Salami or chorizo - 1 silce / cube	0	0	0	0	0	0	0	0	0
Offal (liver, kidneys, pate) - paim size / ½ cup	0	0	Ο	\bigcirc	0	0	Ο	\bigcirc	0
Venison/game - paim size / ½ cup	0	Ο	0	Ο	0	Ο	0	Ο	\bigcirc

EXPLORE Food Frequency Que	estio	nnai	re						
9. Poultry									
 *1. Do you eat poultry e.g. chicken, to No Yes *2. Do you remove the skin from chice Not applicable 	-			oose	one o	nly)			
Always Offen Occasionally Never remove the skin from chicken									
*3. On average, how many servings of one only) (A 'serving' = palm size of chicken or ½ e.g. 1 chicken breast + 2 chicken drum Not applicable	2 cup))							
Less than1 serving 1-3 servings 4-6 servings 7 or more servings									
*4. How often do you usually eat pou	Itry?	<1x / month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Chicken legs or wings - paim size / ½ cup / 1 unit (wing, drumstick)	Ο	Ο	Ο	\bigcirc	Ο	Ο	0	0	0
Chicken breast - paim size / ½ cup / ½ breast Chicken mixed dishes (casserole, stir-fty) - paim size / ½ cup Crumbed chicken (nuggets, patties, schnitzei) - 1 medium / 4 nuggets	0000	000	000	000	000	000	000	000	000
Turkey or quall - paim size / ½ cup Mutton bird or duck - paim size / ½ cup	00	00	00	00	00	00	00	00	00

EXPLORE Food Frequency Questionnaire
10. Fish and Seafood
*1. Do you eat any type of fish or seafood? No Yes
*2. On average, how many servings of fish and seafood (all types; fresh, frozen, tinned) do you eat per week? (Please choose one only) (A 'serving' = 80 - 120g or palm size or small tin (85g)) e.g. 1 fish fillet and 1 small tin of tuna = 2 servings per week.
Not applicable Less than 1 serving 1-3 servings 4-6 servings
 7 or more servings 3. How do you normally cook / eat fish? (You can choose up to 3 options, but please only choose the ones you usually have) Not applicable
Raw / I don't cook It Oven baked / Grilled Deep fried
Shallow fry Micro waved Steamed Poached Smoked

*4. How often do you usually eat sea	rood								
	Never	<1x/ month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Canned Salmon - 1 small can (85-95g)	\bigcirc	\bigcirc	\bigcirc	\bigcirc	\bigcirc	\bigcirc	\bigcirc	\bigcirc	\bigcirc
Canned Tuna - 1 small can (85-95g)	0	0	\bigcirc	\bigcirc	$^{\circ}$	\bigcirc	0	\circ	\circ
Canned Mackerel, sardines, anchovies, herring - 1 small can (85-95g)	0	0	0	0	0	0	0	0	0
Frozen crumbed fish (patties, fillets, cakes, fingers, nuggets) - 1 medium / 4 nuggets	0	0	0	0	0	0	0	0	0
Snapper, Tarakihi, Hoki, Cod, Flounder - paim size / ½ cup	0	\bigcirc	0	\bigcirc	0	\bigcirc	0	0	\bigcirc
Gurnard, Kahawal or Trevally - palm size / ½ cup	0	0	\bigcirc	0	0	0	0	0	0
Lemon fish or Shark - palm size / ½ cup	\odot	\odot	\bigcirc	\bigcirc	\odot	\bigcirc	\odot	\odot	\odot
Tuna - paim size / ½ cup	0	0	0	0	0	0	0	0	0
Salmon, trout or eel - palm size / ½ cup	0	0	0	\bigcirc	0	0	0	0	\odot
*5. How often do you usually eat sea	food?								
	Never	<1x/ month	1-3x /	1x/	2-3x /	4-6x /	Once /	2-3x/	4+ x /
		1101101	month	week	week	week	day	day	day
Shrimp, prawn, lobster or crayfish - 1/2 cup	\bigcirc	0	month	Week	week	week	day	day	
Shrimp, prawn, lobster or crayfish - ½ cup Crab or surumi - ½ cup	00	00	Month	Week	Week	Week	day O	day O	day
	000	000	month O O	Week	Week	Week	day O O		
Crab or surumi - ½ cup	0000	0000		Week	Week O O O O	Week OOOOOOOOOOOOOOOOOOOOOOOOOOOOOOOOOOO			
Crab or surumi - ½ cup Scallops, mussels, oysters, paua or clams - ½ cup	00000	00000						ay 00000	
Crab or surum! - ½ cup Scallops, mussels, oysters, paua or clams - ½ cup Pipi or cockie - ½ cup	000000	000000					\$000000	\$000000	
Crab or surum! - ½ cup Scallops, mussels, oysters, paua or clams - ½ cup Pipi or cockle - ½ cup Kina - ½ cup	00000000	00000000					\$0000000	ay 000000000000000000000000000000000000	

