INTRODUCTION

Tuberculosis, a disease spread by obligate intracellular organism *Mycobacterium tuberculosis*, is probably one of the oldest diseases known to mankind. Tuberculosis is an important issue of concern for public health globally and in the year 2013, nine million new cases of tuberculosis and 1.5 million deaths were reported worldwide due to it.¹ In Pakistan, tuberculosis has been an important area for concern because of its increasing prevalence. According to the WHO, in 2013, the prevalence of all forms of tuberculosis in Pakistan was reported to be 275 cases per 100,000 population and currently Pakistan ranks fifth among countries worst-affected by the disease.² Even though the disease is curable, the treatment is long and has risks of treatment failure or relapse. This highlights
Hypovitaminosis D & associated risk factors in pulmonary TB

In recent years there has been escalating evidence which highlights the significance of vitamin D role in the response of immune system against mycobacterium tuberculosis which is an important predictor in the possible outcomes of tuberculosis. There is up regulation of expression of vitamin D receptor and 25-hydroxyvitamin D-1α-hydroxylase after the activation of a macrophage or monocyte through stimulation of its toll-like receptor 2/1 (TLR2/1) by Mycobacterium tuberculosis or other infectious agents. Research suggests that levels of 25-hydroxyvitamin D \[\text{upto} 30 \text{ ng/ml or higher provides adequate substrate for conversion of 25(OH) D by 1-OHase to its active form, 1, 25 dihydroxyvitamin D}\] which thereafter is able to penetrate into the nucleus, to eventually enhance the expression of cathelicidin which is an anti mycobacterial peptide.

If levels of 25(OH) D in serum are below 20 ng/ml, ability of the monocyte or macrophage to initiate this innate immune response is greatly compromised. Furthermore, the enhanced synthesis of 1,25(OH)\(_2\)D in monocytes and macrophages results in increased ability of activated T lymphocytes to act locally and enhance synthesis of cytokine as well as activate B lymphocytes, which regulate immunoglobulin synthesis. Therefore the factors which contribute towards low plasma levels of Vitamin D are also documented to promote and predispose to pulmonary TB.

The role of host immunity and its influence on the host-pathogen interaction is very important in the possible outcomes after being infected by mycobacterium tuberculosis. The relationship between hypovitaminosis D and predisposition to tuberculosis has been observed in England and African countries but documented studies for Pakistan are scarce. The aim of present study was to find out the frequency of hypovitaminosis D and its associated risk factors in newly diagnosed pulmonary tuberculosis patients prior to administration of standard anti tuberculosis therapy.

**METHODS**

**Subjects:** Eighty newly diagnosed patients having pulmonary tuberculosis were enrolled from the OPD of Ojha Institute of Chest Diseases, Dow University of Health Sciences (DUHS) from November 2010 to March 2011. The patients were of either sex with ages ranging between 18-50 years. They were diagnosed according to the WHO criteria for pulmonary tuberculosis which is Two sputum smear examinations on direct smear microscopy positive for acid fast bacilli (AFB+); one sputum examination positive for AFB and radiographic abnormalities consistent with active pulmonary tuberculosis; or one sputum specimen positive for AFB and one culture positive for AFB. Patients with AFB sputum smear negative; other concomitant ailments such as chronic kidney disease, liver problems, pulmonary silicosis, patients already on anti tuberculosis medication, taking Vitamin D, pregnant or lactating women were excluded from the study.

**Study design:** It was a cross-sectional hospital based study which was conducted after obtaining approval from IRB (DUHS) (Reference number: IRB-165/DUHS-10) and BASR of DUHS (reference number: DUHS/DR/2010/485). After a written informed consent all newly diagnosed pulmonary tuberculosis patients were subjected for evaluation of serum vitamin D levels. Prior to commencing anti tuberculosis therapy, from each patient 10 ml blood sample was collected after an overnight fast and serum Vitamin D level was evaluated by using electrochemiluminescence immunoassay on a Roche Elecsys 10100/201 system. Sociodemographic features were recorded in a specially designed proforma.

