

A STUDY ON ROLE OF PROSTATE SPECIFIC ANTIGEN IN CARCINOMA BREAST

Anand Arumugam¹, N. Naveethalakshmi², K. Rajachidambaram³, C. Divyah Manogari⁴, Samuel R Rakesh⁴

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Corresponding Author:

Dr. N. Naveethalakshmi

Email: drnaveetha@gmail.com

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¹Associate Professor, Department of General Surgery, Dhanalakshmi Srinivasan Medical college Hospital, Siruvachur, Perambalur, Tamil Nadu, India.

²Associate Professor, Department of Biochemistry, Dhanalakshmi Srinivasan Medical college Hospital, Siruvachur, Perambalur, Tamil Nadu, India

³Professor, Department of General Surgery, Dhanalakshmi Srinivasan Medical college Hospital, Siruvachur, Perambalur, Tamil Nadu, India.

⁴Post graduate, Department of General Surgery, Dhanalakshmi Srinivasan Medical college, Perambalur, Tamil Nadu, India.

Abstract

Background: To compare the level of serum prostate specific antigens in patients with carcinoma of breast with normal standardized level and to compare the preoperative and postoperative serum PSA level in patients with carcinoma breast. **Materials and Methods:** This is a prospective randomized control study with inclusion criteria of 50 patients presenting with lump in the breast which proven to be carcinoma through tissue diagnosis and exclusion criteria of patients with lump over breast which proven to be benign through tissue diagnosis and patients with associated ovarian and uterine pathology. This study was conducted in the Department of General Surgery, Dhanalakshmi Srinivasan Medical College & Hospital, and Perambalur for a period of eighteen months. **Result:** Total Number patients enrolled in the study – 50. After statistical analysis, the conclusion made that, there is no significant correlation between serum PSA level and carcinoma breast, The Mean serum PSA level in patients with carcinoma breast was found very low when compared to expected level. The mean serum PSA level between pre-neoadjuvant and post-neoadjuvant has no significant differences, the mean serum PSA level between pre- surgical and post-surgical period has no significant differences. Factors that affect the transport of PSA from tissue to blood may also be considered at this point and also the tumor behavior of the westerner and Asians may be considered for its significant change of PSA. **Conclusion:** There is no significant correlation between serum PSA level and carcinoma breast and no significant difference between Pre surgical and post-surgical serum PSA level in patients with carcinoma breast.

INTRODUCTION

Prostate specific antigen is unique for prostate epithelium numerous studies have demonstrated that female tissue such as breast, endometrium, and ovary are also produce PSA which is similar to prostate since their differentiation and growth are under the control of steroid hormones and PSA is found to be secreted in breast milk of lactating mother and nipple aspirate. Mammary PSA having identical molecular weight and m RNA sequences of seminal PSA. PSA gene expression in breast malignancy found to be under hormonal control since steroid hormone receptor positive breast tumor cell lines T-47D and BT-474 are stimulated by glucocorticoids, mineralocorticoids, progestin's and androgens, hence some amount PSA always will be Present in female serum in the range of 0.1-0.9 ng/lit. The aim of this

study is to analyze the level of serum PSA level in patients with Carcinoma breast and to know its correlation with carcinoma breast.^[1-5]

Aims & Objectives

To compare the level of serum prostate specific antigen in patients with carcinoma breast with normal standardized level and to compare the preoperative and postoperative serum PSA level in patients with carcinoma breast.

MATERIALS AND METHODS

Patients presenting with clinical features of lump in the breast, admitted as in-patient in Department of General Surgery, Dhanalakshmi Srinivasan Medical college from January 2022 to June 2023 will be enrolled in our Prospective randomized control study.

Inclusion Criteria: 50 patients presenting with lump in the breast which proven to be carcinoma through tissue diagnosis.

Exclusion Criteria: Patients with lump over breast which proven to be benign through tissue diagnosis and patients with associated ovarian and uterine pathology.

RESULTS

This study was conducted in the Department of General Surgery, Dhanalakshmi Srinivasan Medical College & Hospital, Perambalur for a period of eighteen months. Patients, who fulfilled the inclusion criteria, were enrolled in this study, after obtaining an informed consent. Total Number patients enrolled in the study – 50.

