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Biochemistry

MEASURE OF SERUM MDA, GST AND VIT-C IN WOMEN WITH CERVICAL CARCINOMA

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Cervical carcinoma (CaCx) is the second most common gynecological cancers globally. Many authors revealed that low levels of antioxidants induce the overproduction of reactive oxygen species (ROS) that result in oxidative stress. In the present study an attempt was made to evaluate the oxidative stress in 20 women with cervical carcinoma by measuring serum Malondialdehyde (MDA), Glutathione-S-transferase (GST) and Ascorbic acid (Vit-C) levels which was compared with 20 age matched non cancerous women. Significantly elevated MDA levels (P<0.001) and lowered GST, Vit-C levels (P<0.0001) were observed in women with cervical carcinoma as compared with healthy controls. From our findings it can be concluded that oxidative stress plays an important role in pathogenesis of cervical carcinoma. Hence supplementation of antioxidants may reduce the oxidative stress and minimize the associated morbidity and mortality.

KEYWORDS: Cervical carcinoma (CaCx), Reactive oxygen species (ROS), Malondialdehyde (MDA), Glutathione-Stransferase (GST) and Ascorbic acid (Vit-C).

INTRODUCTION

Cervical carcinoma is the most prevalent genital tract cancer and is one of the leading causes of deaths due to cancer among women world wide and in India. Several risk factors of cervical cancer such as smoking, oral contraceptives, immune-suppression and infection with HPV are well documented to increase oxidative stress ^(1,2). During infection, free radicals i.e pro-oxidases are generated by which antioxidants are depleted leading to development of oxidative stress ⁽³⁾. The imbalance between these pro-oxidants and anti-oxidants called oxidative stress is assessed by Malondialdehyde (MDA) a lipid peroxidation product ^(4,5).

Glutathione-S-transferase, a secondary antioxidant enzyme helps in the detoxification of ROS by decreasing peroxide levels or it maintains the steady supply of metabolic intermediates glutathione reductase for the primary antioxidant enzymes⁽⁶⁾. Many studies have evidenced an altered expression of GST in plasma and also in tissue biopsy samples of CaCx ⁽⁷⁾

Ascorbic acid reacts with ROS, converting them into semi hydro-ascorbate radicals, reducing the risk of cancer by suppressing free radicals and oxidative stress⁽⁸⁾. Recent studies have shown that ascorbic acid molecule has a direct impact on cervical carcinogenesis⁽⁹⁾.

MATERIALS AND METHODS

The present study was conducted on 40 women in the department of Biochemistry, among them 20 were diagnosed with Cervical carcinoma aged between 35-65 years, selected from the department of Gynecology, Government Maternity Hospital and Sri Venkateswara Medical College, Tirupati and other 20 were age matched healthy non-cancerous controls. written consent was procured from every participant included in the study.

5ml of venous blood was collected from all the participants with aseptic precautions into plain tubes and centrifuged. The obtained serum was analyzed for Malondialdehyde (MDA), Glutathione-Stransferase (GST) and Ascorbic acid (VitC) spectrophotometrically. MDA is determined as a measure of Thiobarbituric acid reactive substances. (TBARS)⁽¹⁰⁾, GST activity was measured using 1, chloro 2, 4 dinitrobenzene as the substrate (CDNB)⁽¹¹⁾ and vitC estimation by Roe,J.H., and Kuether, C.A method⁽¹²⁾.

The data obtained from both groups was expressed as mean \pm S.D., students t-test was used for statistical analysis and p-value < 0.01 for statistical significance as shown in table 1.

Table 1: Mean and SD values of MDA, GST and Vit C in cases and controls

S.no	Parameters	Means ± SD		P Value
		Cases (n:20)	Controls(n:20)	
1	MDA (nmol/L)	5.69 ± 1.61	3.34 ± 0.93	< 0.001
2	GST (IU/L)	36.84 ± 17.17	72.73 ± 19.33	< 0.001
3	Vit C (mg/dl)	0.2 ± 0.07	1.07 ± 0.56	< 0.001

There was a significant raise in MDA levels (P<0.001) and a significant fall in GST and Vit C levels (P<0.001) among CaCx cases when compared with controls.

