Copyright is owned by the Author of the thesis. Permission is given for a copy to be downloaded by an individual for the purpose of research and private study only. The thesis may not be reproduced elsewhere without the permission of the Author.

.

ONCE DAILY MILKING IN LATE LACTATION : EFFECTS ON SOMATIC CELL COUNTS, MILK YIELD AND COMPOSITION OF DAIRY COWS WITH HIGH OR LOW SOMATIC CELLS COUNTS

> A THESIS PRESENTED IN PARTIAL FULFILMENT OF THE REQUIREMENTS FOR THE DEGREE OF MASTER OF AGRICULTURAL SCIENCE IN ANIMAL SCIENCE AT MASSEY UNIVERSITY

> > HALIMA IDD KAMOTE

DEDICATED TO MY MOTHER

,

i

· · · · · · · · · · · ·

ii

.. - .

ACKNOWLEDGEMENTS

The work reported and preparation of this thesis depended on the cooperation of many people to whom I extend my deep appreciation and sincere thanks.

Professors Colin Holmes and Duncan Mackenzie, my two supervisors, provided constructive criticism, tremendous support and advice during the course of the project.

Technical help was offered by Margaret Scott and Geoff Purchas, whereas Martin and Bruce Chesterfield took care of the animals. Dr Robert Holdaway provided a very professional, yet friendly advice on bacteriology assays. Somatic cell counting was carried out at the Livestock Improvement Corporation, Hamilton.

A huge debt of gratitude is due to Dr Patrick Morel who gave the statistical advice and so willingly spent many hours familiarising me with computing.

I feel greatly indebted to my friends Isaka Mashauri, Agnes Nyambo, the Blackwells, and fellow students Room 274, Animal Science Department, who helped ease the tension at crucial times and also made my stay in New Zealand an enjoyable experience.

Special thanks are due to Kathryn Hamilton for conscientious and skilled typing of this manuscript.

Finally, I am very grateful to the Ministry of External Relations and Trade (NZ) for their financial support, and my employer, the Principal Secretary, Ministry of Agriculture and Livestock Development (Tanzania) for granting me a leave of absence enabling me to undertake my studies at Massey University.

-

. ...

TABLE OF CONTENTS

| | | PA | GE |
|------|---------|---|-----|
| TITL | E PAGE | | . i |
| DEDI | CATION | | ii |
| ABST | RACT | | iii |
| ACKN | NOWLED | GEMENTS | v |
| TABL | E OF CC | ONTENTS | vi |
| LIST | OF TABI | LES | ix |
| LIST | OF FIGU | RES | х |
| СНАН | TER 1 | | 1 |
| 1.1 | INTROD | PUCTION | 2 |
| 1.2 | ORGAN | ISATION OF THE THESIS | 3 |
| СНАН | TER TW | O REVIEW OF THE LITERATURE | 4 |
| 2.1 | THE MA | MMARY GLAND | 5 |
| 2.2 | SIGNIFI | CANCE OF SCC | 7 |
| | 2.2.1 | SCC in relation to milk yield | 7 |
| | 2.2.2 | Relationship between SCC and milk composition | 8 |
| | 2.2.3 | SCC and processing properties of milk | 9 |
| 2.3 | SOMATI | C CELLS ORIGIN AND OCCURRENCE IN MILK | 10 |
| 2.4 | INTERPI | RETATION OF SCC | 12 |
| | 2.4.1 | Quarter sample | 12 |
| | 2.4.2 | Composite udder or cow samples | 13 |
| | 2.4.3 | Bulk, herd, milk samples | 13 |
| 2.5 | FACTOR | AS AFFECTING NUMBER OF SOMATIC CELLS IN MILK | 14 |
| ŗ | 2.5.1 | Intramammary infection | 14 |
| | 2.5.2 | Stage of lactation | 15 |
| | 2.5.3 | Age or lactational number | 15 |
| | 2.5.4 | Diurnal variation | 16 |
| | 2.5.5 | Stress related factors | 17 |
| | 2.5.6 | Management factors | 17 |

.

vi

| | | | vii |
|------|-------------------|---|-----|
| | 2.5.7 | Milking frequency | 18 |
| 2.6 | EFFECI | OF MILKING FREQUENCY ON YIELD AND | |
| | СОМРО | SITION OF MILK | 19 |
| | 2.6.1 | Three times milking | 19 |
| | 2.6.2 | Twice daily milking | 21 |
| | 2.6.3 | Once daily milking | 21 |
| 2.7 | PHYSIO | LOGICAL EXPLANATIONS FOR DIFFERENCES IN THE | |
| | EFFECI | TS OF MILKING FREQUENCY | 22 |
| | 2.7.1 | Hormonal effect | 22 |
| | 2.7.2 | Intramammary pressure | 23 |
| | 2.7.3 | Udder storage capacity | 24 |
| | 2.7.4 | Chemical inhibitor | 25 |
| | 2.7.5 | Differentiation and growth of secretory cells | 26 |
| | 2.7.6 | Extended milking intervals as an initial | |
| | | stage of involution | 27 |
| 2.8 | RATION | ALE FOR THE PRESENT STUDY | 29 |
| | | | |
| CHAF | TER 3 | MATERIALS AND METHODS | 30 |
| 3.1 | ANIMA | LS | 31 |
| 3.2 | EXPERI | MENTAL DESIGN AND PROCEDURE | 31 |
| | 3.2.1 | Determination of milk yield and composition | 32 |
| | 3.2.2 | Somatic cell counts | 33 |
| | 3.2.3 | Bacteriology analyses | 33 |
| 3.3 | STATIS | TICAL ANALYSES | 34 |
| | | | |
| CHAF | TER 4 | RESULTS | 36 |
| 4.1 | INCIDE | NCE OF INFECTION | 37 |
| 4.2 | TREATMENT PERIOD | | 37 |
| 4.3 | POST-TREATMENT 43 | | |
| 4.4 | SOMATIC CELLS | | |