**Statistical analysis:** Data was gathered through a detailed proforma completed for each case and ratios, percentages and mean ± SD were calculated through SPSS program 17. P value less than 0.05 was taken statistically significant.

**RESULTS**

Out of 80 study participants, 33 were males and 47 were females. Age ranged from 18 to 50 years with mean age 33.33±12.21 years. Average age of males was 35.03±10.11 and average age of females was 31.93±11.37 years. The average BMI in male patients was 18.1±0.3 and the average BMI in females was found to be 17.53±0.27. Occupation wise five were drivers, six were laborers, three were cooks, twelve were domestic servants, thirty two were housewives, two were government servants, eight were teachers and twelve were jobless. Moreover, 80% of male patients and 16% of female patients were smokers prior to diagnosis of tuberculosis.
Hypovitaminosis was present in all the cases. The prevalence of Vitamin D insufficiency and deficiency was found to be 60.60% and 39.39% in male tuberculosis patients and 61.70% and 38.29% in female tuberculosis patients respectively.

Vitamin D levels in male and female study participants with respect to insufficiency and deficiency are provided in Table-I. The socio-demographic features of study participants are given in Table-II. Levels of serum 25(OH)D observed in different socio-demographic variables are presented in Table-III.

**DISCUSSION**

In this study, we observed Vitamin D deficiency in newly diagnosed pulmonary tuberculosis patients. The high prevalence of vitamin deficiency seen among our study population could be due to nutritional factors as majority of the patients belonged to poor socioeconomic status. However, it is highly unlikely that poor nutrition could be the only reason for this deficiency as it is a well-known fact that only ten percent of the Vitamin D is obtained from diet and the rest (90%) of it is produced in the skin under the influence of ultraviolet sunlight of the sun. Thus, only nutritional aspects are not likely the basis of the high frequency of low vitamin D in TB patients.\(^\text{11,12}\)

The first reports about the possibility of relationship between vitamin D and tuberculosis surfaced twenty years ago,\(^\text{13}\) but since then there have been conflicting reports about any such association in the subsequent studies. A number of studies conducted on Asian and African immigrants in England, African immigrants living in Australia, Ugandan and Korean population have reported very low levels of 25(OH)D and higher prevalence of vitamin D deficiency in TB patients than non-TB individuals.\(^\text{9,10,14,15}\) However, there was contrasting evidence in studies conducted in Tanzania and Vietnam, which showed no considerable difference in 25(OH)D levels between TB cases and matched controls.\(^\text{16,17}\)

Vitamin D deficiency could be an antecedent risk factors for TB in a similar manner as the disease itself could lead to low 25(OH)D levels in TB patients. The correlation between vitamin D and TB is mediated through increased production of cathelicidin and localized action of the 1, 25(OH)\(_2\)D produced in monocytes or macrophages on activated B lymphocytes regulating immunoglobulin

