Vitamin A and zinc deficiencies among tuberculosis patients in Ethiopia

Tibebeslassie Seyoum Keflie, Aregash Samuel, Ashagrie Zewdu Woldegiorgis, Adane Mihret, Markos Abebe, Hans Konrad Biesalski

ABSTRACT

Background: The link between tuberculosis (TB) and malnutrition has long been recognized. Vitamin A and zinc deficiencies may reduce the host defenses and increase the risk for diseases.

Objective: The aim of the present study was to estimate the difference in vitamin A and zinc deficiencies together with dietary intakes among pulmonary TB patients and controls.

Materials and methods: A case-control study design was employed to undertake this study in North Shewa, Ethiopia. Sputum smear examination, high-performance liquid chromatography (HPLC), flame atomic absorption spectrometry (FAAS), and enzyme-linked immunosorbent assay (ELISA) were used to analyse acid fast bacilli (AFB), vitamin A, zinc, and C-reactive protein (CRP), respectively. Dietary intake was assessed using a 24-h recall questionnaire. Mann–Whitney U test, Kruskal–Wallis test, Chi-square, odds ratio (OR), Spearman correlation, and multinomial logistic regression model were computed for data analyses.

Results: In this study, 62 TB cases and 59 controls were included. The proportions of vitamin A deficiencies among TB cases and controls were 56.4% and 39.0%, respectively. All TB cases and 92.5% controls were zinc deficiencies together with protein-energy malnutrition. The odds of TB cases with deficiencies of vitamin A and zinc was 2.3 (95% CI: 1.1 to 4.8) times more likely as compared to the controls. More than 80% of all participants had below average fulfilment of energy and vitamin A intakes.

Conclusion: Vitamin A and zinc deficiencies are severe problems among TB patients. Moreover, undernutrition determines the development of TB. Therefore, the management programs of TB need to address the problems of vitamin A and zinc deficiencies together with protein-energy malnutrition.

Introduction

Tuberculosis (TB) is an ancient disease caused by Mycobacterium tuberculosis. Despite the applied efforts for its control, TB is still one of the major public health challenges. According to the global TB report in 2017, 10.4 million TB cases and 1.67 million TB deaths occurred worldwide in 2016. Of whom, about 70% of the new cases and more than 85% of the deaths were reported from South East Asia and Africa [1]. The report also showed that Ethiopia is one of the top 30 TB burden countries globally with 182,000 new cases and 30,000 deaths in 2016 [1].

The link between TB and malnutrition has long been recognized; malnutrition may predispose people to the development of clinical disease, and TB can contribute to malnutrition [2]. Malnourished people are also susceptible to a new infection since their immune systems are debilitated [3]. As key contributors to immune function and cytokine kinetics, micronutrients such as vitamin A and zinc play a major role in combating TB [4].

Vitamin A is a fat-soluble vitamin which is needed in small quantities for several metabolic activities in the body [5]. Vitamin A and its active metabolites are important for growth and differentiation of a variety of cells, mainly in mucosa-associated epithelia [6]. T and B lymphocytes, macrophages, and generation of antibodies [7]. Zinc is also an essential trace element with diverse physiologic and metabolic implications.
functions [8] such as maintaining immunological integrity, cellular immunity, and antioxidant activity [9]. Because of the absence of specialized zinc storage in the body, a daily intake is required to achieve its steady-state [10]. In the prechemotherapeutic era, cod liver oil rich in vitamin A was used regularly for the treatment of TB to strengthen the host defense system [11].

Vitamin A and zinc deficiencies are common features of pulmonary TB. Deficiencies of both micronutrients can reduce host defenses and immune responses [12]. A study done in Rwanda revealed that 29% of adult TB patients had serum vitamin A levels consistent with deficiency (less than 0.7 μmol/L) [13]. Deficiency in zinc (below 10.7 μmol/L) is thought to be one of the primary causes of morbidity in developing countries and yet little is known about the status of the world [14].

Until recent time, many of the epidemiological studies conducted on vitamin A and zinc deficiencies in Ethiopia were focusing on children and pregnant women [5,15,16]. However, there is a paucity of information on the magnitude of the deficiencies of these micronutrients in TB patients. The available studies were very few [17,18] and conducted long years ago in some pocket areas of North West part of Ethiopia. Therefore, the present study was designed to estimate the difference in vitamin A and zinc deficiencies together with dietary intakes among pulmonary TB patients and non-TB controls.

Materials and methods

Study design, area and population

A facility-based case-control study was conducted in North Shewa Zone of Amhara Regional State, Central Ethiopia. In this study, one referral hospital and five health centers were involved. The study population included TB patients and non-TB controls who were living in the same geographic area. All TB cases who visited the health facilities between March and August 2015 were recruited.

