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**THE INFORMATIONAL NEEDS OF WOMEN
WITH CERVICAL CANCER**

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ABSTRACT

Women diagnosed with cervical cancer are confronted with enormous physical and emotional distress. Information plays a significant role in these women's coping process, enabling them to understand their situation, make adjustment in life and erase their feelings of isolation, and stigmatization. So far, little research has been done in New Zealand to study the information need profile of women with cervical cancer, and the extent to which they are satisfied with the information. The present study offers insight into cervical cancer patients' informational needs, their level of satisfaction with information, and the current ways of information delivery in the region.

Twenty cervical cancer patients from the greater Auckland region and eleven health professionals were recruited for this evaluative study. Semi-structured face-to-face interviews were used to collect data from both patients' and health professionals' perspectives. Results show areas of high informational needs among the participants, and suggest that emphasis should be placed on providing sufficient information in the areas with deficits, such as the causes of cancer, life expectancy, changes in sexual function, how other people live with the disease, and where to get information when needed. Evaluating the participants' use of informational sources reveals that they obtained information from a variety of sources. Some information services, such as patient counseling and support services, the Cancer Society and the oncology district nursing services were helpful but under-utilized by the patients.

Overall, the study shows that most patient participants were satisfied with the information provided by health services in the region. However, the existence of gaps between information needs and satisfaction in certain areas, and the reports of multiple barriers in patients' access to information, indicate that the informational needs of women with cervical cancer had not been adequately met. The findings stress the importance of individual assessment, care coordination and close collaboration among service providers. To provide patients with easily accessible, clearly defined, and readily available information was a recurrent theme in the participants' recommendations to information services in the region.

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TABLE OF CONTENTS

	Page
Acknowledgements	iii
Chapter 1: Introduction	1
1.1 Background of the study	2
1.2 Aim of the study	5
1.3 Structure of the thesis	7
Chapter 2: Literature Review	9
2.1 Patients' informational needs	9
2.2 Sources of information	15
2.3 Barriers to information	16
Chapter 3: Method	19
3.1 Research questions	19
3.2 Participant selection	20
3.3 Data collection	21
3.4 Interview schedule for patient participants	23
3.5 Interview schedule for health professional participants	24
3.6 Data analysis	26
Chapter 4: Results I: Patient Participant Interviews	27
4.1 Demographic characteristics	27
4.2 Informational needs and levels of satisfaction	30
4.3 Sources of information and degrees of helpfulness	34
4.4 Barriers against patients' access to information	38
4.4.1 Participants' knowledge of the informational sources	38
4.4.2 Participants' access to information	40
4.4.3 Difficulties in asking questions	41
4.4.4 Difficulties in understanding the information provided	44
4.4.5 Inconsistent and inaccurate advice to the participants	46
4.4.6 Special needs of new immigrants or people who know little English	46
4.5 Recommendations to the informational services	47

Chapter 5: Results II: Health Professional Interviews	53
5.1 Key informant information	53
5.2 Patients' informational needs	54
5.3 Sources of information for patients	59
5.3.1 Sources within hospital	59
5.3.2 Sources in community	60
5.4 Barriers against patients' access to information	61
5.4.1 Patients' knowledge of the informational sources	61
5.4.2 Patients' access to information	62
5.4.3 Patients' difficulties in asking questions	63
5.4.4 Patients' ability to understand and communicate with health providers	65
5.4.5 Inaccurate and inconsistent advice to patients	67
5.4.6 Barriers against Maori patients' access to information	68
5.5 Recommendations to the informational services	71
Chapter 6: Discussion	74
6.1 Informational needs	74
6.2 Sources of information	76
6.3 Barriers to information	80
6.4 Participants' recommendations	84
6.5 Limitations of the study	86
Chapter 7: Conclusion	89
Appendices:	92
Appendix I: Interview schedule for patient participants	93
Appendix II: Interview schedule for health professional participants	99
Appendix III: Patient participant information sheet	104
Appendix IV: Health professional participant information sheet	107
Appendix V: Participant consent form	110
Appendix VI: Binomial test of proportions	112
References	113

LIST OF FIGURES

	Page
Figure 4.1: Age distribution of patient participants	28
Figure 4.2: Ethnic origin of patient participants	28
Figure 4.3: Education level of patient participants	29
Figure 4.4: Patient participants' overall level of satisfaction	33
Figure 4.5: Sources of information and degrees of helpfulness	36

LIST OF TABLES

	Page
Table 4.1: Patient participants' informational needs and levels of satisfaction	32
Table 4.2: Patient participants' additional information needs and levels of satisfaction	33
Table 4.3: Sources of information and degrees of helpfulness	35
Table 5.1: Key informant information	54
Table 5.2: Information cervical cancer patients requested frequently from key informants (Responses to the 16 items listed in the interview schedule)	55
Table 5.3: Information cervical cancer patients requested frequently from key informants (Additional information supplied by key informants)	56
Table 5.4: Information that the Oncology specialists frequently provided to cervical cancer patients	57
Table 5.5: Information that the nurses in hospital frequently provided to cervical cancer patients	57
Table 5.6: Information that the nurses in community frequently provided to cervical cancer patients	57
Table 5.7: Information that the patient counselors frequently provided to cervical cancer patients	58
Table 5.8: Information that the Maori health workers frequently provided to cervical cancer patients	58