

## Status of Lipid Peroxidation and Role of Serum Zinc & Copper In Subjects With Hypothyroidism

Santasmitta Pal<sup>1</sup>, Sumit Patra<sup>2</sup>, Tapan Mukhopadhyay<sup>3</sup>

<sup>1</sup>Assistant Professor, Dept. of Biochemistry, Medical College, Kolkata

<sup>2</sup> Ex PGT, Dept. of Biochemistry, Medical College, Kolkata

<sup>3</sup>Professor & Head, Dept. of Biochemistry, Medical College, Kolkata

Corresponding Author: Santasmitta Pal

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**Abstract:** Hypothyroidism is a common endocrinopathy which remains associated with free radical generation resulting in oxidative damage to various organs. Zinc and Copper are two essential trace elements for the human body, principally because of its role in antioxidant mechanism. Since oxidative damage has been reported to be related to the development of complications of hypothyroidism, it is prudent to identify the condition, in an attempt to retard the progress of the complications of hypothyroidism. **Aims & objectives:** To estimate serum Malondialdehyde (MDA), Zinc and Copper levels in Primary hypothyroid subjects and compare it with euthyroid subjects.

**Materials and methods:** A cross sectional observational study was conducted in Dept. of Biochemistry in collaboration with Dept. of Endocrinology, Medical College & Hospital, Kolkata from January 2015 to January 2016. The study recruited 75 primary hypothyroid subjects and 75 age and sex matched healthy controls. Serum TSH and fT4 was measured by Chemiluminescence method, while MDA, Zinc and Copper were measured in spectrophotometer.

**Results:** Serum MDA was significantly higher in Primary hypothyroid ( $8.00 \pm 3.95$ ) nmol/ml in comparison to controls ( $1.88 \pm 0.58$ ) nmol/ml. Values of serum zinc and Copper found to be significantly lower in Primary hypothyroid cases compared to euthyroid subjects. ( $p$  value  $< 0.05$ ).

**Conclusion:** In this study it was observed Serum MDA was significantly higher in Primary Hypothyroid subjects along with decreased Zinc and Copper level. Considering the results of this study as well as those done earlier, underlying oxidative damage needs to be restricted. The potential benefits of supplementing Zinc and Copper in Primary Hypothyroid subjects needs to be evaluated further.

**Key words:** MDA, Zinc, Copper, Primary Hypothyroid.

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### I. Introduction

Hypothyroidism is a very common Endocrinopathy worldwide including India where about 42 million Indians are reported to be suffering from spectrum of thyroid disorder, mainly hypothyroidism (1). The physiological effects of thyroid hormones is primarily related to functions of all the tissues of body involving modulation of gene transcription resulting in alteration of basal metabolic rate and energy metabolism affecting carbohydrate, lipid and amino acid metabolism (2).

Free radicals are highly reactive molecules generated by biochemical redox reactions that occur as a part of normal cell metabolism. Free radicals may contribute towards lipid peroxidation and damage macromolecules and cellular structure of the organism, endothelium and erythrocytes. Serum Malondialdehyde (MDA) is the breakdown product of the major chain reactions leading to oxidation of polyunsaturated fatty acids and thus serves as a reliable marker of lipid peroxidation (3,4). Free radicals are eliminated from the body by their interaction with various non-enzymic and enzymic antioxidants such as uric acid, albumin, bilirubin, vitamins E, C, A, Glutathione peroxidase, Superoxide dismutase (SOD) and Catalase (5). Previous clinical and experimental studies showed an altered free radical level (with different results) in hypothyroidism. Some of the studies showed an increase (6,7) while some other showed a decrease (8) or no significant differences (9, 10). Micronutrients like zinc, copper and selenium are also important molecules involving in removal of free radicals through various mechanism namely antioxidant enzyme like Superoxide dismutase, Catalase, Peroxidase etc. Defence against free radical toxicity is related to zinc as a component of SOD (11). There are reports that transport of zinc from the intestine & renal tubules are dependent on thyroid hormones as observed in rat model. (12) Copper in excess is found to be related to oxidative injury in other study (13).

