

CD4 and CD8 Count in Breast Cancer Patient Undergoing Post-Surgery FAC Protocol

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Abstract

Original Research Article

Objective: In this study our main goal is to evaluate the CD4 and CD8 count in breast cancer patient undergoing post-Surgery FAC protocol. **Method:** This quasi-experimental study was carried out at Dept. of Microbiology, Department of radiotherapy & Department of Surgery, Sylhet MAG Osmani Medical College Hospital from January, 01, 2011 to December, 31, 2011. Samples from the 30 cases were collected, first from Department of surgery (before operative procedure) and then, from Department of radiotherapy (after surgery and 3 cycles of adjuvant chemotherapy) from the same patient. **Results:** During the study, left breast was involved in 17 (53.3%) patients and right breast was involved in 13 (43.3%) patients. Regarding histological grading was grade-II in 13 (43.3%) cases, grade-I in 11 (37.7%) and grade-III in 6 (20.0%) cases. CD4+T cell count before chemotherapy was 1023.4 /uL (SD ± 320.7) and CD4+T cell count after chemotherapy was 540.7 /uL (SD ± 266.1). CD4+T cell count was significantly reduced after chemotherapy ($z=9.164$; $p<0.001$). **Conclusion:** from our study we conclude that, CD4 and CD8+ T cell number and their ratio as an assessment of immune function status in breast cancer patients, before and after surgery and adjuvant chemotherapy with the intention of delineating any potential immune aberrations that may result from therapy.

Keywords: breast cancer, CD4/CD8 ratio, chemotherapy.

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INTRODUCTION

Breast cancer incidence and mortality rates remain highest in developed countries compared with developing countries. The incidence is high in North America and Northern European countries, intermediate in Southern European and Latin American countries and low in most Asian and African countries [1]. In the western world breast cancer accounts for 27% of all female cancer and one out of fourteen can expect to develop breast cancer in their life time [2].

In the United States, each year approximately 100,000 new cases are diagnosed and approximately 30,000 patients die of the disease [3]. However, in some regions of the world (North America, Western Europe and Australia) breast cancer mortality is finally beginning to fall, presumably because of the combined action of earlier diagnosis and improved therapy [4].

In countries like India, Japan (though developed), China and those of Africa, the highest recorded incidence rates from the various regional

registries for the year 1990, are 23.5, 33.4, 26.5 and 29.5 per 100,000 respectively; which are more than 50% lower than the few 'low' rates of high-risk populations. Despite the low incidence rates, almost 40% of the 1990 global cancer burden and 45% of deaths due to the disease occurred in the developing countries. In India it is the second most common cancer after cervix, accounting for 19% of the total cancer burden in women. The incidence rate in Asian women has been rising and is associated with a shift towards a more westernized lifestyle. The Bombay Cancer registry has also recorded a consistent rise in the incidence of breast cancer, which has now overtaken cervical cancer, to be the leading cancer in women of Mumbai [5].

In this study our main aim is to evaluate CD4 and CD8 count in breast cancer patient undergoing post-Surgery FAC protocol.

OBJECTIVE

General objective

- To evaluate CD4 and CD8 count and their ratio in breast cancer patient undergoing post-surgery FAC protocol.

Specific objective

- To detect CD8+T cell count before and after chemotherapy.
- To identify histological grading of the patients.

METHODOLOGY

Study type: This was a quasi-experimental study.

Place and period of study: This study was carried out at Dept. of Microbiology, Department of radiotherapy & Department of Surgery, Sylhet MAG Osmani Medical College Hospital from January, 01, 2011 to December, 31, 2011.

Sampling technique: Purposive sampling

Inclusion Criteria

All female diagnosed patients of breast carcinoma –

- Of 25 – 65 years of age.
- Planned for adjuvant FAC protocol after surgery.

Exclusion Criteria

- H/o any complication in breast surgery, which causes delayed wound healing.
- Malignancy complicated with other chronic diseases.
- H/o any immunosuppressive disease, like DM, Rheumatoid arthritis.
- Patients not willing to participate in the study.
- H/O taking any myeloid growth factors (Granulocyte colony stimulating factor, granulocyte-macrophage colony stimulating factor).

