Association of Histopathological Parameters and Axillary Lymphnode Metastasis in Primary Breast Carcinoma

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Abstract

Introduction: The most common malignancy worldwide among females is breast carcinoma and the second most common malignancy in India, next to cervical cancer. A wide range of potential prognostic features have been studied in breast cancer and are mainly divided into two groups i.e. Histopathological and Molecular. The histological features are cost-effective and provide reliable diagnostic and prognostic information in these tumors. Axillary Lymph node status is one of the most important prognostic factors and greatly affects the morbidity and mortality of the patient. Materials and Methods: All breast cancer specimens received in the Department of Pathology over a period of five years. The following histopathological parameters were carefully studied like Tumor size, Histological type, Grade, Presence of necrosis, Inflammatory cell infiltrate, Lymphatic invasion, Blood vessel invasion, Perineural invasion, and other Stromal changes were studied in detail, and association of these histopathological parameters with axillary lymph node metastasis were analyzed. Results: A total of 100 cases were studied, and most of the patients were over the age of 50. The maximum number of cases was in the T2 stage (55%). Infiltrating ductal carcinoma (88%) was the most common type of tumor encountered in the study. The majority of the cases were Grade I tumors. Skin Invasion was seen in 14% and Lymphovascular Invasion was seen in 17% of cases respectively. There was a statistically significant association between the size of the tumor, T stage, Grade of the tumor, necrosis, and inflammatory infiltrate on further analysis. Conclusion: There was a statistically significant correlation between Tumor size, pathological T stage, Grade of the tumor, Necrosis and inflammatory infiltrate with axillary Lymph node metastasis in the present study. Increased tumor size, T stage, higher grade, presence of necrosis and low inflammatory infiltrate are associated with increased axillary Lymph node metastasis.

Keywords: Breast carcinoma- Histopathology- prognostic factors- axillary lymph nodes

Introduction

The most common malignancy worldwide among females is breast carcinoma and in India, the second most common malignancy next to cervical cancer. In India the disease specific mortality of breast cancer is roughly 50% [1]. The incidence of breast cancer in Kolar region is around 6.41% of all malignancies [2]. The worldwide incidence of breast cancer comprises of 10.4% of all malignancies in female population. Breast cancer, a heterogenous disease has varied morphological appearance, molecular features and behavior of response to therapy. It is becoming increasingly important to assess the prognosis of breast cancer in each patient before treatment [1,2].

A wide range of potential prognostic features have been studied. They are mainly divided into two groups i.e. Histopathological and Molecular. Histopathological prognostic features are relatively simple to assess and provide clinically useful prognostic information. Tumor size, grade, histopathological type and axillary lymph node status are most important histopathological features [3-5].

Lymph node staging is an important diagnostic factor and provides prognostic information and showed to be based on histological examination than clinical and radiological examination [4]. Patients with positive...
axillary lymph node shows higher mortality (four to eight
times) than node negative cases. Not only the disease
specific mortality increases in node positive patients but
also the risk of distant recurrence [6].

This study has been taken up to study association of
different histological parameters with axillary Lymph
nodes metastasis, which may help in predicting the
prognosis and guide the treatment of breast carcinoma.

Materials and Methods

All breast cancer specimens received in the
Department of Pathology from R.L.Jalappa Hospital and
Research Center attached to Sri Devaraj Urs Medical
College, Tamaka, and Kolar over a period of five years.
Specimens were cut at 1cm interval and kept in 10%
formalin for overnight fixation, and gross examination
was done according to standard protocol.Sufficient
number of blocks were taken, ensure adequate sampling
and routine paraffin embedding was carried out. Standard
thin sections (4-6 microns) were taken and stained with
routine Haematoxyline & Eosin stain.

Various histopathological parameters were carefully
studied in detail. Assessment of Tumor size was
done According to American joint committee on
cancer; Histological typing was done using WHO.
Grading of breast carcinoma was done using modified
Bloom Richardson scoring system. Semi quantitative
assessment of grading of necrosis was done according to
the study done by Richards CH et al [7]. Inflammatory
cell infiltrate was studied using a study done by Klintrup
et al [8] we also studied lymphatic invasion, Blood vessel
invasion, Perineural invasion, Stromal characteristics.

Inclusion Criteria

All operated breast carcinoma specimens with axillary
lymph node clearance.

