# Importance of vitamin d level in early-stage chronic lymphocytic leukemia and its comparison with healthy population

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#### SUMMARY

Chronic lymphocytic leukemia (CLL) is a chronic lymphoproliferative hematological malignancy accompanied by a Chronic lymphocytic leukemia (CLL) is a chronic lymphoproliferative hematological malignancy accompanied by a monoclonal proliferation of mature B lymphocytes. Vitamin D is vital in calcium hemostasis in serum and skeletal system and is involved in several other cellular processes such as differentiation, proliferation, apoptosis, and angiogenesis. Vitamin D receptors were also demonstrated in normal or malignant hematopoietic cells. The correlation of low serum vitamin D levels with colorectal, breast, and other solid-organ neoplasms is well known. With regard to CLL, vitamin D inadequacy is related to bad prognosis, bad overall survival, and necessity to start treatment early. This study aimed to compare vitamin D levels of patients with early-stage CLL and controls and to examine its relationship with prognostic indicators.

The study included 33 patients with early-stage CLL treated at Atatürk Training and Research Hospital Hematology Clinic between 2015 and 2017 and 34 healthy controls. Age and gender distribution in both groups was similar. Patients with <20 ng/mL vitamin D level were considered as vitamin D deficient.

Vitamin D deficiency was detected in 26/33 patients with CLL and 16/34 controls. A statistically significant difference in vitamin D deficiency was found between these two groups (P = 0.003). CLL cases were compared in terms of RAI stage, splenomegaly, hepatomegaly, and presence of B symptoms. However, no significant difference was observed.

This study demonstrated that vitamin D levels of patients with early-stage CLL were significantly low in comparison with those of healthy population, in line with the literature. More comprehensive studies are required for assessing the effects of low vitamin D levels on the prognosis of patients with early-stage CLL.

Key words: Early-stage CLL, prognosis, vitamin D deficiency

## INTRODUCTION

Chronic lymphocytic leukemia (CLL) is a chronic lymphoproliferative hematological malignancy accompanied by a monoclonal proliferation of mature B lymphocytes. CLL is the most common type of adult leukemia with 5/100,000 incidence rate every year in Western countries (1–3). CLL is a disease that mostly affects elder; 80% of the patients are aged more than 50 years (4,5). However, CLL has a heterogeneous course according to the clinical and molecular characteristics (6,7).

Vitamin D deficiency is quite common around the world. Besides regulating serum calcium and skeletal homeostasis, vitamin D is involved in several other cellular processes including differentiation, proliferation, apoptosis, metastatic potential, and regulation of angiogenesis (8). Vitamin D receptors were also demonstrated in normal or malignant hematopoietic cells. Low serum vitamin D levels are related to colorectal, breast, and other solid-organ neoplasms (9–11).

A double-blind, randomized, and placebo-controlled study demonstrated that the cancer risk of women taking vitamin D replacement daily reduced by 60%–77%. (12). Vitamin D deficiency in patients with CLL has emerged as a new, independent, poor prognostic factor in recent years. (13). Meanwhile, low serum vitamin D levels are related to short overall survival and necessity to start treatment early (14).

TABLE 1: Age and gender characteristics of patients with CLL and healthy controls.

	CLL	Healthy controls
Age	60 (± SD 14)	58 (± SD 11.5)
Gender (F/M)	15 (45.5%)/ 18 (54.5%)	13 (38.9%)/ 21 (61.1%)
Vitamin D levels Normal >20 ng/mL Low <20 ng/mL), n (%)	3 / 33 (22%) 26 / 33 (78%)	18 / 34 (53%) 16 / 34 (47%)

The present study compared serum vitamin D levels of patients with early-stage CLL and normal healthy controls. It also examined the correlation between vitamin D levels and other risk factors such as RAI stage of the disease, B symptoms, and organomegaly.