Statistical Analysis: Group Statistic.

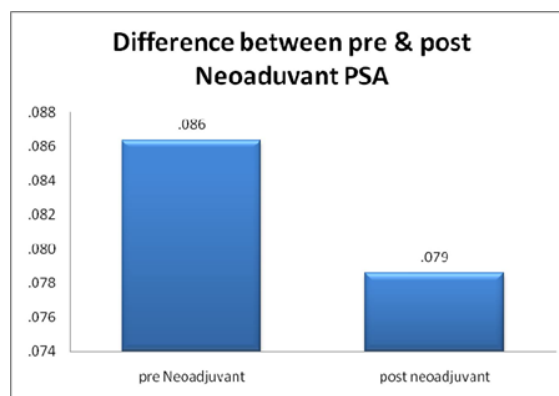


Figure 1: graphical description of mean differences

Table 1: T-Test Group Statistics

PN	N	Mean	Std. Deviation	Std. Error Mean
DIFF Pre-Neoadjuvant	22	.086	.0560	.0119
Post-neoadjuvant	28	.079	.0738	.0140

Table std. deviation and std.mean error of pre neoadjuvant and neoadjuvant

Table 2: Independent Sample Test

	Equality of Variances		t-test for Equality of Means						
	F	Sig.	t	df	Sig. (2-tailed)	Mean Difference	Std. Error Difference	of the Difference	
								Lower	Upper
DIFF Equal Variances assumed	4.125	.048	.411	48	.683	.0078	.0190	-.0304	.0460
DIFF variances not assumed			.424	47.959	.673	.0078	.0184	-.0291	.0447

$P > 0.01$, no significant differences in PSA level between pre neoadjuvant and neoadjuvant

Table 3: Paired Samples Test

	Paired Differences					t	df	Sig. (2-tailed)
	Mean	Std. Deviation	Std. Error Mean	of the Difference				
				Lower	Upper			
Pair 1 PSAPREM RMINgml - PSAPOST MRMINgml	.0036	.1201	.0227	-.0430	.0502	.157	27	.876

a. PN = Neo Adjuvant

$P > 0.01$, no significant differences between PSA level in pre MRM and post MRM

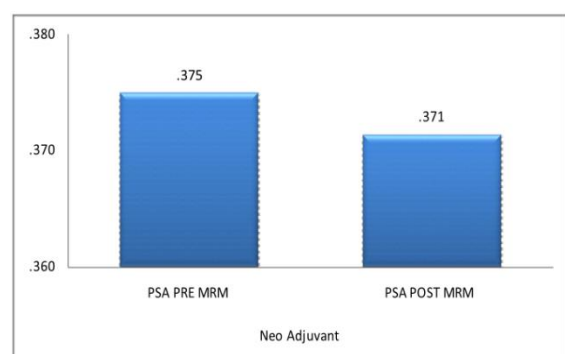


Figure 2: Mean distribution between pre-MRM and post MRM

DISCUSSION

Prostate specific antigen [PSA] is a tumor marker used widely for the diagnosis and monitoring of prostatic adenocarcinoma. The PSA Positivity rate was 28% in the group of all cancer patients. 33% in patients under the age of 50 and 26% in patients at

the age of 50 or older. PSA positive tumor were found in 34% of stage I, 24% of stage II, 18% of stage III and stage IV disease. These findings suggest that PSA production in these tissues may be regulated by mechanism which involve derangement of balance between the various steroid hormone and their receptors and also expression of non-functional receptors or deranged post-receptor pathway. Based on the information presented, PSA can now be regarded as a molecule secreted by tissue in malignant diseases. Studies shown that PSA concentration in cytosol extract has a favorable prognostic indicator in breast cancer, Serum PSA level of breast cancer patients were compared with standardized normal level and pre surgical and postsurgical levels are also been compared, there is no significant correlation between serum PSA level and carcinoma breast and no significant difference between Presurgical and post-surgical serum PSA level.^[6-9]

CONCLUSION

This study conducted in an attempt to know, if serum PSA measurement in Female patients with carcinoma breast have any diagnostic, prognostic or monitoring value. Serum PSA level of breast cancer patients were compared with standardized normal level and pre surgical and post-surgical levels are also been compared. After statistical analysis, the conclusion made that, there is no significant correlation between serum PSA level and carcinoma breast and no significant difference between Pre surgical and post-surgical serum PSA level in patients with carcinoma breast. In prostate, PSA enters the circulation by physical diffusion. Factors that affects the transport of PSA from tissue to blood may also be considered at this point and also the tumor behavior of the westerner and Asians may be considered for its significant change of PSA.

