### POPULATION-BASED STAGE, TREATMENT AND OUTCOMES FOR PATIENTS DIAGNOSED WITH BREAST CANCER IN BRITISH COLUMBIA IN 2002

A. Davidson<sup>1</sup>, S. Chia <sup>2-3</sup>, R. Olson<sup>4</sup>, A. Nichol<sup>6</sup>, C. Speers<sup>2</sup>, A. Coldman<sup>5</sup>, C. Bajdik<sup>3</sup>, R. Woods<sup>2</sup>, and S. Tyldesley <sup>2,6</sup>.

**Background:** The purpose of this study was to describe the stage, treatment, and outcomes at a population level for patients with breast cancer in British Columbia (BC). **Methods:** All incident breast cancer cases registered with Breast Cancer in 2002 were reviewed for information on stage, treatment with primary local surgery, chemotherapy (CT), hormone therapy (HT), and radiotherapy (RT) and outcomes were derived from cancer centre databases for patients referred to provincial cancer centres (85% of cases). For non-referred cases, stage was estimated from manual review of pathology reports available in the provincial tumour registry. Kaplan-Meier curves for BCSS and OS were calculated by stage.

**Results:** There were 2,927 incident cases of breast cancer identified in 2002. Stage distribution was: 0 (*in situ*): 15%, I: 38%, II: 32%, III: 8%, IV: 4% and unknown 3%. Age distribution was: <40: 4%, 40's: 18%, 50's: 25%, 60's: 23%, 70's: 20% and ≥80 years old: 10%. The treatments delivered within 1 year of diagnosis were: RT-56%, CT-32% and HT-57%. The 10-year BCSS rates were for stage: 0: 99.5%, I: 95%, II: 81%, III: 55%, and IV: 4% The 10-year OS rates were, for stage: 0: 89%, I: 81%, II: 68%, III: 43%, and IV: 2%.

**Interpretation:** This analysis provides a Canadian benchmark for treatment rates and 10 year outcomes by stage for all incident cases of breast Cancer. Outcomes in BC compare well to published rates in the USA and Europe.

#### INTRODUCTION

Outcomes for breast cancer patients have improved considerably in recent years. Although breast cancer remains the most common cancer and the second leading cause of cancer deaths in women, the 5-year relative survival rate rose from 75% in the late 1970's to 90% by 2006 (1). Significant factors potentially contributing to these improvements include the availability of breast screening (2-4) and the increasingly multidisciplinary nature of cancer care, appropriate surgery, radiation treatment (5-7), and systemic therapy (8,9). Despite these encouraging improvements in breast cancer outcomes, disparities still exist within and among populations (10-12). Reasons for these differences relate in part to variation in utilization of screening, diagnostic, and treatment modalities. In order to optimally benefit from important advances, an effective provider of cancer care services must ensure that individuals in a given population have equal access to these modalities.

British Columbia has been identified as a population having one of the best breast cancer survival rates in the world (10). There are many potential factors accounting for the favourable outcome. BC is part of a country with a strong economy and a fully publicly funded cancer care system. BC organized the first public screening mammogram program in Canada, which currently screens approximately 50% of its target population (13). Mammography screening is available within the provincial health plan to BC women, free of charge, through the Screening Mammography Program of BC (SMPBC). Coldman and colleagues have previously reported that SMPBC participation is associated with a lower rate of chemotherapy use and an increased rate of breast-conserving surgery (14). BC also has a centralized organized cancer care program through the BC Cancer Agency (BCCA), which provides, for free, all radiotherapy and provides, for free, all chemotherapy and hormone therapy that is

prescribed in the province. Although there are significant geographic issues within the province, which cause some regional variation in access (15), the referral rate for a breast cancer patient to a BCCA centre for the province as a whole was 85% throughout the 2000's.