The aim of this study was to investigate the effect of hypothyroidism on lipid peroxidation and evaluate possible links between trace element concentrations (copper, zinc) and thyroid hormones in comparison to euthyroid subjects.

## II. Materials And Methods

This cross-sectional observational study was conducted at Department of Endocrinology and Department of Biochemistry, Medical College & Hospital, Kolkata from January 2015 to January 2016 after obtaining necessary permission from Institutional Ethics Committee.

Primary Hypothyroidism is very common where there is low FT4 and high TSH. Secondary Hypothyroidism is less common variety where TSH & FT4 all are low. Primary hypothyroid subjects in the age group between 18-70 years and age and sex matched normal subjects were selected after obtaining informed consent from all the subjects. Each individual enrolled in the study underwent a detailed history, clinical examination and laboratory investigation designed for the study. The selected cut-off value for accepting subjects under the study on Hypothyroidism was: TSH above 10.0  $\mu$ IU/ml and FT4 below 0.8 ng/dl. (Williams). Patients suffering from Diabetes Mellitus, chronic renal diseases, chronic infection, pregnancy and patients taking Antioxidant, Zinc or Copper supplementation were excluded from the study. Five ml of fasting blood sample was collected from each individual in sterile plain vial. The blood was allowed to clot and then centrifuged at 3000 rpm for 10 minutes. Serum was separated and stored at  $-20^{\circ}\text{C}$  until analysis. Serum TSH and FT4 were measured by Chemiluminescence method (Siemens Immulite 1000). Serum Zinc and Copper was estimated using commercial kits in semiautoanalyzer (ERBA Chem V+ v2). Serum MDA assay based on its reaction with thiobarbituric acid by SATHO and others (14) in double beam UV-VIS Spectrophotometer (EC-PCI). Data was analysed using Microsoft office Excel and SPSS 20.0 software.

## III. Result

**Table 1 : Determination of central tendency of MDA, Zn and Cu level in test and control groups:**

Biochemical Tests		Test group	Control group
		N=75	N=75
MDA	Mean $\pm$ SD	8.00 $\pm$ 3.95	1.88 $\pm$ 0.58
	Median	6.60	1.98
	IQR	4.2	0.85
Zn	Mean $\pm$ SD	47.59 $\pm$ 32.88	149.68 $\pm$ 71.67
	Median	37.62	112
	IQR	56.66	100
Cu	Mean $\pm$ SD	105.58 $\pm$ 67.20	130.69 $\pm$ 30.44
	Median	73.56	134
	IQR	114.40	24

SD= Standard deviation, IQR= Inter quartile range

Table 1 describes the central tendency of MDA, Zinc & Copper levels in the two groups studied.

**Table 2: Test of normality for data of MDA, Zn and Cu level:**

	Kolmogorov-Smirnov <sup>a</sup>			Shapiro-Wilk		
	Statistic	df	Sig.	Statistic	df	Sig.
MDA	0.181	150	0.000	0.818	150	0.000
Zn	0.164	150	0.000	0.883	150	0.000
Cu	0.088	150	0.006	0.950	150	0.000

a. Lilliefors Significance Correction

The results of both the normality tests, Kolmogorov-Smirnov and Shapiro-Wilk, are significant for all three data sets, i.e. MDA, Zn and Cu levels. These indicate that, these data are not normally distributed. So, parametric test like unpaired t-test were not done to compare the values of these three parameters between test and control groups. Instead, nonparametric test (MANN WHITNEY U TEST) was done.