EXPLORE Food Frequency Questionnaire
11. Fats and Oils
*1. Do you cook meat, chicken, fish, eggs and/or vegetables with fat or oil?
○ No
Ves
2. What type(s) do you use most often? (You can choose up to 3 options, but please
only choose the ones you usually have)
Not applicable
Butter (all varieties)
Margarines (all varieties)
Cooking oils (all varieties)
Lard, Dripping, Coconut oli, Ghee (clarified butter)
Cooking spray
Other (please state)
 * 3. When you use fat or oil to cook, how many servings of fat or oil do you use per dish? (Please choose one only) (A 'serving' = 1 level teaspoon or 5 mL) Not applicable Less than 1 serving 1 serving 2 servings 3 servings
4 servings
5 or more servings
*4. On average, how many servings of fat or oil do you use to cook per week? (Please choose one only)
Not applicable
Less than 1 serving
1-3 servings
4-7 servings
11-14 servings
15 or more servings

EXPLORE Food Frequency Qu	estio	nnai	re						
12. Eggs									
*1. Do you eat eggs?									
◯ No									
⊖ Yes									
*2. On average, not counting eggs u usually eat per week? (Please choose			ng/co	ookin	g, hov	v mar	ıy egg	s do y	/ou
Not applicable		•							
Less than 1 egg									
0 1 egg									
2 eggs									
O 3 eggs									
4 eggs									
5 or more eggs									
*3. How often do you usually eat egg	js?								
	Never	<1x/ month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Whole eggs (hard-bolled, poached, fried, mashed, omelette, scrambled) - 1 egg	0	0	\bigcirc	\bigcirc	0	0	0	0	0
Mixed egg dish (quiche, frittata, other baked egg) - 1 slice	Ο	Ο	0	Ο	Ο	Ο	Ο	Ο	0

EXPLORE Food Frequency Questionnaire 13. Legumes *1. Do you eat legumes e.g. chickpeas/dried peas, soybeans, dried/canned beans, baked beans, lentils or Dahl? ○ № O Yes *2. On average, how many servings of legumes (fresh, frozen, canned, dried) do you eat per week? (Please choose one only) (A 'serving' = 1/2 cup or 125g of cooked legumes) Not applicable C Less than 1 serving 1 serving 2 servings 3 servings 4-5 servings 6-7 servings 8 or more servings *3. How often do you usually eat these legumes? Never <13x</th> 1x 2-3x 4-6x Once 2-3x 4+x 0 Soybeans - ½ cup Tofu - ½ cup Dahl - ½ cup Canned or dried legumes, beans (baked beans, chickpeas, lentils, peas, beans) - ½ cup Hummus - 2 Tbsp