For many decades the BCCA has developed treatment guidelines and disseminated them to all the physicians in the province and is often the first province in Canada to implement new therapeutic agents and regimens. Due to the quality of the outcomes, treatment and stage data, outcomes for patients with breast cancer in British Columbia have been used in international studies assessing prognostic information and to validate prognostic models used in clinics internationally (16) but there has been no comprehensive paper describing outcomes and treatment rates with long term followup. With these considerations in mind, the primary objective of this study was to describe the patient characteristics, stage distribution, stage-specific treatment utilization and outcomes at a population level for all patients diagnosed with breast cancer in BC in 2002. Our second objective was to compare the stage distribution and survival by stage for breast cancer patients in BC to published international reports from American and European databases, where available.

### METHODS

The BCCA has a mandate to deliver cancer care services to the culturally, economically, and geographically heterogeneous population of BC (17). There is a legal requirement to send all pathology reports with a neoplastic diagnosis to the British Columbia Cancer Registry, which thereby captures all incident cases of breast cancer. Death and cause of death information are collected by Vital Statistics. All radiotherapy was provided at one of the four BCCA cancer centres, which are the only providers of

RT in the province, in 2002. All funded anti-cancer drugs are reimbursed by the BCCA, and each drug, dose, and dispensing date is recorded in the BCCA pharmacy data repository since 1998. This information is available to the Breast Cancer Outcomes Unit (BCOU) and its periodic review is an important index of performance for the delivery of breast cancer care to residents of British Columbia.

All incident patients with breast cancer diagnosed between January 1<sup>st</sup> and December 31<sup>st</sup> of 2002 were identified from the BC Cancer Registry. Cases were then linked to radiotherapy records and to the BCCA pharmacy data repository. Pre-treatment prognostic factors such as grade, stage, lymphatic and vascular invasion, estrogen receptor, tumour size, and nodal status, as well as primary surgical therapies, were collected prospectively in the BCOU for the cases referred to the BCCA.

For cases not referred to the BCCA, registry pathology records were reviewed to determine the grade, ER, tumour size, nodal status, and definitive local and regional surgery. Stage was determined for these cases based on the pathology reports. Utilization of radiotherapy (RT), chemotherapy (CT), and hormone therapy (HT) within one year of diagnosis were extracted by stage. For cases not referred to the BCCA, systemic therapy dispensed at other institutions is still captured by the BCCA pharmacy database.

Patients diagnosed with breast cancer in BC during the 2002 year were matched to the SMPBC records and screening information was extracted, including whether or not the cancer was screen-detected. The definition of a screen-detected cancer was a breast cancer diagnosis within one year of an abnormal screen. Patients were considered attendees of the SMPBC if they had a screening mammogram result listed in SMPBC

records. Those who had been screened within the 30 months prior to their diagnosis were considered active attendees.

Overall and disease-specific survivals were calculated using the Kaplan-Meier method. This study was approved by the University of British Columbia BCCA Research Ethics Board.

#### RESULTS

#### Patient Characteristics and Stage Distribution

In 2002, there were 2,927 incident cases of breast cancer diagnosed in the province of British Columbia. Figure 1 demonstrates the stage distribution and Table 1 characterizes the patient population by stage of disease at presentation. The majority of cases, 82%, represented invasive disease, whereas 14% were in situ and 3% unknown. Approximately 70% of tumors were found to be either stage I or stage II at diagnosis. The stage distribution is compared with that of the United States Surveillance Epidemiology and End Results (SEER) in Table 2 (18).

The vast majority (86%) of breast cancers were diagnosed in patients between the ages of 40 and 79. Overall, the median age at diagnosis for all stages was 61, with only 5% of patients younger than 40 and only 10% older than 80. Elderly patients formed a greater percentage of those presenting with more advanced disease than with either in situ or early-stage disease.

Over 50% of all breast cancer patients were attendees of the SMPBC, and the majority of all patients diagnosed with in situ (71%) or stage I disease (63%), had been screened within the 30 months prior to their diagnosis. In contrast, most patients diagnosed with stage II-IV breast cancer were not attendees of the SMPBC. Only 46% of stage II, 35%

 of stage III patients and 27% of stage IV patients had been screened within the 30 months prior to their diagnosis. Of all breast cancers diagnosed, 33% were considered "screen-detected", though this designation applied to 62% of cancers found in the 1,574 breast cancer patients who were attendees of the provincial screening program.