Selection of TB cases and controls

The diagnosis of TB was performed microbiologically and radiologically as per the national standard diagnostic algorithm of Ethiopia [19]. As per the diagnostic algorithm, smear positive pulmonary TB is defined by at least two initial sputum smear examinations positive for acid fast bacilli (AFB) by direct microscopy, or a patient with one initial smear examination positive for AFB by direct microscopy and culture positive, or a patient with one initial smear examination positive for AFB by direct microscopy and radiographic abnormalities consistent with active TB as determined by a clinician.

Smear-negative pulmonary TB is also defined by a patient having symptoms suggestive of TB with at least three initial smear examinations negative for AFB by direct microscopy, and no response to a course of broad-spectrum antibiotics; radiological abnormalities consistent with pulmonary TB; decision by a clinician to treat with a full course of anti-TB; or a patient whose diagnosis is based on culture positive for M. tuberculosis [19,20].

In this study, both smear positive and negative pulmonary TB cases who were willing to participate in the study were included. All selected TB cases were used as a point of contact to recruit controls. Individuals who were apparently healthy, non-TB, living in the same geographic areas with TB cases and interested in participating in the study were selected as controls. As vitamin A and zinc concentrations in the serum are affected by many physiological and pathological states and drugs, TB cases and controls with pregnancy, lactation, chronic or degenerative diseases, and who were taking vitamin A and zinc supplements, corticosteroid drugs and oral contraceptives were excluded from the study.

Duration of treatment

The treatment of TB lasts for a total of six months for newly diagnosed TB patients or for those who had taken previously the anti-TB drugs for less than one month as per the standard treatment guidelines for the general hospital in Ethiopia. The treatment regimen consists of 2 months treatment with Rifampicin (R), Isoniazid (H), Pyrazinamide (Z) and Ethambutol (E) during the intensive phase, followed by four months with Rifampicin and Isoniazid in the continuation phase (2RHZE/4RH) [21].

Data collection

A structured questionnaire, comprised of socio-demographic characteristics, health status and diet factors, was prepared, translated into Amharic (the local language) and administered to each of the study participant. Medical records were also used to collect clinical characteristics.

Anthropometric measurements

Standardized procedures were employed to measure body weight, height, and mid upper arm circumference (MUAC). All study participants were weighed while wearing light clothes using an electronic platform weighing scale to the nearest 0.1 kg. Height was measured to the nearest 0.1 cm by means of a seca stadiometer. Body mass index (BMI) was calculated as body weight (kg) divided by height (m) squared (kg/m²). BMI values of 18.5, 17.0, and 16.0 kg/m² were used as the cut-off values below which patients were classified as having mild, moderate, or severe malnutrition [21]. MUAC was measured half way between the olecranon and acromion processes of the left arm using a flexible non-stretch measuring tape to the nearest 0.1 cm while the arm is hanging relaxed, without compressing the tissues. MUAC less than 23 cm for male and 22 cm for female is used to define undernutrition as per FANTA III [22].

Dietary intake assessment

The dietary intake was assessed by means of a 24-h recall to estimate the intake of energy, protein, vitamin A and zinc. Each 24-h recall was performed using a standardized four-stage protocol [23,24]. First, a complete list of all food and beverages consumed over the 24 h was obtained. Second, detailed descriptions of the food and beverages consumed, including the cooking methods and brand names were recorded, together with the time and place of consumption. Third, estimates of the portion sizes of all consumed food items were determined by referring to household measuring and serving utensils (e.g., spoons, plates or cups), and food packages. Finally, the food recall was reviewed to ensure that all items had been recorded correctly. The amount of nutrient intake from dietary recalls was calculated using nutrisurvey software (Dr.Juergen Erhardt SEAMEO-TROPMED RCCN-University of Indonesia copy right © 2007).

After the dietary intake assessment, the individual dietary diversity score (DDS) was calculated as the number of food groups consumed over the 24-h recall. These food groups were based on the guidelines for measuring the household and individual dietary diversity. The score for individual diet diversity goes from 0 to 9 [25,26]. Considering four food groups as the minimum acceptable dietary diversity, the individual DDS less than four was classified as having a poor dietary diversity [26,27].

Blood collection and serum separation

After overnight fasting, 10 mL of venous blood was withdrawn from antecubital fossa vein into a non-heparinized vacutainer tube between 8:00 and 10:00am at the selected health facilities. Blood samples were allowed to clot for about 1 h in the dark and subjected to centrifugation.
at 4000 rpm for 10 min at room temperature. Samples with visible haemolysis were discarded.

The sera were separated immediately into aliquots of sterile Eppendorf tubes by means of sterile Pasteur pipettes and stored at −20 °C in the laboratory of the health facility. The sera were later transported in the ice box to Armauer Hansen Research Institute (AHRI) where they were stored at −80 °C until analysis. Aliquots of the sera were transferred to Ethiopian Public Health Institute (EPHI) for the analysis of serum vitamin A concentration and to Natural Sciences College of Addis Ababa University for the determination of the serum zinc concentration.