**Table 3: Hypothesis test summary for MDA, Zn and Cu level across test and control groups:**

Null Hypothesis	Test	Sig.	Decision
1. The distribution of MDA is the same across categories of group	Independent-sample Mann-Whitney U Test	0.000	Reject the null hypothesis
2. The distribution of Zn is the same across categories of group	Independent-sample Mann-Whitney U Test	0.000	Reject the null hypothesis
3. The distribution of Cu is the same across categories of group	Independent-sample Mann-Whitney U Test	0.000	Reject the null hypothesis

**The significance level is 0.05**

As the data is not normally distributed as previously shown in Table 2, Nonparametric test (Mann-Whitney U test) was performed to check whether there is significant difference between two groups. It has been found that there is significant difference in distribution of MDA, Zn and Cu levels between the two groups (p value <0.05).

**IV. Discussion**

The data presented in this study show that lipid peroxidation is markedly higher in hypothyroidism than euthyroid subjects (p value < 0.05). In previous studies, different interpretations were given. Venditti et al. (15) in an experimental study on hypothyroid rats showed that Malondialdehyde (MDA) levels did not differ significantly from euthyroid values. Mano et al. (16) found that the concentration of lipid peroxides did not change in hypothyroid rats in comparison with euthyroid animals. Gredilla et al. (17) demonstrated that in vivo and in vitro lipid peroxidation was not altered in the hypothyroid state. Dumitriu et al. (18) showed that the mean malondialdehyde level was significantly higher in hypothyroid patients in comparison to the control group. Yilmaz et al. (19) showed that MDA level of hypothyroid rats was increased in liver, but they were decreased in the tissues of the heart and thyroid. Sawant *et al.* (20) demonstrated that the tissue lipid peroxidation level significantly increased in hypothyroid rats. The result of our study is in conjunction with the results of studies showing that the level of MDA is significantly increased (17, 21). But present results are not in agreement with the other studies (10,16,17,19). The increase in reactive oxygen species induced by thyroid hormone may contribute towards an oxidative stress condition in some tissues with a consequent lipid peroxidative response. Possible sources of elevated free radicals in hypothyroid patients include increased production of radical oxygen species, especially from lipid peroxidation processes and probably decreased antioxidant defence systems.

In our study significant low Zinc levels were observed in hypothyroid subjects in comparison to euthyroid subjects. This is in agreement with other studies. (22,23). This could be possibly caused due to either impaired gastrointestinal absorption of zinc or altered zinc distribution leading to sequestration of zinc in liver and other tissues in hypothyroid subjects. Oliveri et al (24) reported that thyroid hormones did not correlate with indices of zinc status in hypothyroid subjects; although they have observed decreased iodothyronine levels in case of zinc deficiency.

Copper deficiency can exert both a direct effect on the metabolic process and an indirect one disturbing iodine metabolism, and sharply decreasing protein-bound iodine production by the thyroid gland. Our results showed statistically significant decreased serum Cu levels in patients with hypothyroidism when compared with control group (P <0.05). The finding is in agreement with Alturfan et. al, (25), Akcay et. al, (26). But, it is not in conjunction with Aihara et. al, (27) which they found that no significant difference in plasma Cu concentrations between control subjects and patients with thyroid disease, except a higher significant difference in patients with hyperthyroidism. The cause behind increased serum Cu concentration in hyperthyroidism and decreased in hypothyroidism may be attributed to the fact that the most plasma Cu (approximately 93%) is bound to ceruloplasmin and small fraction to albumin (6–7%) or is chelated to amino acids (< 1%), which is diffusible[20,21]. Thyroid hormones enhance the synthesis of lysosomal enzymes in muscle and are necessary for the catabolic response to a variety of stimuli in this tissue and it contributes towards increase in the concentration of free amino acids in plasma (28, 29). These findings may provide one explanation for our data that concentrations of serum Cu was lower in patients with hypothyroidism, however we could not estimate plasma ceruloplasmin levels and erythrocytes Cu concentrations.

So, the observed findings hypothesize that Hypothyroidism is associated often with deficient Zinc and Copper levels in serum and the index of Oxidative stress is also significant. It can be concluded that hypothyroidism is associated with Oxidative Stress as also too decreased zinc and copper levels.

**Conflict of interest**

All authors declare no Conflict of interest.

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