#### **Treatment Characteristics**

The majority of patients with disease ranging from in situ to stage III underwent surgery for their breast cancer (Table 3). Approximately two-thirds of patients with in situ disease underwent breast-conserving surgery (BCS) but only one-third received radiation therapy. Few in situ patients received hormonal treatment within one year of diagnosis (22%)

For patients in the stage I category, again, approximately two-thirds received breastconserving surgery (BCS), with the remainder treated with mastectomy. Of stage I and II patients treated with BCS, 88% also received radiotherapy within 1 year of diagnosis. Patients with stage II disease at presentation were equally likely to undergo mastectomy or BCS, whereas two-thirds of stage III patients underwent mastectomy as their initial surgery. Though the percentage of patients undergoing BCS as the initial surgery decreased from stage I to stage III (from 65% to 15%), the percentage of patients receiving radiation therapy within a year of diagnosis increased (from 59% to 77%). As would perhaps be expected, the percentage of patients that received chemotherapy increased with increasing stage (from 14% of stage I patients to 71% of stage III patients). Approximately 65% of patients with stage I-III cancer were treated with hormonal therapy. When only estrogen receptor-positive (ER+) patients are considered, there is little change to the in situ treatment rate, but 73% of stage I patients received hormonal treatment compared with 86% of stage III and 81% of the stage IV patients.

Models of optimal radiotherapy utilization have been developed (19-20), and Table 4 demonstrates a comparison between the BCCA data from the 2002 cohort and these published ideal utilization rates. The actual BCCA RT rate at 5 years of 59% for all disease stages compares well with the estimated ideal rate of 66% from a Canadian model but is lower than the estimated ideal rate from Australia (83%) (19,20). The difference between actual and ideal rates likely relates to differences in patient preferences are dealt with in the models (19,20). Ideal utilization rates have also been published for chemotherapy (21), and chemotherapy use for all stages is lower in the BCCA 2002 cohort than the ideal published rates, most notably for those patients with stage I disease but also for patients with stage III breast cancer. Endocrine therapy use in BC, however, exceeds the published ideal rate for all stages (22).

We have previously described differences in stage distribution and frequency of types of breast cancer treatments depending on population density within the province (15).

#### **Patient Outcomes**

Disease-specific survival (DSS) and overall survival (OS), for all stages and divided by stage, for breast cancer patients diagnosed in British Columbia in the year 2002 are shown in Figure 2. The 5 year DSS for all stages combined in 2002 was 89% (95% CI: 88 - 90%) and the OS was 83% (95% CI: 81 - 84%). The 10 year DSS was 83.8% (95% CI: 82.4 – 85.2%) and the OS was 70.6% (95% CI: 68.8 - 72.4%).

Initiatives have been undertaken in other parts of the world to evaluate the effectiveness of healthcare service delivery for patients with cancer in their respective countries, and data have been published from these cancer registries {10-12, 22, 23}. In Table 5, relative survival data for breast cancer patients diagnosed in a similar time period in other countries were compared with the British Columbia relative survival rates for the 2002 cohort. The 5-year relative survival rate for the entire cohort of patients diagnosed with breast cancer in BC in 2002 was 90% (95% CI 88 - 91%), which numerically is higher than, or equivalent to, many European countries and similar to the relative survival rates in the United States SEER databases.

#### INTERPRETATION

Outcomes for patients with breast cancer in British Columbia have been used in international studies assessing prognostic information and to validate prognostic models used in clinics throughout the world. This descriptive study of 2,927 breast cancer patients diagnosed in BC in the year 2002 demonstrates a stage distribution heavily weighted toward early-stage disease, particularly stages I and II. Most early-stage cancers were diagnosed in patients aged 40 to 79 years old. The case mix presented here is similar to that reported by the NCCN, a large national database in the United States (18).