Measurements of vitamin A, zinc and C-reactive protein (CRP)

Vitamin A

Serum vitamin A concentration was determined using high performance liquid chromatography (HPLC) as explained in the method of Arroyave et al. [28] with slight modifications. In brief, 100 µL of ethanol and an equal volume of retinyl acetate (reconstituted in ethanol) as internal standard were added into a 2 mL Eppendorf tube containing 100 µL of serum sample. The solution was then mixed for 1 min using vortex mixer. In the first round, 750 µL of n-hexane was added into the solution, vortex mixed for 1 min and centrifuged at 3000 rpm for 10 min. The supernatant was transferred into a new Eppendorf tube. In second round, 750 µL of n-hexane was added and the whole steps were repeated. The pooled supernatant was then evaporated into dryness under a steam of nitrogen, reconstituted in 200 µL of methanol, and transferred into the injection vial.

A Shimadzu HPLC system (Shimadzu, Tokyo) which composed of a reverse phase Supelco LC-18 column (250 × 4.6 mm, 5 µm particle size) and UV–vis detector was used to separate and detect the retinol at 325 nm with a column temperature set at 40 °C. Series of standards (having a concentration of 60, 40, 20, 10 and 5 µg/dL) and samples were loaded into the autosampler tray of the HPLC. Methanol was used as a mobile phase with injection volume of 40 µL, flow rate of 1 mL/min and retention time of 15 min.

Retinol was determined by comparing the retention times with the external standard and quantified based on externally drawn calibration curve using the series concentrations of standards. All extraction procedures were performed under dim light to avoid oxidation of the compound. Vitamin A concentration below 0.7 µmol/L was considered as deficient.

Zinc

Serum zinc concentration was determined by means of flame atomic absorption spectrometry (FAAS) with a micro-sampling technique in AA-7000 Atomic Absorption Spectrophotometer (Shimadzu, Japan) using the method described in Sepehri et al. [29]. In brief, 10 mL of concentrated nitric acid was added to 1 mL of serum in a beaker and heated for 3 h, below boiling point, on a hot plate. When the volumes of the samples reduced to about one-third, 5 mL of 30% hydrogen peroxide solution was added. The samples were further heated almost to dryness at the same temperature. Finally, the residues were dissolved in 50 mL of 1% nitric acid and filtered. The prepared samples were transferred into 50 mL polyethylene tubes for zinc analysis. The concentration of 10.7 µmol/L was used as a cut-off value below which was considered as zinc deficient.

C-reactive protein (CRP)

CRP was measured at AHRI, Ethiopia using enzyme-linked immunosorbent assay (ELISA) (Human CRP ELISA Kit, HK 358, Hycult biotech) method according to the manufacturer’s instruction as indicated in the manual of the kit. For each TB patient, CRP was measured in duplicate. The mean value ≥10 mg/L was considered as CRP positive.

Ethical consideration

This study is a part of our project entitled ‘‘effect of micronutrients on the treatment outcome of TB’’, which has been ethically approved by the ethical approval committee of AHRI- ALERT (All Africa Leprosy Rehabilitation and Training) Centre. Supportive letters were also obtained from the zonal and districts’ health bureau of Amhara Regional State. This study was carried out in accordance with the principle of Helsinki declaration. After explaining the purpose and objective of the study, written informed consent was obtained from each of the study participant.

Statistical analysis

Statistical analysis was conducted using IBM SPSS version 23 statistical program. All continuous data were checked for a normal distribution using a Shapiro-Wilk test. Descriptive statistics, including frequency, proportion, and median (IQR-Inter Quartile Range) were used to describe concentrations of vitamin A and zinc, BMI, and MUAC. Odds ratio (OR) together with 95% confidence interval (CI) was computed to assess the strength of association.

The significance of the difference in continuous and categorical variables between groups was compared using Mann–Whitney U test and Chi-square test, respectively. Kruskal–Wallis test was used to assess the difference in the serum vitamin A and zinc concentrations of TB patients at different duration of treatment. The association between MUAC, BMI, and DDS was examined using Spearman correlation test.

Bivariate analysis was done to explore the crude association between different predictor variables and TB. To control for possible confounding factors and identify factors that are independently associated with TB, multinomial logistic regression analysis was performed for those variables with p-value of less than 0.2 in the bivariate analysis. The major assumptions of logistic regression model were multicollinearity and interaction among independent variables. The absence of multicollinearity was checked using variance inflation factor (VIF)/tolerance and the fitness of logistic regression model was checked using Hosmer-Lemeshow statistical test. Unless specified, p-value < 0.05 was considered as statistically significant.

Results

Socio-demographic characteristics

The present study was conducted on 62 TB patients and 59 controls. The median (IQR) age was 26 (14) years with the minimum and maximum age of 14 and 77 years. Most of the participants were Orthodox Tewahido Christians in religion (89.3%) and farmers in occupation (31.1%) (Table 1).

There was statistically significant difference in sex between TB patients and the controls (P < 0.05). The odds of males with TB was 2.2 (95% CI: 1.05 to 4.48) times more likely as compared to the odds of females. The most (85%) TB afflicted age group was found between 19 and 50 years.

