

International Journal of Medical Science and Current Research (IJMSCR) Available online at: www.ijmscr.com Volume2, Issue 2, Page No: 100-105 March-April 2019



Vitamin D Level and Risk of Community Acquired Pneumonia in adults-an experience at tertiary care hospital

Sheenam Gazala¹, Mohammad Akbar Shah², Bathsheeba Andrabi³, Basharat Kassana⁴, Yasir Bashir⁵, Nusrat Bashir⁶ ¹ Senior Resident Emergency Medicine. ² Assistant Professor Critical Care Medicine

 ¹ Senior Resident Emergency Medicine. ² Assistant Professor Critical Care Medicine ³ Senior Resident Critical Care Medicine ^{1,2,3} SKIMS Srinagar
 ⁴ Consultant J&K Health Services ⁵. Assistant Professor, Medicine GMC, Baramulla

⁶ Assistant Professor, Pathology GMC Srinagar

J&K, India..

*Corresponding Author:

Dr.Nusrat bashir Assistant professor,department of pathology, GMC,Srinagar,J&K,India. 190010

Type of Publication: Original Research Paper Conflicts of Interest: Nil

ABSTRACT

Aims and objectives:

(a)To compare the levels of vitamin D between patients of Community acquired pneumonia and healthy controls.

(b) Association of Vitamin D deficiency with pneumonia.

Materials and methods: It was a prospective case control study.

Results and conclusions: Patients with pneumonia had a statistically significant low Vitamin D levels.

Keywords: Vitamin D, Pneumonia, Community.

INTRODUCTION

Till a few decades ago, vitamin D was thought of only in relation to bone health and calcium homeostasis. It stands at the frontline of current scientific endeavors, being a topic of greatest interest to medical researchers all over the globe. A growing body of evidence, implicating hypovitaminosis D as a risk factor for many diseases Vitamin D is important for the normal activity of skeletal and non-skeletal immune cells and immunity, tissues. bone calcification, and brain processes (1). Management of widespread vitamin D deficiency may fetch profound future health benefits. Vitamin D is involved in the regulation of 1000 human genes. Vitamin D metabolites play an important role in the bodys immunity by induction of phagocyte migration, modulation of the Th1–Th2-cellbalance, and the differentiation of T regulatory cells. (2) (3). It is shown that vitamin D deficiency (<20 ng/mL) is a prevalent condition and may be a key contributor to both acute and chronic infectious diseases including sepsis, pneumonia, urinary tract infections, and surgical site infections. Most cells, suchas B and T lymphocytes, monocytes, and dendritic cells, have specific vitamin D receptors (VDRs). Vitamin D demonstrates its immunomodulatory effects on these cell lines through its effects on the VDR. Deficiency is associated with reduced innate immunity and an increased risk for infections. Vitamin D deficiency

10(

.....

can positively affect a wide variety of microbial infections such as Gram-positive and Gram-negative bacteria, fungi, mycobacteria, and viruses. (4).

Pneumonia is a common lung disease which is responsible for significant morbidity and mortality worldwide [5, 6]. Community-acquired pneumonia (CAP)

is the main type of pneumonia which can also result in high risk of mortality in

Critically ill patients [7-9]. Risk stratification is crucial for the treatment of pneumonia, because it identifies patients with high risk of mortality for whom

Intensive care can be used [10, 11]. Therefore, to improve the accuracy of risk stratification, there is need to find more appropriate factors for predicting the risk of mortality among pneumonia patients. Such findings will provide much help in developing some effective risk stratification tools for evaluating the prognosis of pneumonia patients [12, 13]. Currently, some blood biomarkers, such as

C-reactive protein, procalcitonin and proadrenomedullin are thought to be associated with the survival of pneumonia patients [14-16]. Vitamin D plays important roles in regulating calcium homeostasis and immune response [17-19].