Over 60% of in situ and stage I diagnoses occurred in patients who were attendees of the Screening Mammography Program of British Columbia, whereas the majority of stage III and IV diagnoses occurred in patients who had not been screened. In patients attending the SMPBC, most cancers were screen-detected, including over 70% of in situ and stage I cancers. Most patients with early-stage disease underwent breastconserving surgery, and adjuvant radiation therapy, whereas most patients with stage III breast cancer were treated with mastectomy and adjuvant radiation therapy. Use of both chemotherapy and endocrine therapy increased with increasing stage of disease, up to stage III. Use of radiation therapy and chemotherapy fall slightly below published ideal utilization rates, based on reviews of existing international guidelines, but compare well with other international jurisdictions. Use of endocrine therapy in BC exceeds calculated ideal utilization rates.

In recent years, data has been published from cancer registries in several regions of the world and survival rates in British Columbia are comparable to those reported from other regions such as the US, Europe, and Australia. These findings suggest that the BCCA is meeting its objective of providing timely, evidence-based cancer care services to residents of this province in the context of a widely accessible healthcare system.

Despite the comprehensive nature of the BCCA records, this report has limitations which bear consideration. As mentioned, the referral rate to the BCCA was 85%, so there was not complete data available for all breast cancer patients treated in BC but attempts were made through pathology record review in the BC Cancer Registry to garner information about non-referred cases. In 2002, HER2/*neu* status, an important prognostic and predictive indicator, was not routinely measured in early breast cancer patients because the evidence for trastuzumab (anti-HER 2 antibody) efficacy in adjuvant therapy had yet to emerge. Finally, treatments were recorded as being given, when patients received even one dose of chemotherapy or radiotherapy and a first prescription for endocrine therapy within one year of diagnosis, and do not reflect completion of systemic therapy. The non-adherence rate in British Columbia for adjuvant endocrine therapy has been reported as 40%, but compliance with

chemotherapy and radiotherapy have not been reported (24). Compliance, if different between populations, has the potential to affect comparative patient outcomes, but the primary focus of this report was the description of patient characteristics, treatment, and outcomes in BC.

In conclusion, this report indicates that breast cancer survival rates in BC are comparable to those reported in the literature from other regions of the world. The majority of patients diagnosed with early-stage disease were treated with breast-conserving surgery, as well as adjuvant radiotherapy and hormonal treatment. Continued data collection and periodic reviews are important to ensure that as breast cancer therapy evolves and its delivery to such a diverse population becomes more complex, the publicly funded provincial healthcare system is able to meet the challenges of universal access, and that the subsequent outcomes are comparable to those of any developed region of the world. The favourable results in British Columbia should serve as a benchmark for the rest of the provinces in Canada, and demonstrates the results that can be achieved with a centralized and comprehensive provincial cancer care program.

## REFERENCES

1. Siegel R, Ward E, Brawley O *et al.*: Cancer statistics, 2011: The impact of eliminating socioeconomic and racial disparities on premature cancer deaths. *CA Cancer J Clin* 61:212-236, 2011.

2. Olivotto IA, Mates D, Kan L *et al.*: Prognosis, treatment, and recurrence of breast cancer for women attending or not attending the Screening Mammography Program of British Columbia. *Breast Cancer Res Treat* 54:73-81, 1999.

3. Schopper D, de Wolf C: How effective are breast cancer screening programmes by mammography? Review of the current evidence. *Eur J Cancer* 45: 1916-1923, 2009.

4. Christensen LH, Engolm G, Cortes R *et al.*: Reduced mortality for women with mammography-detected breast cancer in east Denmark and south Sweden. *Eur J Cancer* 43:2773-2780, 2006.

5. Clarke M, Collins R, Darby S *et al.*: Effects of radiotherapy and of the extent of surgery for early breast cancer on local recurrence and 15-year survival: an overview of the randomised trials. *Lancet* 366:2087-2106, 2005.

6. Dragun AE, Huang B, Tucker TC *et al.*: Disparities in the application of adjuvant radiotherapy after breast-conserving therapy for early stage breast cancer: Impact on overall survival. *Cancer* 117(12): 2590-2598, 2011.

7. Martinez SR, Tseng WH, Carter RJ *et al.*: Do radiation use disparities influence survival in patients with advanced breast cancer? *Cancer* Jun 20. doi: 10.1002/cncr.26231. [Epub ahead of print]

8. Early Breast Cancer Trialists' Collaborative Group (EBCTCG): Effects of chemotherapy and hormonal therapy for early breast cancer on recurrence and 15-year survival: an overview of the randomised trials. *Lancet* 365:1687-1717, 2005.