Clinical signs and symptoms of TB

TB patients showed various clinical signs and symptoms. The identified clinical signs and symptoms at the time of diagnosis were fatigue (100%), cough (83.3%), malaise (83.3%), night sweating (83.3%), fever (75%), anorexia (75%), productive cough (75%), chest pain (70.8%), dyspnoea (54.2%), haemoptysis (29.2%) and tachycardia (25%).
Vitamin A and zinc deficiencies

The proportions of vitamin A, zinc and the combination of the two micronutrient deficiencies were 47.9%, 96.5% and 47.8%, respectively. The mean serum retinol and zinc concentrations in females (0.80 ± 0.47 µmol/L and 2.10 ± 1.20 µmol/L, respectively) were not significantly different from that of males (0.77 ± 0.60 µmol/L and 2.50 ± 2.50 µmol/L, respectively) (p > 0.05). The proportions of vitamin A deficiencies among TB patients and the controls were 56.4% and 39.0%, respectively.

All TB patients and 92.5% of the controls were identified as zinc deficient. There were statistically significant differences in all proportions of the deficiencies between TB patients and the controls at p < 0.1. The odds of TB patients who were vitamin A deficient was 2.03 (90% CI: 1.1 to 3.7) times greater than the odds of the controls. In the same line, TB patients had vitamin A and zinc deficiencies 2.3 (95% CI: 1.1 to 4.8) times more likely as compared to the controls.

CRP

About 62% of TB patients were CRP positive. The mean ± SD of retinol and zinc concentration in CRP positive TB patients were 0.62 ± 0.50 µmol/L and 2.05 ± 0.74 µmol/L, respectively. Whereas, these values in CRP negative TB patients were 0.75 ± 0.69 µmol/L and 2.48 ± 1.20 µmol/L, respectively. Excluding CRP positives, the prevalence of vitamin A deficiencies in CRP negative TB patients became 66.7%. Both CRP negative and positive TB patients did not have statistically significant differences in serum retinol (p = 0.25) and zinc (p = 0.09) concentrations.

Durations of TB treatment

Fig. 2 shows the mean serum retinol and zinc concentrations in different durations of TB treatment. During the first month of TB treatment, the retinol level was below the cut-off value of 0.70 µmol/L but increased slightly afterward. Similarly, the zinc level increased slightly up to 2 months of treatment but its concentration throughout the entire courses of the treatment was below the cut-off value of 10.70 µmol/L. Per the result of the analysis of variance, the retinol and zinc concentrations were not significantly different in both within and between the durations of the treatment.

Dietary intakes

The mean DDS was 3.56. Most (84%) of TB patients and the controls (86%) had below average DDS (4.5 out of 9 food groups). DDS had statistically significant correlation with BMI (γ = −0.22; p = 0.019). There was also a significant difference in DDS between males and females (p = 0.031). While analysing the nutrients intake, about 87% of TB patients and 80% of the controls had below 50% fullment of energy intake. Almost all participants had below 50% fullment of vitamin A intake. But, less than 30% of participants had protein and zinc intakes below 50% fullments. In other words, many participants had above the average fullments of protein and zinc intakes, albeit much cereal-based diet.

Nutritional status

More than 29% of the participants had BMI below 18.50 kg/m². The mean BMI of female TB patients (18.80 ± 3.80 kg/m²) was

Table 1

| Type          | Variables | TB-cases (n) | Control (n) | P-value |
|---------------|-----------|--------------|-------------|---------|
| Sex           | Female    | 26           | 36          | 0.036   |
|               | Male      | 36           | 23          |         |
| Residence     | Urban     | 24           | 22          | 0.391   |
|               | Rural     | 29           | 37          |         |
| Age (years)   | 14–18     | 9            | 9           | 0.805   |
|               | 19–30     | 37           | 32          |         |
|               | 31–50     | 12           | 15          |         |
|               | 51–70     | 3            | 3           |         |
| Marital status| Single    | 1            | 0           | 0.807   |
|               | Married   | 30           | 32          |         |
|               | Widowed   | 1            | 1           |         |
| Educational level| Illiterate | 8          | 8           | 0.644   |
|               | Primary   | 20           | 16          |         |
|               | Secondary | 23           | 21          |         |
|               | Tertiary  | 11           | 12          |         |
|               | Religious | 0            | 2           |         |
|               | Teaching  | 56           | 52          | 0.176   |
|               | Christian | 1            | 5           |         |
|               | Muslim    | 1            | 5           |         |
|               | Protestant| 3            | 2           |         |
|               | Catholic  | 2            | 0           |         |
| Occupation    | Student   | 9            | 15          | 0.002   |
|               | Private   | 12           | 19          |         |
|               | Government employees | 8 | 7 |         |
|               | Non-Government employees | 0 | 1 |         |
|               | Farmer    | 29           | 8           |         |
|               | Housewife | 2            | 8           |         |
|               | Retired   | 1            | 0           |         |

Fig. 1 shows a wide-ranging of serum retinol levels. The proportions of severe and, mild to moderate hyporetinolaemia were higher in TB patients than the controls. About 19% of TB patients and 25% of the controls had suboptimal retinol levels, whereas the optimal level in both groups was below 5%.