Exclusion Criteria

Carcinoma of breast in male patients and sarcomas
of breast. Patients on or receiving radiotherapy or
chemotherapy and recurrent tumors were excluded from
the study

Results

100 breast carcinoma cases were studies in the present
study, out of which 53% were above 50 years. 90% of
cases presented with breast Lump, 4% presented with
ulceration and nipple discharge each and 2% presented
with Pagets diasease of breast. 14% of patients had
Tumor size <2 cms, 55% were between 2 to 5cms and
31% were >5 cms. 11% were in T1 stage, 48% were
in T2 stage, 22% were in T3 Stage and19% were in T4
Stage. Histopathological typing was done, 88% of patients
had Infiltrating Ductal Carcinoma, 2% had Infiltrating
Ductal Carcinoma with Lobular Carcinoma component,
2% had Papillary Carcinoma, 1% had Infiltrating Ductal
Carcinoma with Medullary Carcinoma component, 3%
had Medullary Carcinoma, 1% had Adenoid Cystic

Carcinoma, 2% had Infiltrating Ductal Carcinoma with
Mucinous Carcinoma and 1% had Lobular Carcinoma. 46%
of breast carcinoma cases were Grade 1, 39% were in
Grade 2 and 15% were in Grade 3. 28% of cases had Focal,
17% had moderate and 38% had extensive necrosis. In this
study, 6% had No Inflammatory Cell Infiltrate, 37% had
Mild/ Patchy Increase, 28% had Prominent Inflammatory
Reaction and 29% had Florid “Cup Like” Inflammatory
Infiltrate. Lymph node stage distribution, 47% of patients
were in N0 stage, 30% were in N1 stage, 14% were in N2
stage and 9% were in N3 stage.

Association of pathological parameters with axillary
lymph node metastasis:

1. Association between Tumor Size and axillary lymph
node metastasis: there was significant association between
Tumor size and Lymph node stage ($\chi^2 =25.24, df =9,$
p = 0.003) (Figure 1).

2. Association between Pathological T Stage and
axillary Lymph node metastasis: there was significant
association between T stage and Lymph node metastasis.
($\chi^2 =25.24, df =9,$ p = 0.003*) (Figure 2).

3. Association between Grade and axillary Lymph node
is almost similar to the studies done by EL Saghir et. al. [9] with 49.1% cases less than 50 years and mean age of 49.8 years. Similar observation was made by Najjar et. al. [10] with mean age of 49.4 years which is comparable to the present study. In the present study, tumor size, in majority of the cases were between 2 to 5 cms followed by tumor size of more than 5 cms. Similar observations were made by Ogawa Y et. al. [11] and Kwon GY et. al. [12] with highest number of cases between tumor size of 2 to 5 cms.

In our study there was significant association between Tumor size and Lymph node stage as shown in the results. As the tumor size increased, positivity of axillary lymph nodes also increased. These findings are similar to study done by Marwah, et al. [13] Ahmad, et al. [14] Postaci et al [15] and Sukla et al [16]. The risk of axillary lymph node metastasis increases as tumor size increases which suggests that nodal metastasis is indicative of tumour chronology [17].

In our study, as shown in the results, the maximum numbers of cases were in T2 stage (48%) which was similar to the study done by Wang M et al 70. T2 was followed by T3 stage (22%), T4 stage (19%) and T1 stage (11%) respectively. Pistelli M et al [18] in his study on early stage breast cancers observed that maximum number of cases were in T1 stage (53.6%) followed by T2 stage (30.8%) which was contrary to the observations made in the present study. The reason could be that the stage IV tumors were excluded in the study by Pistelli M et al 19 and the study was done only on early breast cancer patients.

The present study showed that there was strong association between histological grades of tumor with axillary metastasis. Grade II and III tumors had more positive axillary lymph nodes and grade 1 tumors showed low rate of axillary metastasis. Various studies have analyzed the importance of histological grade as a prognostic factor in carcinoma of the breast [13,14, 16,17].

Histopathologically-identified tumor necrosis has been recognized as a potential prognostic marker for a variety of tumors. In the present study, necrosis was seen in 83% of cases in comparable to the study done by Krishnamurthy et. al. [19] whereas Carlomango et. al. [20] observed necrosis in only 20.3% of cases. The observations made in the present study were similar to the study done by Matkowski R et. al. [21]. In our study there was a strong association with tumor necrosis and axillary lymph node metastasis. Other Studies have shown that necrosis in the tumor was associated with higher mortality rate, a higher incidence of axillary node metastasis, and a higher mortality rate in patients with axillary node metastasis than were primary malignancy without necrosis.

Several studies outlined the use of the inflammatory infiltrate in breast cancer as a prognostic marker [22, 23]. There was statistically significant correlation between the inflammatory infiltrate and axillary lymph node metastasis i.e. as the inflammatory infiltrate increases, the chance of axillary lymph node metastasis decreases. Aaltomaa et al [22] examined 489 breast cancer patients
with up to 10 years follow-up. They found lymphocyte infiltrate (LI) positively correlated to axillary lymph node status, tumour diameter and histomorphological variables.

