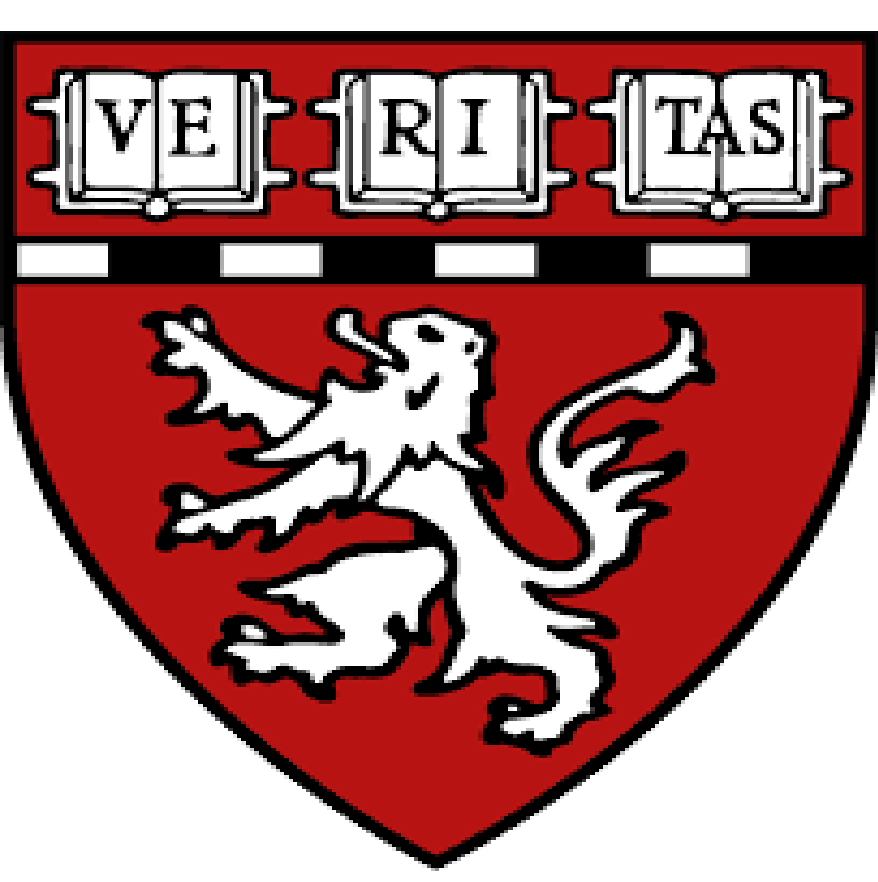




# Implementation of Sentinel Lymph Node Biopsy Following Neo Adjuvant Chemotherapy in Clinically Node Positive Breast Cancer



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## BACKGROUND

- Multiple prospective clinical trials have recently evaluated the feasibility of sentinel lymph node biopsy (SLNB) in patients with clinically positive nodes undergoing neo adjuvant chemotherapy (NCT).
- The data have shown that a low false negative rate can be attained in patients who become clinically node negative after NCT provided that adequate nodal sampling is performed.
- The adoption of this approach into clinical practice may be limited, leaving patients unnecessarily exposed to the morbidity of an axillary node dissection.
- The aim of the present study was to assess the implementation of recent clinical trial data regarding the feasibility of SLNB following NCT for clinically node positive patients, and determine factors associated with variation in clinical practice.

## METHODS

- A retrospective analysis of the National Cancer Database was performed, consisting of women with stage 1-3 invasive breast cancer from 2012 to 2015 with clinically positive nodal disease and who underwent NCT.
- Pathological complete response (pCR) in the nodes was defined as absence of regional lymph node metastasis on final pathology.
- Chi square analyses were performed to compare rates of pCR in patients with axillary lymph node dissection (ALND).