Dietary intake

The mean DDS was 3.56. Most (84%) of TB patients and the controls (86%) had below average DDS (4.5 out of 9 food groups). DDS had statistically significant correlation with BMI (γ = −0.22; p = 0.019). There was also a significant difference in DDS between males and females (p = 0.031). While analysing the nutrients intake, about 87% of TB patients and 80% of the controls had below 50% fulment of energy intake. Almost all participants had below 50% fulment of vitamin A intake. But, less than 30% of participants had protein and zinc intakes below 50% fulments. In other words, many participants had above the average fulments of protein and zinc intakes, albeit much cereal-based diet.

Nutritional status

More than 29% of the participants had BMI below 18.50 kg/m². The mean BMI of female TB patients (18.80 ± 3.80 kg/m²) was
significantly lower than females in the controls (20.40 ± 2.92 kg/m²) (p < 0.05). There was a statistically significant difference in the proportions of undernutrition (BMI < 18.50 kg/m²) between TB patients (40.3%) and the controls (19.3%) (P < 0.05). Undernourished individuals had 2.8 times the odds of TB disease compared to the controls. As defined by MUAC, undernutrition for both male and female was 44.2%. The mean MUAC of females in TB patients (20.60 ± 2.20 cm) was significantly different from those females in the controls (21.80 ± 2.20 cm) at p < 0.10. MUAC was significantly correlated with BMI (\(\gamma = 0.56, P < 0.001\)).

**Determinants of TB**

The multinomial logistic regression model in Table 2 indicated that sex and BMI had statistically significant association with TB patients (p < 0.05). Adjusting for BMI, serum retinol level, religion, and occupation, females had 0.2 times less the odds of TB compared to males. Similarly, undernourished individuals had 3.3 times the odds of TB compared to well-nourished individuals holding sex, serum retinol level, religion and occupation constant.

**Discussion**

In most countries, TB notification is twice as high in men as in women [33]. The study done in central part of Ethiopia also showed a high rate of TB notification in male (60.9%) and people with the age group of 14 to 54 years (87.3%) [20]. In line with this, the odds of males with TB in the present study was 2.2 times more likely as compared to the odds of females. The most (85%) TB afflicted age group was also found between 19 and 50 years. The reason why males and the productive age groups are highly afflicted by TB could be associated with their sociocultural, behavioral and biological components [33].

Micronutrient deficiencies are rampant in Ethiopia. More than 47% of TB patients and controls were affected by the deficiencies of either vitamin A, zinc or their combination. This was corroborated by the report of Keffie et al. [26] who identified the presence of a high risk of micronutrient deficiency in central part of Ethiopia (>60%). According to WHO and International Zinc Nutrition Consultative Group (IZiNCG), the risks of vitamin A and zinc deficiencies are of public health concern when the prevalence of low serum retinol and zinc concentrations are greater than 20% [34]. This implied that the results of the present study were more than twice of the indicator of the public health concern.

Comparing with the control, the odds of TB patients who were vitamin A deficient was 2 times higher. The proportion of vitamin A deficiency in TB patients was 56.4% which was closer to the report of nearly 60% among TB patients in Gondar, Northwest Ethiopia [18]. Severe and mild to moderate types of vitamin A deficiency was more common in TB patients than the controls. The low concentration of vitamin A in the serum of TB patients could be resulted from TB induced anorexia that lead to low consumption of vitamin A rich food items; reduced absorption of dietary vitamin A owing to parasites co-infection; decreased mobilization of hepatic reserves of retinol; rapid utilization of vitamin A by target tissues; and increased urinary losses of vitamin A associated with fever [18,35]. This justification was strengthened by the occurrence of fever and anorexia together with

| Variables                      | β    | Wald | DF  | P-value | Odds Ratio | 95% CI Lower Limit | 95% CI Upper Limit |
|-------------------------------|------|------|-----|---------|------------|--------------------|--------------------|
| Sex                           |      |      |     |         |            |                    |                    |
| Females                       | −1.64| 10   | 1   | 0.002*  | 0.19       | 0.07               | 0.53               |
| Males                         |      |      |     |         |            |                    |                    |
| BMI                           |      |      |     |         |            |                    |                    |
| < 18.50 kg/m²                 | 1.2  | 5.06 | 1   | 0.024*  | 3.33       | 1.17               | 9.50               |
| ≥ 18.50 kg/m²                 |      |      |     |         |            |                    |                    |
| Retinol                                                                     |      |      |     |         |            |                    |                    |
| < 0.70 µmol/L                 | 0.62 | 1.77 | 1   | 0.184   | 1.86       | 0.74               | 4.64               |
| ≥ 0.70 µmol/L                 |      |      |     |         |            |                    |                    |
| Religion                      |      |      |     |         |            |                    |                    |
| Orthodox Tewahido             | 0.7  | 0.22 | 1   | 0.64    | 2.01       | 0.10               | 38.56              |
| Muslim                        | −0.48| 0.07 | 1   | 0.796   | 0.62       | 0.02               | 23.34              |
| Protestant                    |      |      |     |         |            |                    |                    |
| Occupations                   |      |      |     |         |            |                    |                    |
| Students                      | −0.92| 2.01 | 1   | 0.16    | 0.40       | 0.11               | 1.42               |
| Private workers               | −0.84| 1.77 | 1   | 0.183   | 0.43       | 0.13               | 1.48               |
| Government employees          | −0.97| 1.59 | 1   | 0.207   | 0.38       | 0.09               | 1.71               |
| Housewives                    | 1.73 | 2.91 | 1   | 0.088   | 1.26       | 0.35               | 4.51               |
| Farmers                       |      |      |     |         |            |                    |                    |