9. Chia SK, Speers CH, D'yachkova Y *et al.*: The impact of new chemotherapeutic and hormonal agents on survival in a population-based cohort of women with metastatic breast cancer. *Cancer* 110:973-979, 2007.

10. Coleman MP, Forman D, Bryant H *et al.*: Cancer survival in Autralia, Canada, Denmark, Norway, Sweden, and the UK, 1995-2007 (the International Cancer Benchmarking Partnership): an analysis of population-based cancer registry data. *Lancet* 377:127-138, 2011.

11. Verdecchia A, Francisci S, Brenner H *et al.*: Recent cancer survival in Europe: a 2000-2002 period analysis of EUROCARE-4 data. *Lancet Oncol* 8:784-796, 2007.

12. Coleman MP, Quaresma M, Berrino F *et al.*: Cancer survival in five continents: a worldwide population-based study (CONCORD). *Lancet Oncol* 9:730-756, 2008.

13. BC Cancer Agency Screening Mammography Program 2012 Annual Report. http://www.smpbc.ca/NR/rdonlyres/3AF15A7A-B9D4-43C2-A93A-230A634E94AC/60300/SMP\_2012AR\_WEB.pdf (Accessed on line Nov 14, 2012).

14. Coldman AJ, Phillips N, and Speers C: A retrospective study of the effect of participation in screening mammography on the use of chemotherapy and breast conserving surgery. *Int J Cancer* 120:2185-2190, 2007.

15 Olson RA, Nichol A, Caron NR, Olivotto IA, Speers C, Chia S, Davidson A, Coldman A, Bajdik C, Tyldesley S. Effect of community population size on breast cancer screening, stage distribution, treatment use and outcomes Can J Public Health. 2012 Jan-Feb;103(1):46-52.

16. Olivotto IA, Bajdik CD. ADJUVANT! – A validated, web-based, systemic therapy decision aid for early-stage breast cancer. *Am J Oncol Review* 2005;4(Suppl 13):29-31.

17. BC Stats. www.bcstats.gov.bc.ca (accessed July 10, 2011) For Peer Review Only 18. Gloeckler Ries LA, Eisner MP. SEER Cancer Monograph: Cancer Survival Among Adults, Chapter 13: Cancer of the Breast. (accessed on line Nov 14, 2012) http://seer.cancer.gov/publications/survival/surv\_toc.pdf

19. Foroudi F, Tyldesley S, Walker H et al. An evidence-based estimate of appropriate radiotherapy utilization rate for breast cancer. Int J Rad Onc Biol Phys 2002; 53(5): 1240-1253.

20. Delaney F, Barbon M, Jacob S: Estimation of an optimal radiotherapy utilization rate for breast carcinoma. *Cancer* 98:1977-86, 2003.

21. Ng W, Delaney GP, Jacob S, Barton MB: Estimation of optimal chemotherapy utilization rate for breast cancer: setting an evidence-based benchmark for the bestquality cancer care. *Eur J Cancer* 2010; 46: 703-712.

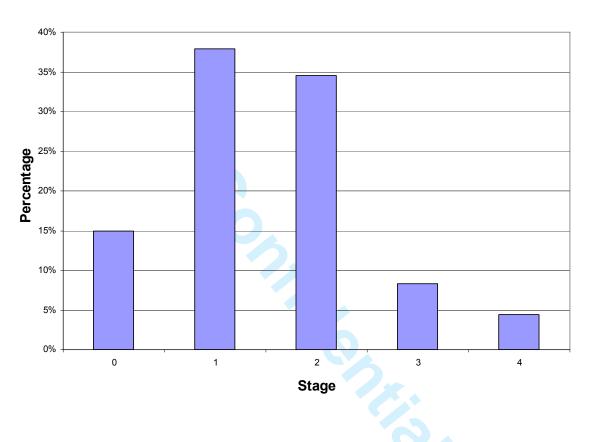
22. Fong A, Ng W, Barton MB, Delaney GP: Estimation of an evidence-based benchmark for the optimal endocrine therapy utilization rate in breast cancer. *Breast* 2010; 19: 345-349.

