Sentinel Node Surgery in Breast Cancer
– Results over ten years in a university hospital

Helle Eilertsen\textsuperscript{a}, Marianne Efskind Harr\textsuperscript{b}, Rolf Kåresen\textsuperscript{c}, Daehoon Park\textsuperscript{d}, Torill Sauer\textsuperscript{e}, Ellen Schlichting\textsuperscript{f}

\textsuperscript{a} University of Oslo, Medical Faculty, Oslo, Norway
\textsuperscript{b} Department of neurosurgery, Oslo University Hospital, Norway
\textsuperscript{c} Department of Breast- and endocrine Surgery, Oslo University Hospital and University of Oslo, Medical Faculty, Oslo, Norway
\textsuperscript{d} Department of Pathology, Vestre Viken Hospital, Drammen, Norway
\textsuperscript{e} Department of Pathology, Oslo University Hospital and University of Oslo, Medical Faculty, Oslo, Norway
\textsuperscript{f} Department of Breast- and endocrine surgery, Oslo University Hospital, Oslo, Norway

Corresponding author:
Ellen Schlichting, Department of Breast- end endocrine Surgery, Oslo University Hospital, Postboks 4956 Nydalen, 0424 Oslo, Norway. elsc@uus.no

Abstract:
Background: Sentinel node biopsy (SNB) was introduced at Ullevål University Hospital in 2000. This article presents results from the first ten years using the method, focusing primarily on extra sentinel metastases in patients with positive sentinel nodes (SN) and axillary recurrences in patients with negative SNs.
Material and methods: A prospective registration of 2762 patients was made from 2000 through 2009.
Results: The median follow-up time was 51 months. The overall detection rate was 93%. 36% of the patients with positive SNs had extra sentinel metastases. These were significantly associated with a macrometastatic SN and a primary tumour > 20 mm. 18% of patients with sentinel metastasis ≤ 2 mm had extra sentinel metastases. 14 patients with negative SN (0.7%) developed axillary recurrence. 32% with a preoperative diagnosis of ductal carcinoma in situ (DCIS) were upstaged to infiltrating carcinoma on final histology. None of the patients with pure DCIS had positive SNs.
Conclusion: Few late events (0.7%) in SN negative axillas demonstrate the safety of the technique.

Key words:
Breast cancer
Ductal carcinoma in situ
Sentinel node
Extra sentinel metastasis
Axillary recurrence
Introduction

As one of the first hospitals in Norway, Ullevål University Hospital introduced sentinel node (SN) surgery as a routine method in March 2000. Between 400 and 500 patients were diagnosed with breast cancer in the hospital per year during the study period. We present the results from the first ten years using the SN technique with a particular focus on four aspects:
- Differences in detection rate on the basis of injection site
- The correlation between size of SN metastasis and the risk of metastasising to extra SN lymph nodes
- The role of SN biopsy in patients with a preoperative diagnosis of ductal carcinoma in situ (DCIS)
- Axillary recurrences in patients with negative SN in the first operation

Patients and methods

Between March 2000 and December 2009, 2762 patients (2751 women and 11 men) with breast cancer underwent SNB at Ullevaal University Hospital. 36 patients had bilateral surgery and thus 2798 SNBs were performed during the study period. The median age was 58 years (range 22-92). 85% of the patients had preoperative diagnosis of DCIS or infiltrating carcinoma by either fine needle aspiration cytology (FNAC) or core biopsy, the rest by surgical biopsy.

Registration of data:
A prospective, scheme based registration was used. The data was kept in an internal hospital database with the acceptance from the relevant authorities.

Inclusion and exclusion criteria:
For the first 5 years patients aged 75 and under with a preoperative estimated tumour size of less than 3 cm were included. The last 5 years also patients with tumours between 3 and 5 cm were included with no upper age limit. The exclusion criteria were for the entire period patients with cytologically and clinically acknowledged axillary metastases and those who had preoperative chemotherapy. During the first 5 years DCIS and multifocal disease were exclusion criteria, but for the last 5 years multifocality and DCIS grade III on cytology or histology were included.

Identification of the sentinel node:
Both blue dye and radioactivity were used to identify the SN. For the first 6 years, blue dye and radioactivity were injected peritumourally. From January 2006 the injection was made periareolarly. Radioactivity was injected the day before surgery, whereas blue dye was injected after the patients were under general anaesthesia. A lymph node was defined as a SN if the radioactivity was at least ten times the background activity, whereas the degree of colour was determined by a subjective assessment by the surgeon.