REFERENCES

1. Libson S, Lippman M. A review of clinical aspects of breast cancer. *International review of psychiatry (Abingdon, England)* 2014;26(1):4–15. doi: 10.3109/09540261.2013.852971. - DOI - PubMed
2. Basile D, Cinausero M, Iacono D, Pelizzari G, Bonotto M, Vitale MG, Gerratana L, Puglisi F. Androgen receptor in estrogen receptor positive breast cancer: beyond expression. *Cancer Treat Rev.* 2017;61:15–22. doi: 10.1016/j.ctrv.2017.09.006. - DOI - PubMed
3. Fujii Rika, Hanamura Toru, Suzuki Takashi, Gohno Tatsuyuki, Shibahara Yukiko, Niwa Toshifumi, Yamaguchi Yuri, Ohnuki Koji, Kakugawa Yoichiro, Hirakawa Hisashi, Ishida Takanori, Sasano Hironobu, Ohuchi Noriaki, Hayashi Shin-ichi. Increased androgen receptor activity and cell proliferation in aromatase inhibitor-resistant breast carcinoma. *The Journal of Steroid Biochemistry and Molecular Biology.* 2014;144:513–522. doi: 10.1016/j.jsbmb.2014.08.019. - DOI - PubMed
4. Hanamura T, Hayashi SI. Overcoming aromatase inhibitor resistance in breast cancer: possible mechanisms and clinical applications. *Breast cancer. Japan: Tokyo; 2017.* - PubMed
5. Hayashi S, Kimura M. Mechanisms of hormonal therapy resistance in breast cancer. *Int J Clin Oncol.* 2015;20(2):262–267. doi: 10.1007/s10147-015-0788-5. - DOI - PubMed
6. De Amicis F, Thirugnansampanthan J, Cui Y, Selever J, Beyer A, Parra I, Weigel NL, Herynk MH, Tsimelzon A, Lewis MT, Chamness GC, Hilsenbeck SG, Ando S, Fuqua SA. Androgen receptor overexpression induces tamoxifen resistance in human breast cancer cells. *Breast Cancer Res Treat.* 2010;121(1):1–11. doi: 10.1007/s10549-009-0436-8. - DOI - PMC - PubMed
7. D'Amato NC, Gordon MA, Babbs B, Spoelstra NS, Carson Butterfield KT, Torkko KC, Phan VT, Barton VN, Rogers TJ, Sartorius CA, Elias A, Gertz J, Jacobsen BM, Richer JK. Cooperative dynamics of AR and ER activity in breast Cancer. *Molecular cancer research : MCR.* 2016;14(11):1054–1067. doi: 10.1158/1541-7786.mcr-16-0167. - DOI - PMC - PubMed
8. Hickey TE, Robinson JL, Carroll JS, Tilley WD. Minireview: the androgen receptor in breast tissues: growth inhibitor, tumor suppressor, oncogene? *Molecular endocrinology (Baltimore, Md)* 2012;26(8):1252–1267. doi: 10.1210/me.2012-1107. - DOI - PMC - PubMed
9. Barton VN, D'Amato NC, Gordon MA, Lind HT, Spoelstra NS, Babbs BL, Heinz RE, Elias A, Jedlicka P, Jacobsen BM, Richer JK. Multiple molecular subtypes of triple-negative breast cancer critically rely on androgen receptor and respond to enzalutamide in vivo. *Mol Cancer Ther.* 2015;14(3):769–778. doi: 10.1158/1535-7163.mct-14-0926. - DOI - PMC - PubMed