Results from previous studies suggest that vitamin D has a protective effect against respiratory tract infections [20, 21]. It has been suggested that vitamin D may exert immunomodulatory effects in patients with pneumonia [22, 23]. Low vitamin D status or vitamin D deficiency is common in the elderly and in critically- ill patients, as well as in patients with pneumonia [24-26]. Some studies reported that low vitamin D status was a risk factor for adverse outcomes among pneumonia patients, but results from some other studies were in disagreement with these reports [27-33]. T Given the importance of pneumonia and because vitamin D deficiency is one of the possible factors involved in susceptibility to pneumonia, further studies on the possible role of vitamin D deficiency in the prevalence or severity of the disease can be useful

Materials and methods:

The study was conducted in emergency and critical care department of our hospital for a time period of 1

year from march 2015 to march 2016 in patients with clinical history of pneumonia.

Study design: IT was a prospective case control study.

Cases: Patients who fulfilled the inclusion criteria

Inclusion criterion:

Age: greater or equal than 20 years.

Presence of new infiltrates on chest radiography.

Presence of one of the following criteria:

- 1. Productive Cough
- 2. Fever > = 38.3 degree centigrade.

3. Chest sign on auscultation.

Exclusion criterion:

Patient with active TB

Immunosuppressed patients.

Controls: Patients more than 20 years attending the hospital for complaints Other than respiratory symptoms. These patients were taken in study after taking proper consent.

A total of 80 patients were taken.40 cases and 40 controls after meeting inclusion criteria.

Methodology:

1- History taking including socio-economic level, residence, and history of upper respiratory tract infection

2- Clinical examination including body temperature, respiratory rate, cyanosis and local chest examination.

3- Routine laboratory investigations including complete blood count (CBC),

C-reactive protein (CRP) (positive above 6 mg/L) and erythrocyte sedimentation rate (ESR).

4- Chest X ray.

5- Measurement of serum Vitamin-D (25OHD): Two ml of venous blood Specimens was collected from a peripheral vein and serum separated by centrifugation and then stored at -20° C and protected from direct exposure to sunlight until the analysis. The timing of blood sample collection within the disease course varied from patient to patient. However, given that

vitamin D has a half-life of between 4 and 8 weeks, the small inconsistency in timing would not be expected to affect the average vitamin D level. Samples were assayed at the end of the enrollment period using a commercially available 25-hydroxy vitamin D3 RIA kit(25-Hydroxyvitamin D 125 RIA Kit; DiaSorin Diagnostics) according to the instructions provided by the manufacturer. All the determinations were performed in the Department of Immunology, Sher-e-Kashmir Institute of Medical Sciences, Soura, Srinagar.

Results:

There were 62.5% males and 37.5% females in the case group and 67.5% males and 32.5% females in the control group. The P value of 0.639 is statistically insignificant.

Hundred percent of cases were found to have deficient 25 hydroxy vitamin D levels whereas 52.5% controls were found to have deficient 25 hydroxy vitamin D levels. The P- value of <0.001 was found to be significant.

The mean serum levels of 25-hydroxy vitamin D levels of cases was found to be 5.14 ± 2.793 ng/ml and of controls it was 15.84 ± 8.519 ng/ml with a significant P-value of <0.001.

The mean serum vitamin D levels among males in case group was 4.81 ± 2.989 ng/ml and in control group was 16.74 ± 8.186 ng/ml. The P- value of <0.001 was found to be significant.

The mean serum vitamin D levels among females in case group was 5.69 ± 2.430 ng/ml and in control group was 13.98 ± 9.223 ng/ml with a significantP-value of <0.001.

Table 1a-(case group)

Total patients(Percentage)	Males(percentage)	Females(percentage)
40(100)	25(62.5)	15(37.5)

Table 1b-(control group)

Total controls (Percentage)	Males(percentage)	Females(percentage
40(100)	27(67.5)	13(32.5)

P Value (0.639)-Insignificant

Table 2(vitamin D sufficiency)

Case group(40)		Control group(40)			
Vitamin-D Sufficent	Vitamin-D Sufficent	In-	Vitamin-D Sufficent	Vitamin-D Sufficent	In-
0(0%)	40(100%)		19(47.5%)	21(52.5%)	
5.14 ± 2.793 ng/ml(Mean value)		15.84 ± 8.519 ng/ml(Mean value)			

P Value (<0001.)-significant.