23. Gondos A, Volker A, Holleczek B *et al.*: Cancer survival in Germany and the United States at the beginning of the 21<sup>st</sup> century: An up-to-date comparison by period analysis. *Int J Cancer* 121:395-400, 2007.

24. Chan A, Speers C, OReilly S, Pickering R, Chia SK. Adjerence of adjuvant hormone therapies in post-menopausal hormone receptor positive (HR+) early stage breast cancer: A Population Based study form British Columbia. Cancer Research 2009; 69(4): 494s.

# **TABLES & FIGURES**





# Stage Distribution percentage

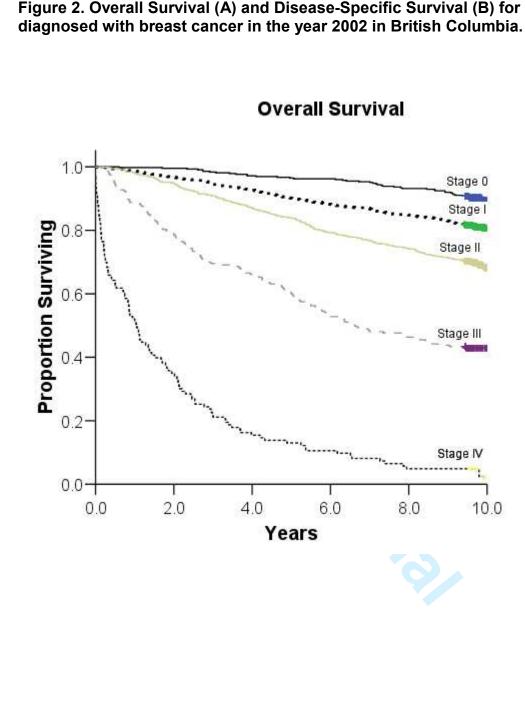
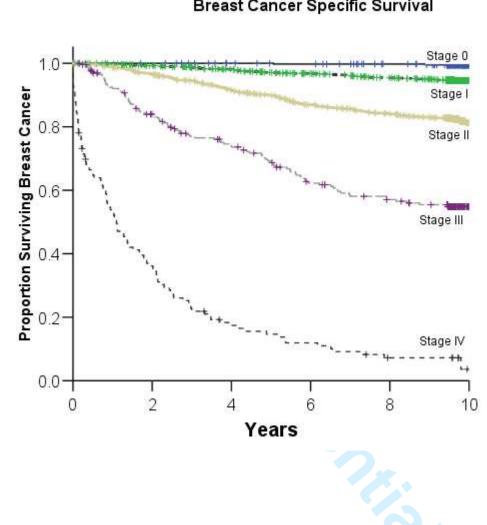


Figure 2. Overall Survival (A) and Disease-Specific Survival (B) for patients



# **Breast Cancer Specific Survival**

Table 1.	<b>Population</b>	<b>Characteristics</b>
----------	-------------------	------------------------