Histopathological examination of sentinel node:
All sentinel nodes were intraoperatively sent to frozen section examination. The nodes were cut in 2 or 3 and 2-3 frozen sections were cut from each cut surface. About 8-12 sections were examined per lymph node. All SNs were later fixated, embedded and stained with hematoxylin-eosin and examined as part of the final histological evaluation. In cases of doubt, immunohistochemical testing was performed.\textsuperscript{1}

**Axillary lymph node dissection:**
Axillary lymph node dissection (ALND) was performed if the SN was positive either intraoperatively or after final histological assessment. In the case of a negative SN intraoperatively, the patient was reoperated if final histology found a metastasis (intraoperatively false negative SN). Patients found to have SNs with isolated tumour cells (metastasis < 0.2 mm) did not undergo ALND, according to the guidelines of the Norwegian Breast Cancer Group.\textsuperscript{2}

**Reevaluation of positive SNs:**
Positive SNs were later re-examined and the largest diameter of metastasis remeasured by an independent pathologist (DP) blinded for other data.

**Definition of axillary recurrence:**
Axillary recurrence was defined as detection of cancer cells in the axilla more than 120 days after the date of primary axillary surgery. The reports of axillary recurrences were found in patient records in the hospital and from reports given by the Norwegian Cancer Registry.

**Statistical analysis:**
Comparison of data between groups was made using chi-square test and Fisher exact test. Multivariate analyses were made using a multiple logistic regression model. A two-tailed p-value of 0.05 or less was considered statistically significant. Analyses were performed using SPSS version 18.0 (SPSS Inc., Chicago, Illinois, USA).

**Results**
Median follow-up time was 51 months (range 0-117). Histopathological tumour characteristics are given in table 1.

**Detection of Sentinel Node**
The detection rate for the entire period was 93% (95% CI 92 - 94). Peritumoural injection of blue dye and radioactivity gave a detection rate of 90% (95% CI 88 - 91), whereas the detection rate with periareolar injection was 96% (95% CI 95 - 97). There was a significant difference in detection rate after change of injection site (p < 0.001).

A total of 30 surgeons were operators. Their detection rate did not differ significantly when grouped according to the number of procedures they performed (Table 2). On multivariate analysis, neither age, tumour grade or tumour size was found to be significant factors in the detection of SN.

A median value of 1 node (range 1-8) was retrieved during the SNB.
27% of the patients in whom SN was not found, had a metastasis to one or more axillary nodes. The median value of metastatic nodes after ALND was 3 (range 1-15).

**Positive sentinel nodes**
A positive SN was on final histopathological examination found in 620 of the 2586 patients (24%) where SN was detected. Largest diameter of SN metastases in each patient categorized in three groups is shown in table 3.
141 patients had an intraoperatively false negative SN and were reoperated with ALND. The negative predictive value of intraoperative SN assessment was 93% (95% CI 92 - 94). Of the patients with an intraoperatively false negative SN, 71% had micrometastasis ≤ 2 mm.
On multivariate analysis, tumour size > 20 mm had a significant association to positive sentinel nodes (p < 0.001), whereas tumour grade and receptor status had no significant correlation.

**Extra SN metastases in patients with positive SN**
Of the patients with positive SN, 36% had metastasized also to extra sentinel lymph nodes.
There was a significant difference in the share of extra SN metastases in correlation with size of the SN metastasis (p < 0.001). The correlation is shown in table 4.
A multivariate analysis comparing SN positive patients with and without extra sentinel metastasis, found that the only other characteristic, apart from size of SN metastasis, was size of primary tumour. A significant difference was seen between patients with tumours smaller than and larger than 20 mm (p = 0.013).

**Positive sentinel node in ductal carcinoma in situ grade III**
There were 225 patients with DCIS grade III on either FNAC or core biopsy. Of these, 72 (32%) had infiltrating carcinoma on final histology. Of the 211 patients with a preoperative DCIS grade III diagnosis that we have data on, 23 had a positive SN (11%), 11 of which had micrometastasis ≤ 2mm. Five of these 23 patients had extra sentinel metastases, but none of these had micrometastasis. None of the patients with DCIS on final histology had a positive SN.

**Axillary recurrences in SN negative patients**
Of those with negative SN, axillary recurrence was seen in 14 patients (0.7%). The median interval of months between time of surgery and registered date of axillary recurrence was 24 (range 4 - 34).
A comparison of tumour characteristics of patients with and without axillary recurrences, showed significant differences for share of grade 3 tumours (p = 0.012), but not for age, size or receptor status (Table 5).