Table 3 (Mean vitamin D levels)

Case group		Control group		
Males	Females	Males	Females	
4.81 ± 2.989 ng/ml	5.69 ±2.430 ng/ml	16.74 ± 8.186 ng/ml	13.98 ± 9.223 ng/ml	

P Value (<0001.)-significant

Dr. Nusrat Bashir et al International Journal of Medical Science and Current Research (IJMSCR)

.....

Discussion: The purpose of this case-control study was to investigate the relationship between vitamin D levels and severity of CAP in a total of 80 subjects (40 cases and 40 controls). The overall prevalence of vitamin D deficiency in the case group was greater than in the control group; however, patients with CAP had a higher rate of severe vitamin D deficiency in comparison with healthy participants. Furthermore, CAP patients with severe vitamin D deficiency had more severe disease (higher CURB-65 score). Although not statistically significant, patients with severe deficiency had a higher rate of ICU admission and a longer duration of hospital stay.

In a study conducted by Mojgan Mamani³⁴ et al In total, 81.2% of the study population had vitamin D levels <30 ng/dL. The risk of pneumonia among subjects with deficient vitamin D levels was 3.69 (95% CI: 1.46, 9.31) times of those with sufficient vitamin D level (P=0.006). Prevalence of severe deficiency of vitamin D in scores three and four of CURB-65 (59.38%), was far more than scores one and two (31.71%). Also, results indicated patients with severe deficiency had a higher risk for ICU 30-day admission, mortality, and longer hospitalization stay, but these were not statistically significant. Similar results were found in our study as well. he results of our study showed that there was a significant association between severe vitamin D deficiency and CAP. Individuals with severe deficiency were 4.16 times more susceptible in acquiring CAP when compared to the people with higher levels of vitamin D. Our result is consistent with the recent studies conducted in other countries,³⁵⁻³⁸ although this relationship was observed for different levels of vitamin D in other studies.

Conclusion: Patients with pneumonia had a statistically significant low Vitamin D levels

References:

- Holick MF. Vitamin D deficiency. N Engl J Med. 2007;19(357(3)):266–281.
- Gombart AF, Borregaard N, Koeffler HP. Human cathelicidin antimicrobial peptide (CAMP) gene is a direct target of the vitamin D receptor and is strongly up-regulated in myeloid cells by 1,25-dihydroxyvitamin D3. FASEB J. 2005;19(9):1067–1077.2.

- Fabri m,stenger S, shin DM ,Yuk jm,liu PT etal.(2011)Vitamin d is required for IFNgamma –mediated antimicrobial activity of human macrophages.Sci Transl Med 3:104ral 102 Pub Med:21998409.
- 4. Youssef DA, Ranasinghe T, Grant WB, Peiris AN. Vitamin D's potential to reduce the risk of hospital-acquired infections. Dermatoendocrinol. 2012;4(2):167–175.
- Cillóniz C, Torres A, Niederman M, Eerden MVD, Chalmers J, Welte T, Blasi F.Community acquired pneumonia related to intracellular pathogens. Intensive Care Med 2016; 42(9): 1374-1386
- Lee JS, Giesler DL, Gellad WF, Fine MJ. Antibiotic Therapy for Adults Hospitalized With CommunityAcquired Pneumonia: A Systematic Review. JAMA 2016; 315(6): 593-602.
- Kishaba T. Community-Acquired Pneumonia Caused by Mycoplasma pneumoniae: How Physical and Radiological Examination Contribute to Successful Diagnosis. Front Med 2016; 3(1):28.
- Phua J, Dean NC, Guo Q, Kuan WS, Lim HF, Lim TK. Severe community-acquired pneumonia: timely management measures in the first 24 hours. Critical Care 2016; 20(1): 237.
- Haq IJ, Battersby AC, Eastham K, Mckean M. Community acquired pneumonia in children. BMJ 2017; 356: j686.
- 10. Aoki T. Risk Stratification of Elderly Community-acquired Pneumonia by Adding Computed Tomography. Intern Med 2016; 55(5): 425-426.
- Kaysin A, Viera AJ. Community-Acquired Pneumonia in Adults: Diagnosis and Management. Am Family Physician 2016; 94(9): 698-706.
- 12. Liapikou A, Torres A. Current treatment of community acquired pneumonia.
- 13. Exp Opin Pharmacother 2013; 14(10): 1319-1332.
- 14. Corrales-Medina VF, Taljaard M, Fine MJ, Dwivedi G, Perry JJ, Musher DM, Chirinos JA. Risk stratification for cardiac complications in patients hospitalized for