	All Stages		Sta	ge 0	Sta	ge I	Sta	ige II	Stag	ge III	Stag	ge IV		age nown
	n	%	n	%	n	%	n	%	n	%	n	%	n	%
Candar	2927	100	424	14.5	1118	38.2	938	32.0	233	8.0	123	4.2	91	3.1
Gender	2000	000/	400	1000/	1110	1000/	000	4000/	220	0.00/	100	000/	07	0.00/
Female	2909	99%	422	100%	1116	100%	933	100%	229	98%	122	99%	87	96%
Age at diagnosis Median (range)	61 (2	7-102)	E9 ('	28-94)	62 (1	29-98)	E9 E/	27-101)	50 (2	30-95)	64 (2	35-96)	74 (2	9-102)
	127	4%	50 (J	1%	38	3%	62	7%	15	6%	04 (3 5	4%	1 1	1%
40 - 49	538	18%	100	24%	176	16%	201	21%	46	20%	9	7%	6	7%
40 – 49 50 – 59	719	25%	124	24 %	261	23%	201	21%	62	20%	33	27%	13	14%
<u> </u>	660	23%	95	29%	201	25 %	185	24 %	40	17%	29	24%	21	23%
70 – 79	583	20%	74	18%	290	20 %	164	18%	35	15%	23	19%	21	23 %
80+	300	10%	25	6%	86	24 % 8%	104	11%	35	15%	23	20%	30	33%
o∪+ Margin status	300	1070	25	070	00	070	100	1170	30	1070	24	2070	30	5570
Positive	123	4%	14	3%	20	2%	39	4%	26	11%	15	12%	9	10%
Negative	2356	81%	371	88%	990	89%	795	85%	168	72%	32	26%	0	-
Close	165	6%	29	7%	56	5%	61	7%	100	7%	32	20 %	0	
Unknown	283	10%	10	2%	52	5%	43	5%	231		73	59%	82	90%
Tumour Size	203	10 /0	10	2 /0	52	570	40	570	201	0 /0	73	3370	02	30 /0
Median (range)	17(0	.1-9.9)	15(0	).1-9.9)	12(0	.1-2.0)	25(0	).1-9.9)	54(0	.1-9.9)	43(0	.4-9.9)	13(1	.1-1.5)
<1.0 cm	529	18%	122	29%	370	33%	27	3%	8	3%	2	2%	0	-
1.0 to 2.0 cm	1110	38%	139	33%	708	63%	227	24%	20	9%	14	11%	2	2%
2.1 to 5.0 cm	822	28%	90	21%	0	-	628	67%	70	30%	34	28%	0	-
>5.0 cm	204	7%	34	8%	0	_	19	2%	115	49%	36	29%	0	_
Unknown	262	9%	39	9%	40	4%	37	4%	20	9%	37	30%	89	98%
ER status		- /-												
Positive	1920	66%	30	7%	943	84%	707	75%	165	71%	63	51%	12	13%
Negative	457	16%	13	3%	139	12%	219	23%	57	25%	26	21%	3	3%
Unknown	550	19%	381	90%	36	3%	12	1%	11	5%	34	28%	76	84%
Grade														
1	784	27%	85	20%	487	44%	177	19%	22	9%	9	7%	4	4%
2	1050	36%	152	36%	413	37%	361	39%	82	35%	35	29%	7	8%
3	875	30%	147	35%	195	17%	383	41%	108	46%	38	31%	4	4%
Unknown	218	7%	40	9%	23	2%	17	2%	21	9%	41	33%	76	84%
LVI														
Positive	519	18%	0	_	77	7%	297	32%	114	49%	28	38%	3	3%
Negative	1747	60%	18	4%	994	89%	600	64%	85	36%	40	33%	10	11%
Unknown	661	23%	406	96%	47	4%	41	4%	34	15%	55	45%	78	86%

# Positive nodes														
0	1439	49%	71	17%	1005	90%	339	36%	17	7%	7	6%	0	-
1-3	497	17%	0	-	0	-	422	45%	65	28%	10	8%	0	-
4+	252	9%	0	-	0	-	124	13%	107	46%	20	16%	1	1%
Positive (# unk)	3	0%	0	-	0	-	0	-	3	1%	0	-	0	-
Nodal status unk	736	25%	353	83%	113	10%	53	6%	41	18%	86	70%	90	99%
SMPBC attender														
Yes	1574	54%	302	71%	704	63%	431	46%	81	35%	33	27%	23	25%
No	1353	46%	122	29%	414	37%	507	54%	152	65%	90	73%	68	75%
Screen detected*														
Yes	971	62%	238	79%	499	71%	189	44%	25	31%	11	33%	9	39%
No	603	38%	64	21%	205	29%	242	56%	56	69%	22	67%	14	61%

\* Screen detected defined as diagnosis of breast cancer within 1 year of an abnormal screen. For patients with synchronous bilateral disease, the first diagnosis was used to define the screen detected variable, which was then assigned to both diagnoses.