| Variables | Frequency | Percentage (%) |
|-----------|-----------|----------------|
| Gender    |           |                |
| Male      | 33        | 41.25          |
| Female    | 47        | 58.75          |
| Age       |           |                |
| <20       | 39        | 48.75          |
| 20-30     | 20        | 25             |
| 30-40     | 13        | 16.25          |
| 40-50     | 8         | 10             |
| BMI       |           |                |
| <16       | 45        | 56.25          |
| 16-17     | 16        | 20             |
| 17.18.5   | 18        | 22.5           |
| 18.5-25   | 1         | 1.25           |
| Level of Education | | |
| Illiterate | 55 | 68.75 |
| Literate | 25 | 31.25 |
| a) Primary level | a) 18 | a) 72 |
| b) Secondary level | b) 07 | b) 28 |
| Occupation | | |
| House wife | 32 | 40 |
| Laborer | 6 | 7.5 |
| Driver | 5 | 6.25 |
| Cook | 3 | 3.75 |
| Domestic servant | 12 | 15 |
| Teacher | 8 | 10 |
| Government Servant | 2 | 2.5 |
| Jobless | 12 | 15 |
| Ethnicity | | |
| Punjabi | 07 | 8.75 |
| Pushto | 10 | 12.5 |
| Hazarwal | 09 | 11.25 |
| Siraiki | 24 | 30 |
| Sindhi | 14 | 17.5 |
| Kashmiri | 07 | 8.75 |
| Urdu speaking | 07 | 8.75 |
| Bengali | 02 | 2.5 |
| SES | | |
| less than 6,000 per month | 61 | 76.25 |
| 6,000 to 25,000 per month | 19 | 23.75 |
| Smoking status | | |
| Yes | 42 | 52.5 |
| No | 38 | 47.5 |
production activated T lymphocytes controlling cytokine synthesis. In our study, mean BMI for both male and female tuberculosis patients was below normal levels that is an indicator of the meager health and nutritional condition of these participants. None of the patients in the study had white collar jobs and all the patients belonged to lower SES. Sixty four (80%) tuberculosis patients were extremely poor and earned less than 6,000 rupees per month which further confirms the poor nutritional status and poor hygienic conditions of these patients. The positive connection between BMI and serum 25(OH)D in our study is not similar to negative relationships observed in studies conducted in western countries which are elucidated by sequestration of fat soluble vitamin D in adipose tissue. We suggest that the positive association between the two variables is due to the reason that low BMI in these patients reveals a prolonged period of less sun exposure and hence, decreased vitamin D status.

In male tuberculosis patients vitamin D insufficiency (52.63%) was slightly more common than Vitamin D deficiency (47.36%). In contrast, in female patients, vitamin D deficiency was 2.2 times more prevalent than vitamin D insufficiency. This could be a result of the poor sun exposure in female tuberculosis patients because of social and cultural reasons as most of them were house wives and wore veil (burqa) whereas most of the male patients had moderate sun exposure.

In a study conducted in Egypt, plasma levels of Vitamin D were found to be low at the time of diagnosis of pulmonary TB which decreased further at four and six months during treatment. These findings are in accordance with our results. Furthermore, in similar studies conducted in northern India and rural areas around Lahore, Pakistan, prevalence of Vitamin D deficiency was higher; and Vitamin D insufficiency was less compared to our findings. A reason for this finding could be due to the fact that both quoted studies study were conducted in very poor rural population where poor nutrition and very low BMI was reported. In another study reported in Tanzania, Vitamin D levels in TB patients were again found to be considerably lower than matched controls but plasma levels of Vitamin D were found to be negatively associated with weight and BMI of the TB patients. This finding was in contrast to our findings where plasma levels of Vitamin D were positively associated with the weight and BMI of the TB patients. Reason for this difference could be the fact that low BMI in our population was due to malnutrition whereas weight loss in the study conducted in Tanzania was artificially induced through surgical means which could have resulted in less sequestration of the fat soluble vitamin after the removal of excessive fat.

