

## Brief Communication

# Lower plasma levels of selenium and glutathione in smear-positive tuberculosis patients in Malawi

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## Abstract

**Objective:** The objective of this study was to investigate selenium and glutathione levels in pulmonary tuberculosis patients and controls in Malawi.

**Methods:** The case-control study included 19 patients and 15 apparently healthy controls for whom, levels of selenium and glutathione were measured.

**Results:** The plasma selenium levels were significantly lower in tuberculosis patients compared with the controls ( $p = 0.02$ ).

**Conclusion:** No significant differences were observed for the plasma glutathione levels ( $p = 0.39$ ). [*Ethiop. J. Health Dev.* 2011;25;(3):230-232]

## Background

Both protein-energy malnutrition and deficiencies in micronutrients are associated with a significant impairment of various interactions and functions of the cell-mediated immune system (1). A study in Malawi showed that selenium deficiency occurred in 87 % of tuberculosis (TB) patients (2) while a study in Ethiopia, on antioxidants status in untreated TB patients, showed that the patients were deficient in glutathione (GSH) compared to healthy controls (3). Selenium via selenoproteins and GSH are important endogenous antioxidants that protect against oxidative damage and regulate immune functions (4, 5). The objective of this study was thus to investigate plasma levels of selenium and GSH in pulmonary TB patients and controls in Malawi.

## Methods

A case-control study was conducted in Malawi between September and December 2006. Fifty-four patients were asked to participate of whom 42 accepted. Fifteen patients did not have follow-up samples, 6 withdrew the consent without any reason and data from 2 patients was not included due to inappropriate sample collection. For the controls, 47 were asked to participate and 25 consented. Six controls did not have follow-up samples and 4 withdrew the consent without any reason. The final sample set consisted of 19 patients and 15 controls aged 15-55 years.

TB was confirmed when a patient was smear-positive on at least two of three sputum specimens. The study was done 2-8 weeks after onset of treatment because then they had resumed their ordinary diet. Patients with a previous history of TB and extra pulmonary TB patients were excluded. Controls were randomly selected according to the "random-walk method" (6). Matching of

age and sex was done and adjusted for the statistical analysis.

Ethical clearance was obtained from the Regional Committee for Medical Research Ethics, Norway and the Malawi College of Medicine Research Ethics Committee, Malawi. Participants were enrolled after signing a written informed consent form.

Statistical analysis was done using SPSS for Windows 14.0 (Statistical Package for the Social Sciences, Chicago, IL, USA)

## Procedure for Blood Testing

Seven ml blood was drawn into tubes containing heparin from non-fasting participants. Two ml blood was then transferred to an eppendorf vial containing 50  $\mu$ l 2.0 M serine borate buffer for glutathione measurements. Blood samples were protected from bright light.

The samples were stored for 40-43 minutes and then centrifuged in Labofuge 200 (Osterode, Germany) at 2500 g for 10 minutes. All samples were frozen at -20 °C for 3 weeks and then at -80 °C until transferred on dry ice to Norway for analysis. The procedure for GSH analysis has been described by Bohn et al. (7). Selenium was measured with PerkinElmer, (Elan® DRC™ II Inductively Coupled Plasma Mass Spectrometry instrument, Concord, Canada).

## Results

Characteristics of study participants are shown in table 1. Adjusting for age and sex, TB patients had significantly lower levels of selenium compared to the controls ( $p = 0.02$ ). No significant difference was observed for GSH ( $p = 0.39$ ) (Table 1).