NB: The reference category is the controls.

*Statistically significant at p < 0.05.
fatigue, cough, malaise, and night sweating in more than 74% of TB patients in the present study.

Likewise, there was a significant difference in the proportion of zinc deficiency among TB patients and controls. All TB patients were identified as zinc deficient. This result was much higher than the report of the previous study done in Northwest Ethiopia (32.1%) [17]. Such a proportional difference could be owing to variations in the dietary patterns and seasons together with the availability and accessibility of food items rich in zinc. In line with the present study, the low serum zinc concentration in TB patients was also reported in Indonesia [24], Ecuador [36], Malaysia [37], and Korea [38].

The lowered concentration of zinc among TB patients could probably be associated with reduction in hepatic production of a zinc carrier protein (α2-macroglobulin); redistribution of zinc from blood circulation to other tissues; and/or a rise in the production of metallothionein, a protein that transports zinc to the liver [17,37,39]. The concomitant deficiencies of vitamin A and zinc were very high in TB patients as compared to the control. This could be due to the high magnitude of zinc deficiency in TB patients. Vitamin A and zinc deficiencies are usually occurred together, as zinc deficiency impairs the synthesis of retinal binding proteins and reduces plasma retinal concentration [40].

Several studies indicated the essence of measuring CRP in TB patients to get a clear picture on the serum retinol and zinc concentrations [18,36,41]. CRP is a tissue protein which is produced at the time of acute phase response and its concentration changes rapidly as a result of infection or inflammation. Visser et al. [41] showed the relationship between the low retinol concentration in active TB and the acute phase response. Likewise, Cuevas and Koyanagi [9] described the low concentration of zinc in TB patients with and without a raised CRP.

In the present study, we did not observe any difference in the concentrations of vitamin A and zinc in the serum of TB patients with and without CRP. In line with our finding, Karyadi, et al. [24] reported that retinol concentration did not correlate significantly with markers of acute phase response. This suggested that CRP is not the best option to control the change in the acute phase response in the concentrations of vitamin A and zinc in the serum of TB patients.

We observed a fluctuation of vitamin A and zinc concentrations in TB patients at different durations of anti-TB treatment. During the first month, the retinol concentration was below the cut-off value but later increased slightly. The gradual increment of zinc concentration was also observed up to the second month of the treatment, but such a concentration was below the cut-off value throughout the whole course of anti-TB treatment. The improvement of vitamin A was in accordance with the reports of Mugusi et al. [7] in Tanzania and Kasu et al. [18] in Ethiopia. It is believed generally as patients improve and fever subsides, the loss of vitamin A declines, and its serum concentration resume to the reference ranges [18].

Despite the low improvement, our observation on the changes of zinc concentration was corroborated by the findings of Khanna et al. [42] and Kasu et al. [17]. On the contrary, Edem and others [39] revealed that trace elements and vitamins were lowest at 2 and 4 months of anti-TB drug therapy. This may be due to drug-micronutrient interactions or drug induced nutrient depletion. Further, the effects of anti-TB drugs on the absorption of zinc was presented by Karyadi et al. [11] in their study done in Indonesia. Ethambutol was shown in rats to increase not only zinc absorption but also urinary zinc losses, resulting in reduced circulating zinc concentrations [43].

In the comparison made on nutritional status as measured by BMI and MUAC, the results were significantly lower in TB patients than the controls. Undernourishment in TB patients was almost three times higher than that of the controls. In agreement with our results, several studies demonstrated undernutrition as a well-recognized clinical sign of active TB [24,44,45]. Pakasi et al. [46] described undernutrition as a reflection of two processes. One would be protein-energy malnutrition (which severely affects host defense) and the other was wasting due to the catabolism induced by the acute phase response. The causes of undernutrition in patients with active TB could be anorexia, impaired absorption of nutrients, or increased catabolism associated with the inflammatory and immune response [24,44]. Our study identified undernutrition and sex as significant determinants of active TB.