| Variable                          | n (%)  | Serum 25(OH) D, ng/ml | p-value |
|-----------------------------------|--------|-----------------------|---------|
| Gender                            |        |                       |         |
| Male                              | 37 (46.25) * | 19.45 * | 0.002   |
| Female                            | 43 (53.75) | 9.41     |         |
| Age                               |        |                       | 0.2     |
| <20                               | 39 (48.75) | 8.34     |         |
| 20-30                             | 20 (25) | 12.22     |         |
| 30-40                             | 13 (16.25) | 13.34    |         |
| 40-50                             | 8 (10)  | 9.13     |         |
| Ethnicity                         |        |                       |         |
| Punjabi                           | 07 (8.75) | 8.22     |         |
| Pushto                            | 10 (12.5) | 11.34    | 0.045   |
| Hazarawal                         | 09 (11.25) | 9.45     |         |
| Siraiki                           | 24 (30) * | 9.33 *   |         |
| Sindhi                            | 14 (17.5) | 12.38    |         |
| Kashmiri                          | 07 (8.75) | 12.5     |         |
| Urdu speaking                     | 07 (8.75) | 10.76    |         |
| Bengali                           | 02 (2.5)  | 10.88    |         |
| Occupation                        |        |                       |         |
| House wife                        | 32 (40) * | 9.87 *   |         |
| Laborer                           | 6 (7.5)  | 12.34    |         |
| Driver                            | 5 (6.25)  | 10.11    |         |
| Cook                              | 3 (3.75)  | 11.10    | 0.001   |
| Domestic servant                  | 12 (15)  | 12.29    |         |
| Teacher                           | 8 (10)   | 10.55    |         |
| Government servant                | 2 (2.5)  | 9.76     |         |
| Jobless                           | 12 (15)  | 8.76     |         |
| Marital status                    |        |                       |         |
| Married                           | 44 (55)  | 12.33    | 0.21    |
| Single                            | 36 (35)  | 11.34    |         |
| Separate/Widowed                  | 0        |           |         |
| SES                               |        |                       |         |
| less than 6,000 per month         | 61 (76)  | 12.37    | 0.018   |
| 6,000 to 25,000 per month         | 19 (24)  | 15.11    |         |
| Smoking                           |        |                       |         |
| Yes                               | 42 (52.5) | 13.97    | 0.85    |
| No                                | 38 (47.5) | 14.35    |         |
| BMI kg/m2                         |        |                       |         |
| <16                               | 45 (56.25) * | 9.89 * | 0.028   |
| 16-17                             | 16 (20)  | 12.7     |         |
| 17.18.5                           | 18 (22.5) | 11.41    |         |
| 18.5-25                           | 1 (1.25)  | 11.05    |         |

* = statistically significant.
One way ANOVA test.
In our study, all patients showed deranged Vitamin D levels prior to the administration of standard anti-tuberculosis therapy which means that this derangement was not drug induced. The fact that matched controls were not enrolled for comparison of the relevant variables can be considered a limitation of the study. An association of low plasma levels of Vitamin D with pulmonary tuberculosis was found but a causal relationship could not be established which is another limitation of our study.

CONCLUSION

Frequency of hypovitaminosis D is found to be 100% with insufficiency (61.25%) and deficiency (38.75%) in newly diagnosed pulmonary tuberculosis patients. Whereas gender, ethnicity, occupation and BMI were the risk factors associated with it. The fact that all tuberculosis patients showed derangements in vitamin D levels implies that vitamin D should be administered as adjuvant therapy with standard anti tuberculosis therapy. This might be helpful in reducing the severity of disease and preventing complications.

Conflict of Interest: None.