| | | | viii | |
|-----------------|---|-------|------|--|
| CHAI | PTER 5 GENERAL DISCUSSION AND CONCLUSION | • • | 46 | |
| _ 5.1 | SCC - PREDICTIVE ABILITY FOR INFECTED AND | | | |
| | UNINFECTED QUARTERS | | 47 | |
| 5.2 | INCIDENCE OF INFECTION | • • • | 49 | |
| 5.3 | MILK YIELD AND COMPOSITION | ••• | 51 | |
| 5.4 | SOMATIC CELLS | | 54 | |
| | | | | |
| BIBLIOGRAPHY 58 | | | | |

ī

.

- --

LIST OF TABLES

. . . .

.

| | PA | AGE |
|---------------------------|---|-----|
| Table 3.1 | A simple chronological plan of the experiment | 31 |
| Table 4.1 | Prevalence of infection within quarters and cows in | |
| | two treatment groups at the beginning and end of the experiment | 38 |
| Table 4.2 | Daily means for milk yield and composition and somatic cells in cows with high SCC or low SCC over a 4 week treatment period during which half of the cows of each group were milked once daily and the other half | |
| 77 1.1. 4 0 | twice daily). | 39 |
| Table 4.3 | Daily means for milk yield and composition and somatic cells in cows with high SCC or low SCC over a 2 week post-treatment period during which all cows were | |
| | milked twice daily. | 44 |

1

.

LIST OF FIGURES

. . .

| Figure 4.1 | The yield of cows with a high or low initial cell count during a four week period during which half the cows from each group were milked twice daily and the other half once daily, and during a 2 week post-treatment period when all cows were milked twice daily 40 a. Milk b. Fat c. Protein d. Lactose |
|------------|---|
| Figure 4.2 | Somatic cell count of cows with high or low initial somatic cell count during a four week period during which half the cows from each group were milked twice daily and the other half once daily, and during a 2 week post-treatment period when all cows were milked twice daily |

ł

PAGE

CHAPTER 1

.

.

· · · · · ·

. . . . **.**

CHAPTER ONE

1.1 INTRODUCTION

Somatic cell count (SCC) expresses the estimated total number of 'somatic' (body) cells present in one ml of milk. Mastitis infections are characterised by an elevation in the number of somatic cells in milk, which can be used as an indirect measure of infection (Nickerson and Heald, 1982; Holdaway, 1990). SCC is widely used for monitoring udder health in dairy herds. Also, in many countries, it is used as one criterion for milk payment to producers. Thus the degree of association between SCC and prevalence of mastitis is an important parameter.

Mastitis is inflammation of the udder mainly associated with bacterial infection (Dodd, 1971). On the basis of severity of inflammation of the mammary gland, two broad categories of mastitis are recognised: clinical mastitis where physical examination of the udder or the milk reveal abnormalities, and subclinical mastitis where mammary gland inflammation exists in the absence of visible signs. The subclinical form is detected by tests such as SCC applied to the milk to detect the effects of inflammation.

Research on SCC response to infection has resulted in the recommendation by the International Dairy Federation (IDF) of 500,000 cells/ml as a threshold value for mastitis diagnosis (Tolle, 1975). However, several workers report inaccuracies in diagnosis with the IDF definition, because infections are often associated with SCC below the threshold (Berning and Shook, 1992). Elevation of SCC can also occur in response to other factors related to the cow or management (Ward and Schulz, 1972). Moreover, threshold values for SCC tend to vary depending on whether there has been a low or high incidence of udder infection in the herd (Holdaway, 1990). Nevertheless, it is certain that intramammary infection will increase SCC. For this reason SCC is a useful parameter for the detection of subclinical mastitis.

Indirect methods which determine the SCC of milk samples such as the California Mastitis Test (CMT) and Wisconsin Mastitis Test (WMT) have been available for some time, as has the direct microscopic SCC procedure (Schalm *et al.*, 1971). More recently, automated devices for rapid determination of SCC in milk samples have become available. The two most commonly used are the Coulter Milk Cell Counter which counts particles as they flow through an electric field, and the Fossomatic, which stains cells with fluorescent dye and then counts the number of fluorescing particles. Both devices are capable of rapid, inexpensive determination of SCC in large numbers of samples (Heeschen, 1975). The ability to correctly interpret these SCC data, however, depends on an understanding of various factors which may affect them, plus the interaction between their effects.

1.2 ORGANISATION OF THE THESIS

This thesis consists of five chapters. Chapter One is an introductory chapter presenting highlights of the degree of association between SCC and udder infection.

Chapter Two contains a detailed review of literature relating to the importance of SCC and infection to the dairy industry; SCC interpretation; the origin of somatic cells, mechanism of their movement into the milk, and possible factors affecting their concentration in milk; effects of milking frequency on yield and composition of milk; and physiological explanations for milking frequency response. This chapter ends by presenting the objectives of the present study.

Experimental procedures and results are given in Chapters Three and Four respectively, discussion of the results, a general conclusion and suggestions for further investigation are presented in Chapter Five.

.. . . .