Study Design

- A total number of thirty breast cancer patients were enrolled for the study, and from them consecutive two blood samples were taken. The first sample was taken before surgery, when they were admitted into surgery ward. And the next sample was taken

after surgery and the third cycle of the scheduled six cycled adjuvant FAC (5-Fluouracil, Adriamycin, Cyclophosphamide) protocol chemotherapy .Samples were selected purposively, according to inclusion and exclusion criteria.

Data Collection Procedure

Samples from the cases were collected, first from Department of surgery (before operative procedure) and then, from Department of radiotherapy (after surgery and 3 cycles of adjuvant chemotherapy) from the same patient. Prior ethical permission was obtained from the E.C. of Sylhet MAG Osmani Medical College. Written informed consent was obtained from each patient Detailed history from the patient was taken.

Statistical Analysis

All data will be checked and analyzed with the help of SPSS (statistical package for social sciences) with version 12.0. Statistical significance will be calculated using paired T test.

RESULTS

In table-1 shows age distribution of the patients where 14 (46.7%) patients were in the age group of 41 to 50 years, 9 (30.0%) patients were in the age group of 31 to 40 years, 4 (13.3%) patients were in the age group of 51 to 60 years, 2 (6.7%) patients were in the age group up to 30 years and 1 (3.3%) patient was in the age group above 60 years. The following table is given below in detail:

Table-9: Distribution of respondents by age group (n=30)

Age group	Frequency	Percentage
≤30 years	2	6.7
31-40 years	9	30.0
41-50 years	14	46.7
51-60 years	4	13.3
>60 years	1	3.3

In figure-1 shows distribution of patients according to side of involvement. Left breast was involved in 17 (53.3%) patients and right breast was involved in 13 (43.3%) patients. The following table is given below in detail:

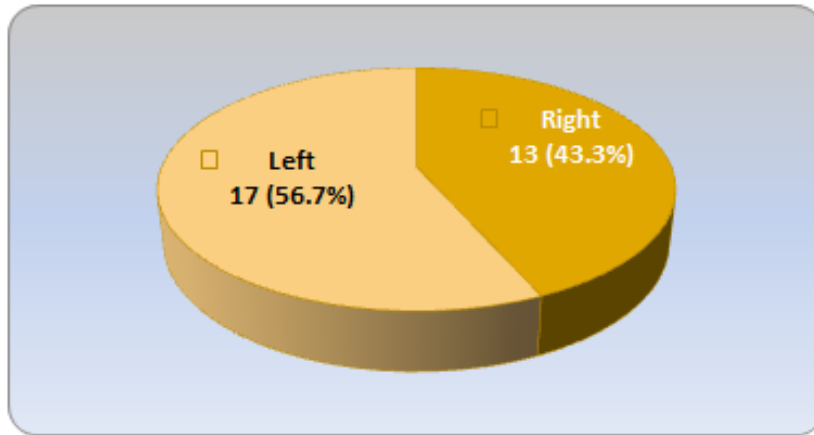


Fig-1: Distribution of patients according to side of involvement (n=30)

In figure-2 shows distribution of respondents according to histological grading. Regarding histological grading was grade-II in 13 (43.3%) cases,

grade-I in 11 (37.7%) and grade-III in 6 (20.0%) cases. The following figure is given below in detail:

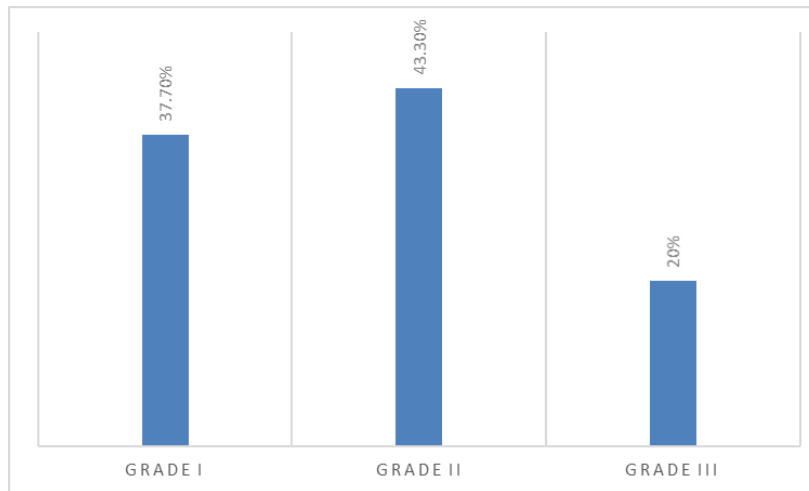


Fig-2: Distribution of respondents according to histological grading (n=30)