.

Dr. Nusrat Bashir et al International Journal of Medical Science and Current Research (IJMSCR)

community-acquired pneumonia. Mayo Clin Proc 2014; 89(1): 60-68.

- 15. Florin TA, Ambroggio L. Biomarkers for CommunityAcquired Pneumonia in the Emergency Department. Curr Infect Dis Rep 2014; 16(12): 451.
- 16. Agnello L, Bellia C, Di GM, Lo SB, Calvaruso L, Bivona G, Scazzone C, Dones P, Ciaccio M. Utility of serum procalcitonin and C-reactive protein in severity assessment of community-acquired pneumonia in children. Clin Biochem 2016; 49(1-2): 47-50.
- 17. Andersen SB, Baunbæk EG, Jensen AV, Petersen PT, Rohde G, Ravn P. Failure of CRP decline within three days of hospitalization is associated with poor prognosis of Community-acquired Pneumonia. Infect Dis 2017: 251-260.
- 18. Bikle DD. Extraskeletal actions of vitamin D. Ann N Y Acad Sci 2016; 1376(1): 29-52.
- 19. da Costa DS HJ, Ferreira TB, Kasahara TM, Barros PO, Monteiro C, et al.
- 20. Vitamin D modulates different IL-17secreting T cell subsets in multiple sclerosis patients. J Neuroimmunol 2016; 299(1): 8-18.
- 21. Groseanu L, Bojinca V, Gudu T, Saulescu I, Predeteanu D, Balanescu A,
- 22. Berghea F, Opris D, Borangiu A, Constantinescu C, Negru M, Ionescu R. Low vitamin D status in systemic sclerosis and the impact on disease phenotype. Eur J Rheumatol 2016; 3(2): 50-55.
- 23. Gonçalves de Carvalho CM, Ribeiro SM. Aging, lowgrade systemic inflammation and vitamin D: a minireview. Eur J Clin Nutr 2017; 71(4):434-440.
- 24. Kempker JA, West KG, Kempker RR, Siwamogsatham O, Alvarez JA, Tangpricha V, Ziegler TR, Martin GS. Vitamin D Status and the Risk for Hospital-Acquired Infections in Critically Ill Adults: A Prospective Cohort Study. Plos One 2015; 10(4): e0122136.
- 25. Watkins RR, Lemonovich TL, Salata RA. An update on the association of vitamin D deficiency with common infectious diseases. Can J Physiol Pharmacol 2015; 93(5): 363-368.