Multivariate analysis showed that LI was independently related to axillary lymph node status and was able to predict recurrence free survival as well as breast cancer related survival.

In conclusion, in the present study, there was a statistically significant correlation between Tumor size, pathological T stage, Grade of the tumor, Necrosis and inflammatory infiltrate with axillary Lymph node metastasis. increased tumor size, T stage, higher grade, presence of necrosis and low inflammatory infiltrate are associated with increased axillary Lymph node metastasis. Hence, These histopathological factors can be used as prognostic markers in patients with breast cancer in a resource limited setting.

References

1. Carter D, Schnitt S, Millis R. The Breast. In: Mills SE, Greenmon J, Hornick JL, Longacre TA, Reuter VE editors. Sternberg’s Diagnostic Surgery Pathology. 6th ed. Philadelphia: Wolters Kluwer. 2015;:317-84.

2. Kalyani R, Das S, Singh B, Kumar H. Cancer profile in the department of pathology of sri devaraj urs medical college, Kolar: A ten years study. Indian Journal of Cancer. 2010;47(2):160. https://doi.org/10.4103/0019-509x.63011

3. Ellis I, Lee A, Pinder S, Rakha A. Tumors of the breast. In: World Health Organization. World Health Organization Classification of Tumours.Pathology and Genetics: Tumours of the Breast and Female Genital Organs. 2003:38-39.

4. Sinha S, Nath J, Mukherjee A, Chatterjee T. Predictive and Prognostic Factors in Breast Cancer and their Association with ERPR HER2/neu expression. I. Carcinog Mutagen. 2016;7:1-4.

5. Hoda SA, Brogi E, Koerner FC, Rosen PP. Rosen’s Breast Pathology. 3rd ed. Philadelphia: Wolters Kluwer; 2013.p.235-513.

6. Fisher B, Bauer M, Wickerham DL, Redmond CK, Fisher ER, Cruz AB, Foster R, Gardner B, Lerner H, Margozele R, Poisson R, Shibata H, Volk H. Relation of number of positive axillary nodes to the prognosis of patients with primary breast cancer. An NSABP update. Cancer. 1983 Nov 01;52(9):1551-1557. https://doi.org/10.1002/1097-0142(19831101)52:9<1551::aid-cncr2820520902>3.0.co;2-3

7. Richards CH, Flegg KM, SD Roxburgh C, Going JJ, Mohammed Z, Horgan PG, McMillan DC. The relationships between cellular components of the peritumoural inflammatory response, clinicopathological characteristics and survival in patients with primary operable colorectal cancer. British Journal of Cancer. 2012 05 17;106(12):2010-2015. https://doi.org/10.1038/bjc.2012.211

8. Klintrup K, Mäkinen JM, Kauppila S, Väre PO, Mellko J, Tuominen H, Tuppurainen K, Mäkelä J, Karttunen TJ, Mäkinen MJ. Inflammation and prognosis in colorectal cancer. European Journal of Cancer. 2005 Nov;41(17):2645-2654. https://doi.org/10.1016/j.ejca.2005.07.017

9. El Saghir NS, Shamseddine AL, Gaera F, Bikhazi K, Rahal B, Salem ZM et.al. Age distribution of breast cancer in Lebanon: increased percentages and age adjusted incidence rates of younger-aged groups at presentation. J Med Liban 2002; 50: 3-9.

10. Najjar H, Easson A. Age at diagnosis of breast cancer in Arab nations. International Journal of Surgery. 2010;8(6):448-452. https://doi.org/10.1016/j.ijjsu.2010.05.012

11. Erdogan N, Dengizmen A, Akyildizgedem A, Sahin E, Tetikku RUS. Angiogenesis in breast cancers without lymph node metastasis. Turk J Pathol 2010; 26: 136-9.