DISCUSSION

Oxidative stress is an imbalance between production of ROS in cells and tissues and the capability of the biological mechanisms to detoxify these reactive products (13,16).

Oxidative stress plays an important role in development of carcinogenesis by generating free radicals (or) by reducing antioxidant levels (14,16).

In our study an attempt was made to assess the extent of oxidative stress in terms of lipid peroxidation product MDA, antioxidant enzyme Glutathione-S Transferase activity and the antioxidant vitamin, Ascorbic acid levels in women with cacx. Significantly elevated MDA levels (P<0.001) (4.5.15.16), and lowered GST and Ascorbic acid levels (P<0.001) This confers that low levels of antioxidants in these patients may be due to their over use in scavenging lipid-peroxides as well as sequestration by tumor cells.

CONCLUSION

Our study concludes that oxidative stress plays a significant role in the pathogenesis of CaCx and it emphasizes that there is a need for supplementation of Antioxidants which may reduce the oxidative stress and minimize the associated morbidity and mortality.

REFERENCES

- S.S Beevi, M.H. Rashee, A. Geetha Evidence of Oxidative and nitrosative stress in patients with cervical squamous cell carcinoma, clin. chim. Acta, 375 (2007), PP. 119-213.
- T. Finkel. oxidant signals and oxidative stress cuss. opin. cell Biol., 15 (2003), PP. 247-254
- Ebrahimi, S, Soltani, A, Hashemy, SI. Oxidative stress in cervical cancer pathogenesis and resistance to therapy. J Cell Biochem. 2019;120:6868-77.
- 4. Naidu, M. S. K., Suryakar, A. N., Swami, S. C., Katkam, R. V., & Kumbar, K. M

- Oxidative stress and antioxidant status in cervical cancer patients. Indian Journal of Clinical Biochemistry, (2007). 22(2), 140-144.
- Gutteridge JMC. Lipid peroxidation and antioxidants as biomarkers of tissue damage. Clin. Chem. 1995;41:1819-28.
- Jelic M, Mandic A, Kladar N, Sudji J, Bozin B, Srdjenovic B. Lipid Peroxidation, Antioxidative Defense and Level of 8-hydroxy-2-deoxyguanosine in Cervical Cancer Patients. J Med Biochem. 2018 Jul;37(3):336-45.
- Manjur, Balasubramanian, Nalini N. Oxidative stress and tumor markers in cervical cancer patients. J Biochem Mol Biophys. 2002 Dec;6(6):387-90. 7.
- A.C Carr and B. Frei. Does Vit. C as pro-oxidant under physiological conditions. FASEB Journal. 1999;13:1007-24.
- Journal, 1999;13:1007-24. Hwang JH, Kim MK, Lee JK, Dietary supplements reduce the risk of cervical intra epithelial neoplasia. Int. J Gynecol cancer. 2010; 20(3): 398-403 Draper HH, Hadley M. Malondialdehyde determination as index of lipid peroxidation. Methods Enzymol. 1990; 186:421 Weissman S. Red Cell Metabolism. A Manual of Biochemical Methods. 2nd Edition The

- Welssinari S. Red Cell Metatoristic. 1976 Jul;49(3):310-11.

 Roe, JH and Kuether, C.A.: J. Bio I. chem., 147: 399, 1943

 Gabriele Pizzino, Natasha Irrea, Mariapaola Cucinotta, et al. Oxidative stress:Harms and Benefits for Human Health. Oxid Med Cell Longevity. 2017 Jul: 8416763

 Sharhan S, Norman H, Fatimah A, et al. Antioxidant intake and status and oxidative stress in relations to breast cancer risk: A case control study. Asian Pac. J, Cancer pre-2008: 9: 343-50.
- 2008: 9; 343-50.
 Gonçalves TL, Erthal F, Corte CL, Müller LG, Piovezan CM, Nogueira CW,et al. Involvement of oxidative stress in the pre-malignant and malignant states of cervical cancer in women. Clin Biochem. 2005;38 (12):1071–75.
 Shah SR, Shaheen B Shaikh, Shaheena Yassir. Altered Total Antioxidant Capacity and
- Malondialdehyde in Cervical Cancer Patients and Effect of Chemoradiation. Sch Int J Biochem. 2021;4(2): 14-19