REFERENCES

1. WHO publishes Global tuberculosis report 2013; 2013. Available: http://www.who.int/tb/publications/global_report/en/
2. World Health Organization. Epidemiology, Global tuberculosis control, surveillance, planning and financing. WHO Report. 2011;p.6-33.
3. Nava-Aguilera E, Andersson N, Harris E, Mitchell S, Hamel C, Shea B, et al. Risk factors associated with recent transmission of tuberculosis: systematic review and meta-analysis. Int J Tuberc Lung Dis. 2009;13:17–26.
4. Martineau AR, Honecker F, Wilkinson RJ, Griffiths CJ. Vitamin D in the treatment of pulmonary tuberculosis. J Steroid Biochem Mol Biol. 2007;103:793–798.
5. Martineau AR, Hall BM, Maunsell ZJ, Newton SM, Davidson RN, Packe GE, et al. Vitamin D induces antimycobacterial immunity in vivo and in vitro. Thorax. 2005;60:i43. doi:10.1164/rcm.200701-007OC
6. Martineau AR, Wilkinson KA, Newton SM, Floto RA, Norman AW, Skolimowska K, et al. IFN-gamma and TNF-independent vitamin D-inducible human suppression of mycobacteria: the role of cathelicidin LL-37. J Immunol. 2007;178:7190–7198.
7. Kim JH, Park JS, Cho YJ, Yoon HI, Song JH, Lee, CT, et al. Low serum 25-hydroxyvitamin D level: An independent risk factor for tuberculosis? Clin Nutr. 2013;32:614–619. doi:10.1016/j.clnu.2013.11.014.
8. Nnoaham KE, Clarke A. Low serum vitamin D levels and tuberculosis: a systematic review and meta-analysis. Int J Epidemiol. 2008;37:113–119. doi:10.1093/ije/dyn247.
9. Wejse C, Olesen R, Rabna P, Kaestel P, Gustafson P. Serum 25-hydroxyvitamin D in a West African population of tuberculosis patients and unmatched healthy controls. Am J Clin Nutr. 2007;86:1376–1383.
10. Wilkinson RJ, Llewelyn M, Toossi Z, Patel P, Pasvol G, Lalvani A, et al. Influence of vitamin D deficiency and vitamin D receptor polymorphisms on tuberculosis among Gujarati Asians in west London: a case-control study. Lancet. 2000;355:618–621.
11. Hewison M. Vitamin D and the immune system: new perspectives on an old theme. Endocrinol Metab Clin North Am. 2010;39:365–379. doi:10.1016/j.men.2010.02.010
12. Chocano-Bedoya P, Roonenberg AG: Vitamin D and tuberculosis. Nutr Rev. 2009;67(5):289-293. doi:10.1111/j.1753-4887.2009.00195.x
13. Grange JM, Davies PD, Brown RC, Woodhead JS, Kardjito T. A study of vitamin D levels in Indonesian patients with untreated pulmonary tuberculosis. Tubercle. 1985;66:187–191.
14. Kibirige D, Mutebi E, Sekeotileko R, Woroedria W, Mayanja-Kizza H. Vitamin D deficiency among adult patients with tuberculosis: a cross sectional study from a national referral hospital in Uganda. BMC Research Notes. 2013;6:293. doi:10.1186/1756-0509-6-293
15. Hong JY, Kim SY, Chung KS, Kim EY, Jung JY, Park MJ, et al. Association between vitamin D deficiency and tuberculosis in a Korean population. Int J Tuberc Lung Dis. 2014;18:73–78. doi:10.5588/ijtld.13.0536.
16. Friis H, Range N, Changalucha J, PrayGod G, Jeremiah K, Faurch-Jepsen D, et al. Vitamin D Status among Pulmonary TB Patients and Non-TB Controls: A Cross-Sectional Study from Mwanza, Tanzania. PLoS ONE. 2015;10(12):e014142. doi:10.1371/journal.pone.014142.
17. Ho-Pham LT, Nguyen ND, Nguyen TT, Nguyen DH, Bui PK, Nguyen VN, et al. Association between vitamin D insufficiency and tuberculosis in a vietnamese population. BMC Infect Dis. 2010;10:306. doi:10.1186/1471-2334-10-306
18. Sita-Lumsden A, Lathporn G, Swamathan R, Milburn HJ. Reactivation of tuberculosis and vitamin D deficiency: the contribution of diet and exposure to sunlight. Thorax. 2007;62:1003–1007. doi:10.1136/thx.2006.070060
19. Edem VF, Ige O, Arinola OG. Plasma vitamins and essential trace elements in newly diagnosed pulmonary tuberculosis patients and at different durations of anti-tuberculosis chemotherapy. Egyptian Journal of Chest Diseases and Tuberculosis. 2015;64:675–679.
20. Karoli R, Fatima J, Gupta SS, Shukla V, Moidurrehman, Manhar M. Vitamin D Deficiency in Medical Patients at a Teaching Hospital in North India. J Assoc Physicians India. 2015;63:35-39.
21. Junaid K, Rehman A, Saeed T, Jolliffe DA, Wood K, Martineau AR. Genotype-independent association between profound vitamin D deficiency and delayed sputum smear conversion in pulmonary tuberculosis. BMC Infectious Diseases. 2015;15:275. doi:10.1186/s12879-015-1018-5

Authors’ Contributions:

Fahad Azam: Conceived the idea, designed the study; collected, analysed and interpreted data; drafted the manuscript and made the changes suggested by reviewers for the final version of the manuscript.

Abida Shaheen: Analysed and interpreted the data; drafted the manuscript and approved the final version of the manuscript.

Rabia Arshad: Conceived and designed the study, collected, analysed interpreted data; drafted the manuscript.

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