- 26. Borella E, Nesher G, Israeli E, Shoenfeld Y. Vitamin D: a new anti-infective agent? Ann N Y Acad Sci 2014; 1317(1): 76-83.
- 27. Esposito S, Lelii M. Vitamin D and respiratory tract infections in childhood. BMC Infect Dis 2015; 15(1): 487.
- 28. Inamo Y, Hasegawa M, Saito K, Hayashi R, Ishikawa T, Yoshino Y, Hashimoto K, Fuchigami T. Serum vitamin D concentrations and associated severity of acute lower respiratory tract infections in Japanese hospitalized children. Pediatr Int 2011; 53(2): 199-201.
- 29. Remmelts HH, Spoorenberg SM, Oosterheert JJ, Bos WJ, de Groot MC, van de Garde EM. The role of vitamin D supplementation in the risk of developing pneumonia: three independent case-control studies. Thorax 2013; 68(11): 990-996.
- 30. Sudfeld CR, Giovannucci EL, Isanaka S, Aboud S, Mugusi FM, Wang M, Chalamilla G, Fawzi WW. Vitamin D Status and Incidence of Pulmonary Tuberculosis, Opportunistic Infections, and Wasting Among HIV- Infected Tanzanian Adults Initiating Antiretroviral Therapy. J Infect Dis 2013; 207(3): 378-385.
- 31. Muhe L, Lulseged S, Mason KE, Simoes EA. Casecontrol study of the role of nutritional rickets in the risk of developing pneumonia in Ethiopian children. Lancet 1997; 349(9068): 1801-1804.
- 32. Leow L, Simpson T, Cursons R, Karalus N, Hancox RJ. Vitamin D, innate immunity and outcomes in community acquired pneumonia. Respirology 2011; 16(4): 611-616.
- 33. Remmelts HH, van de Garde EM, Meijvis SC, Peelen EL, Damoiseaux JG, Grutters JC, Biesma DH, Bos WJ, Rijkers GT. Addition of vitamin D status to prognostic scores improves the prediction of outcome in community-acquired pneumonia. Clin Infect Dis 2012; 55(11): 1488-1494.
- 34. Kim HJ, Jang JG, Hong KS, Park JK, Choi EY. Relationship between serum vitamin D concentrations and clinical outcome of community-acquired pneumonia. Int J Tuberculosis Lung Dis 2015; 19(6): 729-734.

Dr. Nusrat Bashir et al International Journal of Medical Science and Current Research (IJMSCR)

-
 - 35. 32. Haliloglu M, Bilgili B, Haliloglu O, Gogas YD, Cinel I. Vitamin D level is associated with mortality predictors in ventilator-associated pneumonia caused by Acinetobacter baumannii. J Infecti Dev Ctries 2016; 10(6):567-574.
 - 36. 33. Holter JC, Ueland T, Norseth J, Brunborg C, Frøland SS, Husebye E, Aukrust P,Heggelund L. Vitamin D Status and Long-Term Mortality in Community-AcquiredPneumonia: Secondary Data Analysis from a Prospective Cohort. PloS One 2016;11(7): e0158536
 - 37. Association between serum concentration of 25-hydroxyvitamin D and communityacquired pneumonia: a case-control study Mojgan Mamani,^{1,2}Neda Muceli,²Hamid Reza Ghasemi Basir,³Maryam Vasheghani,⁴ and Jalal Poorolajal⁵Int J Gen Med. 2017; 10: 423–429.
 - 38. Kim HJ, Jang JG, Hong KS, Park JK, Choi EY. Relationship between serum vitamin D

concentrations and clinical outcome of community-acquired pneumonia. Int J Tuberc Lung Dis. 2015;19(6):729–734.

- 39. Quraishi SA, Bittner EA, Christopher KB, Camargo CA., Jr Vitamin D status and community-acquired pneumonia: results from the third National Health and Nutrition Examination Survey. PloS One. 2013;8(11):e81120.
- 40. Aregbesola A, Voutilainen S, Nurmi T, Virtanen JK, Ronkainen K, Tuomainen TP. Serum 25-hydroxyvitamin D3 and the risk of pneumonia in an ageing general population. J Epidemiol Community Health. 2013;67(6):533–536.
- 41. Wayse V, Yousafzai A, Mogale K, Filteau S. Association of subclinical vitamin D deficiency with severe acute lower respiratory infection in Indian children under 5 y. Eur J Clin Nutr.