**Discussion**
Our rate of axillary recurrences supports the conclusion from other studies: A negative SN is a reliable indicator of axillary status and omitting axillary dissection after negative sentinel node biopsy is safe.12,15,16,32-34

**Detection rate**
Our overall detection rate is slightly lower than what is reported in other studies3-6, but meets the requirements set by the Norwegian Breast Cancer Group2. The
A significant difference in detection rate after change of injection site validates the change in guidelines made by the Norwegian Breast Cancer Group. A hypothesis that the detection rate was low because of a large number of inexperienced surgeons at our teaching hospital failed, when we found no significant differences in identification rate between the experienced and inexperienced surgeons. Further, no clinical or tumour characteristics were found to be significant in detecting SN. This suggests that the method is less dependent of the surgeon and the patient population and more dependent on qualities within the method itself. Thus an explanation for the relative low detection rate might be our stringent definition for a SN to have at least ten times the background radioactivity. Most publications on the method does not define this level and might thus have included nodes we excluded due to too low levels of activity.

Extra sentinel node metastases
Macrometastasis to SN and size of primary tumour are significant predictors for the involvement of extra SN metastasis. Similar findings have been reported in other studies. 18% of our patients with a SN metastasis ≤ 2 mm had further nodal involvement. A metaanalysis of 25 studies found an incidence of further nodal involvement in 20% of patients with micrometastasis or isolated tumour cells to SN. The literature has for years been contradictory about micrometastases` influence on recurrence and survival. Guiliano et al concluded in their randomized controlled trial (RCT) that ALND might no longer be justified for women with small breast carcinomas and a positive SN as long as a set of specified adjuvant therapies are given. The Norwegian Breast Cancer Group recently decided that patients with SN metastasis ≤ 2 mm should avoid having ALND if breast preserving therapy including radiotherapy to the breast and lower axilla is carried out.

Sentinel node metastasis in ductal carcinoma in situ
The value of doing SNB in DCIS has been disputed; some authors are in favour, others against. Most, however, agree that it should be done in patients who undergo mastectomy, because this eradicates the possibility to later perform SNB if preoperative diagnosis of DCIS is changed to infiltrating carcinoma. None of the preoperative DCIS patients in our material who on final histology was found to have true DCIS had positive SNs, whereas 11% of the patients that were upstaged to infiltrating carcinoma did. Based on these results we find that omitting SNB in patients with pure DCIS on final histology could be justified, whereas patients that on final histology have an infiltrating component should undergo SNB. The large share of patients in our material (32%) who were upstaged from a preoperative diagnosis of DCIS grade III by FNAC or core biopsy to infiltrating carcinoma on final histology, shows that invasive foci are often overseen in the preoperative evaluation, and that many patients will have to return for a second operation if SN is not done in the primary operation. Some studies have looked at risk factors among preoperative DCIS patients that can predict presence of an invasive component and thus who could benefit from having SNB performed as part of the initial operation. The studies have contradicting results and a firm conclusion is at the present time not possible to make.

Axillary recurrences
An axillary recurrence rate of 0.7% with a median follow up time of 51 months is acceptable and in accordance with results from other studies. A recent study with a median follow-up time the same as ours, found a recurrence rate of 0.7%\textsuperscript{27}, another reports an axillary recurrence rate of 0.6% after a median follow up time of 37 months.\textsuperscript{28} A systematic review and meta-analysis of 48 studies found a recurrence rate for axillary metastasis in clinically node negative women with a primary negative SN of 0.3% after a median follow up time of 34 months.\textsuperscript{29}

Our results indicate that longer follow-up time does not increase the rate of recurrence much, in accordance with other studies which observe that the majority of recurrences happens during the first years after diagnosis.\textsuperscript{5,14,30}

Apart from a significant higher share of grade 3 tumours, we found no other significant characteristics in the axillary recurrence population. Similar results were reported by Kiluk et al\textsuperscript{31} and Bergquist et al\textsuperscript{28}. However, we had very few cases of recurrences in each risk category, and the lack of significant relation of recurrence to the other risk factors may be due to the low number ($n = 14$) and thus lack of statistical power.

One explanation of the low axillary recurrence rate might be that we have a low-risk population with a high percentage of good prognostic factors such as small tumour size (median 15 mm), oestrogen (84%) and progesterone (68%) positivity and a low frequency of poor prognostic factors, such as lymph node involvement (24%) and Her2 positivity (9%) (Table 1). This selection is probably explained by the introduction of mammography screening for women aged 50-69 in the hospital’s referral area in 1996, four years prior to the introduction of the SN procedure.

### Conclusion

Few recurrences (0.7%) in SN negative axillas demonstrate the safety of the technique. Apart from a significant higher share of grade 3 tumours, we found no other significant characteristics in the axillary recurrence population. The median time to recurrence was 24 months.

The SN procedure seems unnecessary in patients with histology proven DCIS. The rather large share of patients with extra SN-metastases, also in those with micrometastasis (18%), shows that removal of SN alone might not be sufficient for local control of the axilla.