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Table 1: Characteristics of the study group, mean values of Selenium and GSH levels in Patients and controls\*\* and multivariate linear regression analysis in both groups\*

	TB patients n = 19 n (%)	Controls n = 15 n (%)	Coefficient	95% CI	P-value
Women	6	5			
Men	13	10			
Age, years, mean [SD]	28.0±7.0	32.1±10.5			
Illiterate	5 (26)	5 (33)			
Farmers	14 (74)	12 (80)			
Body Mass Index, mean [SD]	18.5±2.0	21.3±1.9			0.80
Selenium** (µmol/L)	1.10±0.22	1.23±0.19			0.10
GSH** (µmol/L)	3.85±1.68	4.48 ±1.52			0.29
Group* (selenium)			-0.17	-0.31-0.02	0.02
Group* (GSH)			-0.53	-1.81-0.73	0.39

\*\*Mean±Standard deviation [SD] GSH=Glutathione

\*Adjusted for age and sex. CI= Confidence Interval

### Discussion

After adjusting for age and sex, selenium concentration in TB patients was 0.22 µmol/L lower than the controls ( $p = 0.02$ ). The difference is significant and in agreement with Kassu et al. (8). The concentration of GSH in TB patients was 0.53 µmol/L lower than the controls ( $p = 0.39$ ). This result is not in agreement with results found in Ethiopia (3) where the GSH concentrations were significantly lower in TB patients. The possible explanations could be the degradation of GSH by enzyme gamma-glutamyl transferase during sample collection and preparation (9), small sample size and selection bias. We have avoided degradation of GSH by using serine borate buffer that inhibits enzyme gamma-glutamyl transferase (9).

Selenium is a trace mineral, involved in several key metabolic activities via selenoproteins. These enzymes are essential to protect against oxidative damage and to regulate immune functions (4). GSH is an antioxidant and protects cells from toxic effects of reactive oxygen species. It plays an important role in regulating the cellular immunity and protects against mycobacteria (5). The reason for lower selenium and GSH levels in TB patients is unclear, but could be explained by the increased oxidative stress and the reduced levels of antioxidant capacity found in this group (3). Other factors like variation in diet, absorption, the severity of the disease and being HIV positive may also have influenced the plasma levels of selenium and GSH.

Limitations of the study were the small sample size and no statistical correction for HIV status in both groups. Further studies are hence needed to understand the observed deficiencies of selenium and GSH in TB patients.

### Conclusion

This study showed that TB patients had lower plasma levels of selenium compared with controls.

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### References

1. Chandra RK. Nutrition and the immune system: An introduction. *Am J Clin Nutr* 1999;66(2):460S-3S.
2. Van Lettow M, Harries AD, Kumweda JJ, Zijlstra EE, Clark TD, Taha TE, et al. Micronutrient malnutrition and wasting in adults with pulmonary tuberculosis with and without HIV co-infection in Malawi. *BMC Infect Dis* 2004;21:4(1):61
3. Madebo T, Lindtjorn B, Aukrust P, Berge RK. Circulating antioxidants and lipid peroxidation products in untreated tuberculosis patients in Ethiopia. *Am J Clin Nutr* 2003;78(1):117-22.
4. Ryan-Harshman M, Aldoori W. The relevance of selenium to immunity, cancer, and infectious/inflammatory diseases. *Can J Diet Pract Res* 2005;66(2):98-102.
5. Venketaraman V, Dayaram YK, Amin AG, Ngo R, Green RM, Talaue MT, et al. Role of glutathione in macrophage control of mycobacteria. *Infect Immun* 2003;71(4):1864-71.
6. Gibson RSFEL. An Interactive 24-hour recall for assessing the adequacy of iron and zinc intakes in developing countries. 1999. ILSI Press. Ref Type: Pamphlet

7. Bohn SK, Smeland S, Sakhi AK, Thoresen M, Russnes KM, Tausjo J, et al. Post-radiotherapy plasma total glutathione is associated to outcome in Patients with head and neck squamous cell carcinoma. *Cancer Let* 2006;238(2):240-7.
8. Kassu A, Yabutani T, Mahmud ZH, Mohammad A, Nguyen N, Huong BT, et al. Alterations in serum levels of trace elements in tuberculosis and HIV Infections. *Eur J Clin Nutr* 2006;60(5):580-6.
9. Sakhi AK Russnes KM, Smeland S, Blomhoff R, Gundersen TE. Simultaneous quantification of reduced and oxidized glutathione in plasma using a two-dimensiounal chromatographic system with parallel porous graphitized carbon columns coupled with fluorescence and coulometric electrochemical detection. *J Chromatogr A* 2006;1104(1-2):179-89.