XPLORE Food Frequency Que	estio	nnai	re						
4. Vegetables									
*1. Do you eat vegetables? No Yes									
*2. On average, how many servings o per day? Do NOT include vegetable jui (A 'serving' = 1 medium potato / kumar lettuce)	ices. ra or 1	(Pleas ½ cup	se cho cook	oose	one or	ıly)	-	-	eat
e.g. 2 medium potatoes + ½ cup of pea	is = 3	servi	ngs						
Not applicable									
Less than 1 serving									
1 serving									
2 servings									
3 servings									
4 or more servings									
*3. How often do you usually eat thes	se veg	getabl	es?						
	Never	<1x/	1-3x /	1x /	2-3x /	4-6x /	Once /	2-3x/	4+ x /
Potato (bolled, mashed, baked, roasted) - 1 medium / ½ cup	\bigcirc	month	month	week	week	week	day	day	day
Pumpkin (bolled, mashed, baked, roasted) - 1/2 cup	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ
Kumara (bolled, mashed, baked, roasted) - 1 medium / ½ cup	Ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	ŏ	Ŏ
Mixed frozen vegetables - ½ cup	Ο	Ο	Ο	0	Ο	Ο	0	Ο	0
Green beans - ½ cup	\bigcirc	00000000	\bigcirc	00000000	00000000	00000000	ŏ	\bigcirc	00000000
Silver beet, spinach - ½ cup	0000000	\bigcirc	0000000	\bigcirc	\bigcirc	Ο	0	\bigcirc	0
Carrots - 1 medium / ½ cup	\bigcirc	\bigcirc	\bigcirc	\bigcirc	0	\bigcirc	0	\bigcirc	0
Sweet com - 1 medium cob / ½ cup	0	0	0	0	0	0	0	0	0
Mushrooms - 1/2 cup	\bigcirc	0	0	0	0	0	0	\bigcirc	0
Tomatoes - 1 medium / ½ cup	\bigcirc	0	0	0	\bigcirc	0	000000	000000	0
					0	0	\cap	\cap	0
Beetroot - 1 medium / ½ cup	\bigcirc	\odot	\odot	\odot	\odot	\bigcirc	\circ	\bigcirc	\odot

	Never	<1x/	1-3x /	1x/	2-3x /	4-6x /	Once /	2-3x /	4+ x /
		month	month	week	week	week	day	day	day
Green bananas (plantain) - 1 medium / ½ cup	\odot	\odot	\odot	0	\odot	\odot	0	\odot	0
Sprouts (alfalfa, mung) - ½ cup	\circ	\circ	$^{\circ}$	\odot	0	0	0	\odot	0
Pacific Island yams - 1 medium / ½ cup	\bigcirc	\odot	0	\odot	0	0	0	\bigcirc	0
Turnips, swedes, parsnip or yams - ½ cup	0	Ο	0	0	Ο	0	0	\bigcirc	0
Onions, celery or leeks - ¼ cup	0	0	0	0	0	0	0	0	0
Cauliflower, broccoil or broccoflower - ½ cup	Ο	Ο	Ο	Ο	Ο	Ο	0	Ο	0
Brussel sprouts, cabbage, red cabbage or kale - ½ cup	0	0	0	0	0	0	0	0	0
Courgette/zucchini, marrow, eggpiant, squash, kamo kamo, asparagus, cucumber - ½ cup	Ο	0	Ο	Ο	Ο	0	0	Ο	0
Capsicum (peppers) - ½ medium / ¼ cup	\bigcirc								
Avocado - % avocado	Ó	Ó	Ó	Ó	Ó	Ó	Ó	Ó	Ó
Lettuce greens (mesculin, cos, loeberg) - ½ cup	Ó	Ó	Ó	Ó	Ó	Ó	Ó	Ó	0
Other green leafy vegetables (whitioof, watercress, taro leaves, puha) - ½ cup	0	0	0	0	0	0	0	0	0