The dietary intake assessment revealed that the mean DDS was 3.56, and above 80% of TB patients and the controls had below the average DDS. These results were in accordance with the previous study done in the same area [26]. A diet of at least 4 DDS was valid as nutritionally adequate, but below 4 DDS represented a poor dietary diversity [26,47]. This suggested that both TB patients and controls had poor dietary diversity. The intakes of energy and vitamin A rich foods items were also poor. More than 85% of TB patients could not fulfill even half of their daily requirements of energy and vitamin A. Poor dietary intake of vitamin A rich food is an important predictor of vitamin A deficiency [16].

Although the intakes of protein and zinc looked better, the main dietary sources were cereal based. The previous study in the same area also indicated a high reliance on the consumption of cereals, vegetables and legumes with less animal source foods [26]. As our body does not store zinc, it requires a constant dietary intake. Most plant-based diets are not good sources of zinc owing to the presence of phytate, a component of plants that chelates zinc and prevents its absorption [48]. The low content of zinc in Ethiopian soil could be another factor for the inadequate intake of zinc [15]. Hence, plant- based diets with low animal source proteins were accounted for the high deficiency of zinc in both TB patients and controls.

**Limitation**

Although this study had the strength of dealing with the serum vitamin A and zinc concentrations in TB patients together with CRP, nutritional status and dietary intakes, it was a facility-based study and the participants could not represent the general population. In addition, the 24-h recall method, which was used to assess the dietary intake, could probably miss some of the feeding behaviours particularly in the case of altered dietary patterns.

**Conclusion**

In conclusion, vitamin A and zinc deficiencies are severe problems among TB patients and non-TB controls. The cause-effect relationship between undernutrition and TB is just like the puzzle of the hen or the egg, of the two which comes first. Our study underlines that undernutrition determines the development of TB. Most TB patients have DDS below average and, very poor dietary intakes of vitamin A, zinc, protein, and energy. During the duration of anti-TB treatment, the concentrations of vitamin A and zinc improve but not to the extent of curbing their deficiencies. Therefore, to claim success, TB management program needs to give emphasis on addressing the problems of vitamin A and zinc deficiencies together with protein-energy malnutrition. In the future, we recommend further community-based studies to be conducted at different parts of Ethiopia to substantiate our findings.

**Acknowledgment**

We are grateful to all study participants, data collectors, health facilities, and the health bureau of North Shewa Zone of Amhara Region. We are also indebted to Mr. Alexander Koza, Mr. Hulumtaye Tefera, Mr. Meseret Woldeyohannes and Mr.Debebe Haileu for their technical assistance during the processes of the laboratory analyses. This study was financially supported by the Dr. Hermann Eiselen Ph.D. grant from the foundation fiat panis. The first author obtained a scholarship from Food Security Centre (FSC) of University of Hohenheim, which is supported by the German Academic Exchange Service (DAAD) with funds of the Federal Ministry of Economic Cooperation and Development (BMZ) of Germany, and thus, highly acknowledged.

32
Conflict of interest

The authors declared that they have no competing interest.

Author’s contribution

TSK designed the study, collected the data, analyzed, interpreted and drafted the manuscript. AS, AZ, AM, MA and HKB critically reviewed the manuscript. All the authors read and approved the manuscript.

Supplementary materials

Supplementary materials associated with this article can be found, in the online version, at doi:10.1016/j.jctube.2018.05.002.

References

[1] WHO. Global TB report 2017. Annex 4: TB burden estimates, notifications and treatment outcomes for individual countries and territories, WHO regions and the world. 2017. Accessed on February 26, 2018. http://www.who.int/tb/publications/global_report/gtbr2017_annex4.pdf?ua=1.

[2] Macallan DC. Malnutrition in tuberculosis. Diagn Microbiol Infect Dis 1999;34(2):153–7.

[3] Sinclair D, Abba K, Grobler L, Sudaasnanam TD. Nutritional supplements for people being treated for active tuberculosis. Cochrane Database Syst Rev 2011:11.

[4] Bhakaram P. Micronutrient malnutrition, infection, and immunity: An overview. Nutr Rev 2002;60:540–5.

[5] Demissie T, Ali A, Mekonnen Y, Haider J, Umetu M. Magnitude and distribution of vitamin A deficiency in Ethiopia. Food Nutr Bull 2010;31(2).

[6] Biesalski HK, Donatou N. New aspects in vitamin A metabolism: the role of retinyl esters as systemic and local sources for retinol in mucous epithelia. J Nutr 2000;130(12 suppl 1):3458–50.

[7] Mugusi FM, Rusizoka O, Habib N, Fawzi W. Vitamin A status of patients presenting with pulmonary tuberculosis and asymptomatic HIV-infected individuals, Dar es Salaam, Tanzania. Int J Tuberc Lung Dis 2003;7(8):804–7.

[8] Hambidge M. Zinc and health: current status and future directions. J Nutr 2000;130(Suppl 13):1344–9.