The detection rate increased after change of injection site from peritumoral to periareolar.

### Acknowledgements:
Siri Larønningen, Cancer Registry, Norway
References:


Table 1. Tumour characteristics of 2798 breast cancer patients. Missing data given in numbers only. The percentages thus show the distribution of the parameters in known cases. *The registration of Her2 started in 2004.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Number</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Median invasive tumour size (n = 2476, missing 41)</td>
<td>15 mm (range 0.5, 86)</td>
<td></td>
</tr>
<tr>
<td>Histological type (n = 2776)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ductal</td>
<td>1507</td>
<td>54%</td>
</tr>
<tr>
<td>Lobular</td>
<td>220</td>
<td>8%</td>
</tr>
<tr>
<td>DCIS</td>
<td>246</td>
<td>9%</td>
</tr>
<tr>
<td>Ductal + DCIS</td>
<td>623</td>
<td>22%</td>
</tr>
<tr>
<td>Other</td>
<td>180</td>
<td>7%</td>
</tr>
<tr>
<td>Missing</td>
<td>22</td>
<td></td>
</tr>
<tr>
<td>Histological grading of invasive ductal carcinomas (n = 1499)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>469</td>
<td>31%</td>
</tr>
<tr>
<td>2</td>
<td>695</td>
<td>47%</td>
</tr>
<tr>
<td>3</td>
<td>335</td>
<td>22%</td>
</tr>
<tr>
<td>Missing</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Oestrogen receptor status: (n = 2456)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive</td>
<td>2059</td>
<td>84%</td>
</tr>
<tr>
<td>Negative</td>
<td>397</td>
<td>16%</td>
</tr>
<tr>
<td>Missing</td>
<td>61</td>
<td></td>
</tr>
<tr>
<td>Progesterone receptor status: (n = 2444)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive</td>
<td>1662</td>
<td>68%</td>
</tr>
<tr>
<td>Negative</td>
<td>782</td>
<td>32%</td>
</tr>
<tr>
<td>Missing</td>
<td>73</td>
<td></td>
</tr>
<tr>
<td>Her2 status (n = 1459*)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive</td>
<td>138</td>
<td>9%</td>
</tr>
<tr>
<td>Negative</td>
<td>1321</td>
<td>91%</td>
</tr>
<tr>
<td>Missing</td>
<td>39</td>
<td></td>
</tr>
<tr>
<td>Total SN procedures per surgeon</td>
<td>≤ 30</td>
<td>&gt;30 ≤100</td>
</tr>
<tr>
<td>--------------------------------</td>
<td>------</td>
<td>----------</td>
</tr>
<tr>
<td>Det. Rate [95% CI]</td>
<td>91% [86 - 95]</td>
<td>94% [91 - 97]</td>
</tr>
</tbody>
</table>

Table 2. Detection rate related to total number of sentinel node (SN) procedures done by each surgeon.

<table>
<thead>
<tr>
<th>Size</th>
<th>≤ 2 mm</th>
<th>&gt;2 ≤ 5 mm</th>
<th>&gt; 5 mm</th>
</tr>
</thead>
<tbody>
<tr>
<td>Share of SN metastases (n = 620, 16 missing )</td>
<td>170 (28%)</td>
<td>158 (26%)</td>
<td>276 (46%)</td>
</tr>
</tbody>
</table>

Table 3. Largest diameter of sentinel node (SN) metastases.

<table>
<thead>
<tr>
<th>Size (mm)</th>
<th>≤ 2</th>
<th>&gt;2 ≤5</th>
<th>&gt; 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Extra SN metastasis [95% CI]</td>
<td>17% [12 - 23]</td>
<td>33% [26 - 40]</td>
<td>49% [43 - 55]</td>
</tr>
</tbody>
</table>

Table 4. Share of extra sentinel node (SN) metastasis related to size of sentinel node metastasis.

<table>
<thead>
<tr>
<th>Patients with negative SN, n = 1966</th>
<th>Patients with axillary recurrence, n = 14</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Median age (years)</td>
<td>59</td>
<td>58</td>
</tr>
<tr>
<td>Grade 3</td>
<td>217/995 (22%)</td>
<td>7/14 (50%)</td>
</tr>
<tr>
<td>Oestrogen receptor negative</td>
<td>281/1667 (17%)</td>
<td>4/12 (33%)</td>
</tr>
<tr>
<td>Progesterone receptor negative</td>
<td>544/1661 (33%)</td>
<td>6/12 (50%)</td>
</tr>
<tr>
<td>Median invasive tumour size (mm)</td>
<td>14</td>
<td>13</td>
</tr>
</tbody>
</table>

Table 5. Tumour characteristics in patients with and without axillary recurrence.