EXPLORE Food Frequency Que	estio	nnai	re							
15. Fruit										
 *1. Do you eat fruit? No Yes *2. On average, how many servings of eat per day? Do NOT include fruit juice (A 'serving' = 1 medium or 2 small piece e.g. 1 apple + 2 small apricots = 2 serving) 	e. (Ple ses of	ase c	hoose	e one	only)			l) do y	you	-
Not applicable Less than one serving 1 serving 2 servings 3 or more servings	•									
*3. How often do you usually eat the	S e fru i Never	<1x/ month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day	
Apple - 1 medium / ½ cup	0	\bigcirc	\bigcirc				\bigcirc	\bigcirc		
Pear - 1 medium / ½ cup	Õ	Õ	Õ	Õ	Õ	Õ	Õ	Õ	Õ	
Banana - 1 medium / ½ cup	\bigcirc	\bigcirc	0	0	0	0	\bigcirc	\bigcirc	\bigcirc	
Orange, mandarin, tangelo, grapefruit - 1 medium / 2 small	0	0	0	0	0	0	0	0	0	
Peach, nectarine, plum or apricot - 1 medium / ½ cup / 2 small	0	0	0	0	0	0	0	0	0	
Mango, paw-paw or persimmons / ½ cup	0	0	0	0	0	0	0	0	0	
Pineappie - ½ cup	0	0	0	0	0	0	0	0	0	
Grapes - ½ cup / 8-10 grapes	0	0	0	0	Q	0	0	0	0	
Strawberries, other berries, cherries - 1/2 cup	0	0	O	Ō	0	Q	0	0	O	
Meion (watermeion, rockmeion) - % cup	Q	Õ	000	Ŏ	00000	00000	00000	00	000000	
Kiwifruit - 1 medium / 2 small	Q	0	Q	Q	Q	Q	Q	Q	Q	
Feljoas - 1 medium / 2 smail	Õ	Õ	Õ	Õ	Õ	Õ	Õ	Õ	Õ	
Tamarillos- 1 medium / ½ cup	Q	0	Q	0	0	Õ	Q	Õ	Q	
Suitanas, raisins or currants - 1 small box	Ő	Q	Q	Q	Q	0	õ	Q	Q	
Other dried fruit (apricots, prunes, dates) - 4 pieces	\odot	\odot	\odot	\odot	\odot	\odot	\odot	\odot	\bigcirc	

EXPLORE Food Frequency Que	estio	nnai	e						
16. Drinks									
*1. On average, how many drinks do (A 'serving' = 250 mL or one cup/glass	•	iave p	er da	y? (Pl	ease	choo	se one	e only)
Less than 1 serving									
4-5 servings									
6-8 servings									
9-10 servings									
11 or more servings									
st2. How often do you usually have th	iese d								
	Never	<1x / month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Instant soup (Cup of soup) - 250 mL / 1 cup	Q	Q	0	Q	Q	0	Q	Q	Q
Fruit juice (Just Juice, Fresh-up, Charile's, Rio Gold) - 250 mL / 1 cup/glass	0	0	0	Ο	0	Ο	Ο	0	0
Fruit drink (Choice, Rio Spilce) - 250 mL / 1 cup/glass	\bigcirc	0	0	0	0	0	\bigcirc	\bigcirc	0
Vegetable juice (tomato juice, V8 juice) - 250 mL / 1 cup/glass	0	0	0	0	0	0	0	0	0
loed Tea (Lipton Ice tea) - 250 mL / 1 cup/glass	Q	Q	Q	Q	Q	Q	Q	Q	Q
Cordial or Powdered drinks (Thriftee, Raro, Vita-fresh) - 250 mL / 1 cup/glass	0	0	Ο	Ο	Ο	Ο	Ο	0	0
Low-calorie cordial - 250 mL / 1 cup/glass	0	0	0	0	0	0	0	0	0
Energy drinks small-medium can (V, Red Buli) - 250-350 mL	×	0	0	Q	0	0	Q	0	0
Energy drinks large can (Monster, Mother, Demon, large V) - 450-550 mL	0	0	0	0	0	0	0	0	0
Sugar-free Energy drinks (sugar-free V, Monster, Red Bull) - 1 small can	0	0	0	0	0	0	0	0	0
Diet soft/fizzy/carbonated drink (diet sprite) - 250 mL / 1 cup/glass	0	\bigcirc	0	0	0	0	0	0	0
Sofufizzy/carbonated drinks (Coke, Sprite) - 250 mL / 1 cup/glass	0	0	0	0	0	0	0	0	0
Sport's drinks (Gatorade, Powerade) - 1 bottle	Q	Q	0	0	Q	Q	Q	Q	Q
Flavoured water (Mizone, H2Go flavoured) - 1 bottle	0	0	0	0	0	Q	0	Q	Ő
Water (unflavoured mineral water, soda water, tap water) - 250 mL / 1 cup/glass	0	0	0	0	0	0	0	0	0