ER = Estrogen Receptor

LVI = Lymphovascular Invasion

Unk = Unknown

SMPBC = Screening Mammography Program of British Columbia

# Table 2. Comparison of stage distribution for British Columbia and SurveillanceEpidemiology and End Results breast cancer cases.

Stage	BC % of Cases	SEER % of Cases
In Situ	14%	15%
I	38%	42%
II	32%	32%
III	8%	7%
IV	4%	4%
Unknown	3%	-

### **Table 3. Treatment Characteristics**

	All Stage		Stage 0		Stage I		Stage II		Stage III		Sta	ge IV	Stage Unknown	
	n	%	n	%	n	%	n	%	n	%	n	%	n	%
	2927	100	424	14.5	1118	38.2	938	32.0	233	8.0	123	4.2	91	3.1
Initial Surgery														
None	121	4%	1	0%	7	1%	13	1%	22	9%	65	53%	13	14%
BCS	1510	52%	281	66%	726	65%	445	47%	34	15%	22	18%	2	2%
Mastectomy	1086	37%	104	25%	344	31%	458	49%	154	66%	22	18%	4	4%
Unknown	210	7%	38	9%	41	3%	22	2%	23	10%	14	11%	72	79%
RT within 1 year of diagnosis	1649	56%	159	38%	655	59%	599	64%	179	77%	57	46%	0	-
RT within 5 years of diagnosis	1719	59%	167	39%	679	61%	619	66%	184	79%	65	52%	NA	
RT within 1 year of BCS	1223	81%	155	55%	639	88%	394	88.5%	26	77%	NA		NA	
Chemotherapy (CT) within 1 year of diagnosis	928	31%	0	-	159	14%	543	58%	166	71%	53	43%	7	8%
CT within 5 years of diagnosis	1008	34%	0	-	177	16%	558	59%	168	72%	61	49%	NA	
Hormonal therapy within 1 year of diagnosis (All)	1664	57%	95	22%	709	63%	610	65%	156	67%	63	51%	31	34%
HT within 5 years of diagnosis	1777	61%	109	26%	734	66%	657	70%	169	73%	67	54%	NA	
Hormonal therapy within 1 year of diagnosis (ER+)	2283	78%	89	21%	816	73%	788	84%	200	86%	100	81%	NA	

HT = Hormone therapy

ER+ = Estrogen Receptor-Positive

		All Stages	Stage I	Stage II	Stage III	Stage IV
	BCCA	59	61	66	79	52
RT (%)	Ideal (19)	66	69	82	95	64
	Ideal (20)	83	84	84	91	47
Chemotherapy	BCCA	34	16	59	72	49
(%)	Ideal (21)	59	56	56	90	29
Hormonal	BCCA	61	66	70	73	54
therapy (%)	Ideal (22)	64	NR	NR	NR	NR

# Table 4. Comparison of 5 year BCCA and optimal use of RT and systemic therapies.

# Table 5A. Comparison of British Columbia outcomes for patients diagnosed in 2002 with data from the United States SEER database.

Stage	5 Year OS	5 Year RSR	5 Year RSR		
	BC	BC	USA		
			SEER (18)		
In Situ	96%	103%	100%		
I	90%	98%	100%		
П	84%	91%	86%		
III	60%	65%	57%		
IV	12%	13%	20%		
All Cases	83%	90%	89%		

OS = Overall Survival BC = British Columbia RSR= Relative Survival Rate SEER = Surveillance Epidemiology and End Results

Table 5B. Comparison of British Columbia outcomes for patients diagnosed in2002 with data from other cancer registries in the United States and Europe.

5 Year RSR BC	5 Year RSR USA SEER (18)	5 Year RSR ICBP Denmark (10)	5 Year RSR ICBP Norway (10)	5 Year RSR ICBP Sweden (10)	5 Year RSR ICBP UK (10)	5 Year OS EUROCARE (11)	Germany 5 Year RSR (23)
90%	89%	82%	84%	89.3%	78.8%	79%	81%

RSR= Relative Survival Rate

SEER = Surveillance Epidemiology and End Results

ICBP = International Cancer Benchmark Project

UK- United Kingdom