12. Kwon GY, Lee SD, Park ES. Mast Cell and Macrophage Counts and Microvessel Density in Invasive Breast Carcinoma-Comparison Analysis with Clinicopathological Parameters. Cancer Research and Treatment. 2005;37(2):103. https://doi.org/10.4143/ctr.2005.37.2.103

13. Batra A, Marwah N, Marwah S, Gupta V, Shyak S, Sen R. Correlation of proliferative index with various clinicopathologic prognostic parameters in primary breast carcinoma: A study from North India. Journal of Cancer Research and Therapeutics. 2018;14(3):537. https://doi.org/10.4103/0973-1482.167614

14. Ahmad Z, Khurshid A, Qureshi A, Idriss R, Asghar N, Kayani N. Breast carcinoma grading, estimation of tumor size, axillary lymph node status, staging, and Nottingham prognostic index scoring on mastectomy specimens. Indian Journal of Pathology and Microbiology. 2009;52(4):477. https://doi.org/10.4103/0377-4929.56123

15. Postaci H, Zengel B, Yararbas U, Usla A, Eliyatkin N, Akpinar G, Cengiz F, Durusoy R. Sentinel Lymph Node Biopsy in Breast Cancer: Predictors of Axillary and Non-Sentinel Lymph Node Involvement. Balkan Medical Journal. 2013 Dec 30;30(4):415-421. https://doi.org/10.5152/bmj.2013.9591

16. Shukla A, Jain S, Swankar M. Correlation of axillary lymph nodes involvement and Nottingham prognostic index with various histopathologic prognostic factors in invasive breast carcinoma. Int Surg J. 2019;6:1187-93.

17. Hartveit F. Axillary metastasis in breast cancer: When, how, and why?., Seminars in Surgical Oncology. 1989;5(2):126-136. https://doi.org/10.1002/sso.2980050209

18. Pistelli M, Pagliacci A, Battelli N. Prognostic factors in early stage triple negative breast cancer: Lessons and limits from clinical practice. Anticancer Res. 2013;33:2737-42.

19. Krishnamurthy J, Kumar P. Significance of prognostic indicators in infiltrating duct carcinoma breast: Scenario in developing country. Indian Journal of Cancer. 2016;53(1):34. https://doi.org/10.4103/0019-509x.180834

20. Carrolmagno C, Perrone F, Lauria R, De Laurentiis M, Gallo C, Morabito A, Pettinato G, Panico L, Belleri T, Apicella A, Petrella G, Bianco R, De Placido S. Prognostic Significance of Necrosis, Elastosis, Fibrosis and Inflammatory Cell Reaction in Operable Breast Cancer. Oncology. 1995;52(4):272-277. https://doi.org/10.1159/000222472

21. Matkowski R, Gisterek I, Halon A, Lacko A, Szewczyk K, Staszek U, et al. The prognostic role of tumor-infiltrating CD4 and CD8 T lymphocytes in breast cancer. Anticancer Res. 2009;29:2445-52.

22. Allen MD, Jones LJ. The role of inflammation in progression of breast cancer: Friend or foe? (Review). International Journal of Oncology. 2015 07 08;47(3):797-805. https://doi.org/10.3892/ijo.2015.3075

23. Aaltomaa S, Lipponen P, Eskelinen M, Kosma V, Marin S, editors. Sternberg’s Diagnostic Surgery Pathology. 6th ed. Philadelphia: Elsevier. 2013:;57-145.

24. Fletcher CDM. Diagnostic Histopathology of Tumors. 4th ed. Philadelphia: Elsevier. 2013:;57-145.

25. Alhava E, Syrjänen K. Lymphocyte infiltrates as a prognostic variable in female breast cancer. European Journal of Cancer. 1992 04;28(4-5):859-864. https://doi.org/10.1016/0959-8049(92)90134-n

26. Hartveit F. Axillary metastasis in breast cancer: When, how, and why?. Seminars in Surgical Oncology. 1989;5(2):126-136. https://doi.org/10.1002/sso.2980050209

27. Petrellia G, Bianco R, De Placido S. Prognostic Significance of Necrosis, Elastosis, Fibrosis and Inflammatory Cell Reaction in Operable Breast Cancer. Oncology. 1995;52(4):272-277. https://doi.org/10.1159/000222472

28. Matkowski R, Gisterek I, Halon A, Lacko A, Szewczyk K, Staszek U, et al. The prognostic role of tumor-infiltrating CD4 and CD8 T lymphocytes in breast cancer. Anticancer Res. 2009;29:2445-52.

29. Allen MD, Jones LJ. The role of inflammation in progression of breast cancer: Friend or foe? (Review). International Journal of Oncology. 2015 07 08;47(3):797-805. https://doi.org/10.3892/ijo.2015.3075

30. Aaltomaa S, Lipponen P, Eskelinen M, Kosma V, Marin S, editors. Sternberg’s Diagnostic Surgery Pathology. 6th ed. Philadelphia: Elsevier. 2013:;57-145.

31. Alhava E, Syrjänen K. Lymphocyte infiltrates as a prognostic variable in female breast cancer. European Journal of Cancer. 1992 04;28(4-5):859-864. https://doi.org/10.1016/0959-8049(92)90134-n

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