17. Dressings and Sauces

*1. How often do you usually have these dressings or sauces?									
	Never	<1x / month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Butter (all varieties) - 1 tsp	0	\bigcirc	\bigcirc	\bigcirc	0	\bigcirc	0	\bigcirc	0
Margarine (all varieties) - 1 tsp	\odot	\circ	\circ	\bigcirc	0	\odot	\circ	\circ	0
Oil (all varieties) - 1 tsp	\odot	\odot	\odot	\bigcirc	0	\odot	\odot	\bigcirc	\odot
Cream or sour cream - 1 Tbsp	0	0	0	0	Ο	0	0	0	0
Mayonnaise or creamy dressings (aioli, tartae sauce) - 1 Tbsp	0	0	0	0	0	0	0	0	0
Low fat/calorie dressing (reduced fat mayonnaise) - 1 Tbsp	Ο	\bigcirc	0	\bigcirc	Ο	Ο	0	\bigcirc	0
Salad dressing (french, Italian) - 1 Tbsp	0	\odot	\odot	\odot	0	\odot	0	\bigcirc	\odot
Sauces (tomato, BBQ, sweet chill, mint) - 1 Tbsp	0	0	0	0	0	0	0	0	0
Mustard - 1 Tbsp	\odot	\bigcirc	\odot	\bigcirc	0	0	\bigcirc	\bigcirc	\bigcirc
Soy sauce - 1 Tbsp	0	0	0	0	0	0	0	0	0
Chutney or relish - 1 Tbsp	\odot	\odot	\odot	\bigcirc	0	0	\odot	\odot	\bigcirc
Gravy homemade - ¼ cup	0	0	0	0	0	0	0	0	0
Instant Gravy (e.g. Maggi) - ¼ cup	0	0	0	0	0	0	0	0	0
White sauce/cheese sauce - ¼ cup	Ο	Ο	Ο	Ο	Ο	Ο	Ο	Ο	0

18. Miscellaneous - Cakes, Biscuits and Puddings

*1. How often do you usually eat these baked products?

	Never	<1x/	1-3x /	1x/	2-3x /	4-6x /	Once /	2-3x /	4+ x /
		month	month	week	week	week	day	day	day
Cakes, loaves, sweet muttins - 1 slice / 1 muttin	\bigcirc	\odot	\bigcirc	\bigcirc	0	\bigcirc	0	\bigcirc	0
Sweet ples or pastries, tarts, doughnuts - 1 medium	0	0	\odot	0	\odot	0	$^{\circ}$	$^{\circ}$	\circ
Other puddings or desserts - not including milk-based puddings (sticky date pudding, paviova) - ½ cup	0	0	0	0	0	0	0	0	0
Plain biscuits, cookles (Round wine, Ginger nut) - 2 biscuits	0	0	\odot	0	\odot	0	0	0	0
Fancy biscuits (chocolate, cream) - 2 biscuits	\bigcirc	\odot	0	\bigcirc	0	\bigcirc	0	\bigcirc	0

EXPLORE Food Frequency Questionnaire

19. Miscellaneous

*1. How often do you usually eat these other foods?

	Never	<1x / month	1-3x / month	1x / week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Jelly - ½ cup	0	0	0	\bigcirc	0	\bigcirc	Ó	Ó	Ó
Ice blocks - 1 Ice block	0	Ο	0	0	Ο	0	0	Ο	0
Lollies - 2 Iollies	0	0	0	\bigcirc	0	0	0	\bigcirc	\bigcirc
Chocolate - including chocolate bars (Moro bars) - 1 small bar	0	0	0	0	0	0	0	0	0
Sugar added to food and drinks - 1 level tsp	0	0	0	0	0	0	0	0	0
Jam, honey, marmalade or syrup - 1 level tsp	0	0	0	0	0	0	0	0	0
Vegemite or marmite - 1 level tsp	0	0	0	0	Q	0	0	0	0
Peanut butter or other nut spreads - 1 level Tbsp	0	00	0	0	000	0	00	\odot	0
Brazil nuts or wainuts - 2	0	0	0	O		0	0	\bigcirc	Ő
Peanuts - 10	0	0	0	0	0	0	0	0	0
Other nuts (almonds, cashew, pistachio, macadamia) - 10	0	ŏ	\odot	\bigcirc	0	\bigcirc	\bigcirc	\bigcirc	0
Seeds (pumpkin, sunflower)	0	Q	00	0	Q	00	0	0	0
Muesil bars - 1 bar	000000000000	000	0	Õ	000000	0	\bigcirc	000000	0000
Coconut cream - ¼ cup	0	0	0	00	0	0	0	0	0
Coconut milk - % cup	0	\bigcirc	\odot	0	0	0	\odot	0	0
Lite coconut milk - % cup	0	0	\bigcirc	\bigcirc	0	Q	Õ	0	0
Potato crisps, com chips, Twisties - 1/2 cup / handful	0	0	0	0	0	0	0	\bigcirc	0
*2. Do you use salt in cooking?									
O Never									
Rarely									
◯ Sometimes									
Always									
*3. Do you use salt at the table?									
· ·									
Never									
Rarely									
Sometimes									
Usually									
Always									
<u> </u>									