Table-2 showed the comparison of CD4+T cell count before and after chemotherapy. CD4+T cell count before chemotherapy was 1023.4 /uL (SD ± 320.7) and CD4+T cell count after chemotherapy was 540.7 /uL

(SD ± 266.1). CD4+T cell count was significantly reduced after chemotherapy (z=9.164; p<0.001). The following table is given below in detail:

Table-2: Comparison of CD4+T cell count before and after chemotherapy (n=30)

Parameter	CD4+T cells /uL		*p-value
	Before Chemotherapy	After Chemotherapy	
Mean	1023.4	540.7	p<0.001
SD	320.7	266.1	
Range	476-1875	121-1081	

SD=Standard deviation

*Pair-t test was applied to analysed the data.

Table-3 showed the comparison of CD8+T cell count before and after chemotherapy. CD8+T cell count before chemotherapy was 669.1/uL (SD ± 437.5) and

CD8+T cell count after chemotherapy was 318.5 /uL (SD ± 212.7). CD8+T cell count was significantly reduced after chemotherapy (Z=4.418; p<0.001).

Table-3: Comparison of CD8+T cell count before and after chemotherapy (n=30)

Parameter	CD8+T cells / uL		*p-value
	Before Chemotherapy	After Chemotherapy	
Mean	669.1	437.5	p<0.001
SD	318.5	212.7	
Range	293-1707	110-821	

SD=Standard deviation

*Pair-t test was applied to analysed the data.

DISCUSSION

Site of lesion of the present study is 56.7% (17 patients) in left and 43.3% (13 patients) in right breast involvement, which is consistent with the studies done by others. One study found that the tumor were located 52% in the left side and 48% in the right side of breast [6]. Another study found 52.2% in left side and 47.8% in right side of their study patients [7]. But other found in his observation that 76% in right side and 24% in left side of breast carcinoma patients [8] which disagree with other study[9].

In this study, histological grades were as follows- 37.7% Grade - I, 43.3% Grade - II & 20.0% Grade- III. But various Authors show various information. As Malik I A (2002) found that 12% Grade - I, 53% Grade - II & 35% Grade - III. One report described in their study as 6% Grade - I, 61% Grade- II & 33% Grade – III [10]. Another report found in their study, 5.3% Grade - I, 64% Grade - II & 30.7% Grade - III [11]. These variations occur due to geographical presentation, genetic involvement, sample size and also for study design. In this study, family history of breast cancer was positive in 26.7% cases and no family history found in 73.3% cases.

In the study discussed here, CD4+T cell count before surgery & chemotherapy was 1023.4 /uL (SD \pm 320.7) and CD4+T cell count during chemotherapy was 540.7 /uL (SD \pm 266.1). CD4+T cell count was significantly reduced during chemotherapy than before ($z=9.164$; $p<0.001$). And the CD8+T cell count before surgery and chemotherapy was 669.1/uL (SD \pm 437.5) and CD8+T cell count during chemotherapy was 318.5 /uL (SD \pm 212.7). CD8+T cell count was significantly reduced during chemotherapy ($Z=4.418$; $p<0.001$). One study showed the CD4+T cell decreased from a mean of 588/mm³ before chemotherapy to 105 /mm³ during chemotherapy ($P= .0002$) and C8+ T cells decreased from a mean of 382 /mm³ before chemotherapy to 150 /mm³ during chemotherapy ($P = .0009$) [10]. In our study, we were working on the patients undergoing adjuvant FAC protocol. The chemotherapy protocol was specific for all patients. But in Mackall *et al.* [11], the protocol was not specific; three different protocols were used there. Moreover, the patients also were not specific, three different categories of tumors studied there (brain tumour, sarcoma and NHL). But in our study, all patients were of breast carcinoma.

CONCLUSION

From our study we conclude that, CD4 and CD8+ T cell number and their ratio as an assessment of immune function status in breast cancer patients, before and after surgery and adjuvant chemotherapy with the intention of delineating any potential immune aberrations that may result from therapy.

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