STUDY POPULATION CHARACTERISTICS FOR PATIENTS UNDERGOING NEOADJUVANT CHEMOTHERAPY WITH CLINICALLY NODE POSITIVE DISEASE

| Characteristic                            | No SLNB<br>n=5574 |       | SLNB alone<br>n=1405 |       | SLNB + ALND<br>n=1859 |       |          |
|---|-------------------|-------|----------------------|-------|-----------------------|-------|----------|
|   | n                 | %     | n                    | %     | n                     | %     |          |
| <b>Age group (years)</b>                  |                   |       |                      |       |                       |       |          |
| <40                                       | 808               | 60.3  | 249                  | 18.58 | 283                   | 21.12 | p=0.0002 |
| 40-54                                     | 2,322             | 61.58 | 638                  | 16.92 | 811                   | 21.51 |          |
| 55-69                                     | 2,039             | 65.37 | 438                  | 14.04 | 642                   | 20.58 |          |
| >=70                                      | 405               | 66.61 | 80                   | 13.16 | 123                   | 20.23 |          |
| <b>Income</b>                             |                   |       |                      |       |                       |       |          |
| High SES                                  | 1,853             | 59.07 | 550                  | 17.53 | 734                   | 23.4  | p<0.001  |
| Low SES                                   | 3,721             | 65.27 | 855                  | 15    | 1125                  | 19.73 |          |
| <b>Insurance</b>                          |                   |       |                      |       |                       |       |          |
| Not insured                               | 310               | 74.7  | 44                   | 10.6  | 61                    | 14.7  | p<0.001  |
| Private insurance                         | 3,392             | 60.11 | 987                  | 17.49 | 1264                  | 22.4  |          |
| Public                                    | 1,872             | 67.34 | 374                  | 13.45 | 534                   | 19.21 |          |
| <b>Facility type</b>                      |                   |       |                      |       |                       |       |          |
| Academic/Research                         | 1,695             | 65.88 | 351                  | 13.64 | 527                   | 20.48 | p=0.0002 |
| Other                                     | 3,879             | 61.92 | 1054                 | 16.82 | 1332                  | 21.26 |          |
| <b>Volume</b>                             |                   |       |                      |       |                       |       |          |
| Low volume                                | 364               | 66.3  | 84                   | 15.3  | 101                   | 18.4  | p=0.074  |
| Medium volume                             | 1,741             | 64.6  | 403                  | 14.95 | 551                   | 20.45 |          |
| High volume                               | 3,469             | 62.01 | 918                  | 16.41 | 1207                  | 21.58 |          |
| <b>Grade</b>                              |                   |       |                      |       |                       |       |          |
| Low/intermediate                          | 2,849             | 62.19 | 715                  | 15.61 | 1017                  | 22.2  | p=0.0201 |
| High                                      | 2,725             | 64.01 | 690                  | 16.21 | 842                   | 19.78 |          |
| <b>CDCC</b>                               |                   |       |                      |       |                       |       |          |
| CDCC 0                                    | 4,818             | 62.45 | 1,279                | 16.58 | 1618                  | 20.97 | p<0.001  |
| CDCC >=1                                  | 756               | 13.56 | 126                  | 8.97  | 241                   | 12.96 |          |
| <b>Markers</b>                            |                   |       |                      |       |                       |       |          |
| HR <sup>C</sup> -positive / HER2-negative | 3,829             | 64.24 | 842                  | 14.13 | 1289                  | 21.63 | p<0.001  |
| HR-negative / HER2-positive               | 26                | 60.47 | 5                    | 11.63 | 12                    | 27.91 |          |
| HR-positive / HER2-positive               | 1,596             | 60.14 | 523                  | 19.71 | 535                   | 20.16 |          |
| Triple negative                           | 87                | 67.97 | 27                   | 21.09 | 14                    | 10.94 |          |
| unknown                                   | 36                | 67.92 | 8                    | 15.09 | 9                     | 16.98 |          |

PATHOLOGICAL COMPLETE RESPONSE (pCR) IN PATIENTS UNDERGOING NEOADJUVANT CHEMOTHERAPY IN CLINICALLY NODE POSITIVE DISEASE

| Characteristic                            | pCR |       | No pCR |       |          |
|---|-----|-------|--------|-------|----------|
|   | n   | %     | n      | %     |          |
| <b>Age group (years)</b>                  |     |       |        |       |          |
| <40                                       | 84  | 29.68 | 199    | 70.32 | p=0.0011 |
| 40-54                                     | 182 | 22.44 | 629    | 77.56 |          |
| 55-69                                     | 115 | 17.91 | 527    | 82.09 |          |
| >=70                                      | 28  | 22.76 | 95     | 77.24 |          |
| <b>Clinical N</b>                         |     |       |        |       |          |
| C1  | 346 | 22.05 | 1223   | 77.95 | P=0.0171 |
| C2  | 31  | 16.58 | 156    | 83.42 |          |
| C3  | 32  | 31.07 | 71     | 68.93 |          |
| <b>Pathological T</b>                     |     |       |        |       |          |
| T0  | 155 | 72.77 | 58     | 27.23 | p<0.001  |
| T1  | 188 | 22.41 | 651    | 77.59 |          |
| T2  | 56  | 9.62  | 526    | 90.38 |          |
| T3/T4                                     | 10  | 4.44  | 215    | 95.56 |          |
| <b>Grade</b>                              |     |       |        |       |          |
| Low/intermediate                          | 154 | 15.14 | 863    | 84.86 | p<0.001  |
| High                                      | 255 | 30.29 | 587    | 69.71 |          |
| <b>CDCC</b>                               |     |       |        |       |          |
| CDCC 0                                    | 372 | 22.99 | 1246   | 77.01 | p=0.0076 |
| CDCC >=1                                  | 37  | 15.35 | 204    | 84.65 |          |
| <b>Markers</b>                            |     |       |        |       |          |
| HR <sup>C</sup> -positive / HER2-negative | 194 | 15.05 | 1095   | 84.95 | p<0.001  |
| Others                                    | 215 | 37.72 | 355    | 62.28 |          |

## RESULTS

- A greater proportion of ALNDs were performed in non-academic institutions (1,332, 71.7%).
- Age 40-54 years (811, 44%), patients low comorbidity scores (1,618, 87%), and lower socioeconomic levels (1,125, 60%) were all also associated with higher rates of ALND.
- Patients with hormone receptor positive/Her 2 negative tumors (1,289, 69%), and tumors of low/intermediate grade (1,017, 55%), were more likely to undergo ALND.
- Of 1,857 women, 22% had a pCR.

## CONCLUSIONS

- Despite the promising results shown in prospective, randomized clinical trials, a large proportion of patients with complete nodal response to NCT still undergo ALND.
- These patients are susceptible to the morbidity associated with this procedure.
- Over 20% of patients with clinically positive nodes treated with NCT were found to be pN0 at the time of ALND.
- Specific facility, patient, and tumor characteristics were associated with clinical practice variation in the use of SLNB following NCT.
- Although a portion of patients may have had clinical indications for ALND, our findings suggest that barriers may exist in the translation of clinical trial findings to routine practice.
- Efforts to address these potential barriers may result in better outcomes for patients treated for breast cancer.