EXPLORE Food Frequency Questionnaire									
20. Miscellaneous - Takeaways									
*1. On average, how often do you eat	take	away	s per v	week	? (Ple	ase cl	noose	one o	only)
Less than 1 times									
1-2 times 3-4 times									
4-5 times									
*2. How often do you usually eat thes	e tak	eawa	y food	ls?					
	Never	<1x / month	1-3x / month	1x/ week	2-3x / week	4-6x / week	Once / day	2-3x / day	4+ x / day
Meat ple, sausage roll, other savourles - 1 ple / 2 small sausage rolls or savourles	0	0	0	0	0	0	0	0	0
Hot potato chips, kumara chips, french fries, wedges - $\%$ cup	0	0	0	0	0	0	0	0	0
Chinese - 1 serve	0	0	0	0	0	0	0	0	0
Indian - 1 serve	0	0	0	0	0	0	0	0	0
Thai - 1 serve	0	\odot	\odot	0	0	0	0	\odot	0
Pizza - 1 medium slice	0	0	0	0	Ο	0	0	0	0
Burgers - 1 medium burger	0	\odot	\odot	0	0	0	0	\odot	\odot
Battered fish - 1 piece	0	\circ	\circ	0	Ο	0	0	\circ	0
Fried chicken (KFC, Country fried chicken) - 1 medium piece	\bigcirc	0	\odot	\bigcirc	\odot	\bigcirc	0	\bigcirc	\odot
Bread based (Kebab, sandwiches, wraps, Pita Pit, Subway) - 1 medium	0	0	0	0	0	0	0	0	0

21. Other

* 1. Are there any other foods or drinks that you can think of that you have on a regular basis that was not covered by this questionnaire?

○ № ○ Yes

EXPLORE Food Frequency Questionnaire

22. Other

1. Please list these foods and drinks including; the serving size, and how many times per week you eat or drink these items (e.g. Pizza, 4 slices, one time per week)

*

7

Appendix D. Assumptions made when entering the New Zealand Women's Food Frequency Questionnaire (NZWFFQ).

Participants choice	Item entered into FoodWorks
Milk	
Oat milk	So Good, Soy milk
Full cream milk	4% of the amount entered as cream, the rest
	milk, whole
Lite 1.5% fat and Trim 0.5% fat	Milk, lite 1.5% fat
Whole 3.3% fat and lite 1.5% fat	Milk, whole, 3.3% fat
Not applicable/nothing chosen	Milk, whole, 0m
Spreads	
Monounsaturated fat margarine e.g. olive, rice	Margarine, Mono, 70% fat, Olivio Bertolli
bran, canola oil	Classico
Polyunsaturated fat margarine e.g. sunflower oil	Margarine, Poly, 60% fat, Sunrise
spreads	
Light monounsaturated fat margarine e.g. Olivio	Margarine, Mono, 55% fat, Olivani Light
Spread Light	
Light polyunsaturated fat margarine e.g. Flora	Margarine, Poly,50% fat, Flora Light
Spread Light	
Plant sterol enriched margarine e.g. Pro Active,	Spread, Rice bran oil, Alfa one
Logical spreads	
Light plant sterol enriched margarine e.g. Pro	Spread, Rice bran oil, Light, Alfa one
Active spread light	
Butter and margarine blend e.g. Country soft,	Dairy blend, Butter Canola
butter lea	
Not applicable/nothing chosen	Margarine, Mono, 70% fat, Olivio Bertolli
	Classico, 0m
Oils	1
Lard, dripping, coconut oil, ghee (clarified butter)	Lard
Cooking spray	Oil, composite
Not applicable/nothing chosen	Oil, composite, 0m
Butter and cooking oil	Oil, composite