[9] Cuevas L, Koyanagi A. Zinc and infection: a review. Ann Trop Paediatr 2005;25:49–60.

[10] Bonaventura P, Benedetti G, Albaride F, Misser P. Zinc and its role in immunity and inflammation. Autoimmun Rev 2015;14:277–85.

[11] Karyadi E, West CE, Schultink W, Nelwan RH, Gross R, Amin Z, Dolmans WMV, van der Meer JW, Hautvast JG, West CE. Poor micronutrient status of active pulmonary tuberculosis patients in Indonesia. J Nutr 2000;130:2953–8.

[12] FAO. Food and Agriculture Organization. Guidelines for measuring household food insecurity and inadequate dietary diversity. 2007. Version 2. Rome, Italy.

[13] Kellie TS, Samuel A, Christine L, Noor D, Biesalski HK. Dietary Patterns and Risk of Micronutrient Deficiencies: their Implication for Nutritional Intervention in Ethiopia. J Nutr Health Food Sci 2017;6(1):1–16. http://dx.doi.org/10.15226/jnhfs.2018.001120.

[14] WHO. World Health Organization. Indicators for assessing infant and young child feeding practices Part 1 Definitions. Food and nutrition technical assistance; 2007.

[15] Arroyave G, Chichector CO, Flores H, Glover J, Mejia L, Olsen JA, Simpson KL, Underwood BA. Biochemical methodology for the assessment of vitamin A status: a report of the international vitamin a consultant group. Washington DC: Nutrition Foundation; 1982.

[16] Sepehri Z, Mirzaei N, Sargazi A, Sargazi A, Mishkar AP, Kiani Z, Oskooe HO, Arefi D, Ghami S. Essential and toxic metals in serum of individuals with active pulmonary tuberculosis in an endemic region. J Clin Tuberculosis Other Mycobact Dis 2017;6:8–13. http://dx.doi.org/10.1016/j.jctube.2017.01.001.

[17] WHO. World Health Organization. Global prevalence of vitamin A deficiency in populations at risk 1995–2005: WHO global database on vitamin a deficiency. Geneva, Switzerland: World Health Organization; 2009.

[18] Olang B, Abdallah Z, Neshati R, Ali MA, Naghavi M, Agnetta Yrgev. Vitamin A status in pregnant women in Iran in 2001 and its relationship with province and gestational age. Food Nutr Rev 2014;58:25707. http://dx.doi.org/10.3402/fnr.v58.25707.

[19] Kassaye T, Receveur O, Johns T, Becklake MR. Prevalence of vitamin A deficiency in children aged 6–9 years in Wukro, northern Ethiopia. Bull World Health Organ 2001;79:415–21.

[20] Neyrolles O, Quintana-Murci L. Sexual inequality in tuberculosis. PLoS Med 2009;6(12):e1000199 http://dx.doi.org/10.1371/journal.pmed.1000199.

[21] WHO. World Health Organization. Indicators for assessing vitamin A deficiency and their application in monitoring and evaluation of intervention programs. 1996.

[22] Stephens CB. Vitamin A, infection, and immune function. Annu Rev Nutr 2001;21:167–92.

[23] Koyanagi A, Kuffo D, Gresely L, Shenkian A, Cuevas LE. Relationships between serum concentrations of C-reactive protein and micronutrients, in patients with tuberculosis. Ann Trop Med Parasitol 2004;98:391–9.

[24] Rohini K, Srikumar PS, Sazena K, Kumar MA. Alteration in the levels of serum micronutrients in tuberculosis patients. Int J Biol Med Res 2011;2(1):25707.

[25] Choi R, Kim HT, Lim Y, Kim MJ, Kwon OJ, Jeon K, Park HY, Jeong BH, Koh JW, Lee SY. Serum concentrations of trace elements in patients with tuberculosis and its association with dietary, serum albumin and C-reactive protein concentrations. Nutr Res 2011;31(4):303.

[26] Kassaye T, raceeveur O, Johns T, Becklake MR. Prevalence of vitamin A deficiency in children aged 6–9 years in Wukro, northern Ethiopia. Bull World Health Organ 2001;79:415–21.

[27] Gibson RS. Nutritional assessment. New York, NY: Oxford University Press; 1993.

[28] Paton NI, Chua YK, Earnest A, Chee CB. Randomized controlled trial of nutritional interventions for children and adolescents with pulmonary tuberculosis patients in some selected public health facilities of Addis Ababa, Ethiopia. A cross-sectional study. BMC Nutr 2017;3:93–100.

[29] Habte TY, Krawinkel M. Dietary diversity score: a measure of nutritional adequacy in Ethiopia. J Nutrit Health Food Sci 2018;6(1):1–7. http://dx.doi.org/10.15744/2393-6177.

[30] WHO. World Health Organization. Indicators for assessing infant and young child feeding practices Part 1 Definitions. Food and nutrition technical assistance; 2007.

[31] Supplementary materials associated with this article can be found, in the online version, at doi:10.1016/j.jctube.2018.05.002.