## MATERIALS AND METHODS

This study included 33 patients with early-stage CLL at Atatürk Training and Research Hospital Hematology Clinic between 2015 and 2017. It also involved 34 healthy controls whose vitamin D level was checked during their routine follow-up. Vitamin D deficiency was considered as 20 ng/mL or lower. The studied blood samples were venous blood drawn after 8 hours of fasting. Vitamin D levels were checked by the liquid chromatography method. Data collected from computer registers were statistically analyzed using SPSS version 16.0 (SPSS Inc., IL, USA). Categorical variables were compared using the chi-square test, whereas numerical variables were compared using the Mann–Whitney U test. P values less than 0.05 were considered significant.

### RESULTS

Age, gender, and vitamin D levels of patients with CLL and healthy controls are shown in table 1. No statistically significant differences were found between two groups with respect to age and gender distribution. The average follow-up duration of patients with CLL was reported as 44 months (12–102 months). Details on stages of the patients, B symptoms (fever, night sweating, and weight loss), and organomegaly are summarized in Table 2.

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Further, 14 of 26 (53.8%) patients with CLL having low vitamin D level were males and 12 (46.2%) were females. Also, 13 (50%) of the patients with CLL having low vitamin D level were aged more than 65 years. With regard to RAI stages, 14 (53.9%) patients were at stage 2, 8 (30.8%) were at stage 1, and 4 (15.4%) were at stage 0. Moreover, 7 (26.9%) of the patients had hepatomegaly, 12 (46.2%) had splenomegaly, and 3 (11.5%) had B symptoms. Out of seven (21.2%) patients with CLL having normal vitamin D levels, four (47.1%) were at RAI stage 2 and three (42.9%) were at RAI stage 1. Five (71.4%) of the patients were aged 65 years or above; four (57.1%) were males and three (42.9%) were females. Three (42.9%) patients having normal vitamin D level had B symptoms.

Also, 10 of 16 (62.5%) controls having low vitamin D level were females and 6 (37%) were males. Six (37%) of these patients were aged 65 years or above.

Vitamin D deficiency was detected in 26 (78.8%) patients with CLL 16 (47%) controls. A statistically significant difference was observed between two groups in terms of vitamin D deficiency (P = 0.003). Patients with CLL were compared in terms of RAI

TABLE 2: Characteristics of patients with CLL.		
RAI stage (stage 0/1/2)	n: 4 (12.1%)/n: 19 (57.6%)/n: 10 (30.3%)	
Presence of B symptoms	5 (15.2%)	
Splenomegaly	14 (42.4%)	
Hepatomegaly	5 (5.2%)	
Hemoglobin	13.3 g/L(7.80–17.50 g/L)	
Thrombocyte count	191,000 ´ 106/L (10,000–335,000 ´ 106/L)	
Leukocyte count	14,900 ´ 106/L (1700–13,400 ´ 106/L)	
Lactate dehydrogenase	227 (140–396)	

stage, splenomegaly, hepatomegaly, and presence of B symptoms according to vitamin D levels, but no significant difference was observed.

## DISCUSSION

CLL is the most common type of adult-age leukemias. It is characterized by the increase in monoclonal B lymphocytosis (15). Usually, treatment and cure are not expected in patients with CLL. The treatment is focused on keeping the disease under control and establishing control over symptoms rather than on curing the disease. Therefore, it is significant to know the factors affecting prognosis in patient with noncurative CLL because the disease has a slow progression. Shanafelt et al. detected vitamin D deficiency in 30.5% of patients with CLL demonstrated a decline in the duration between vitamin D deficiency and the start of the treatment and total survival (14). They also reported that knowing vitamin D deficiency as a modifiable factor might be significant in patients recently diagnosed with B-CLL (14). Serum vitamin D deficiency is common around the world. It is encountered in the ratio of 25%–50% during routine clinical examination (16).

Studies on vitamin D deficiency in patients with CLL are rather limited. Parveen et al. detected vitamin D deficiency in 56.7% of patients with CLL (17). Vitamin D deficiency is detected in 82.2% of patients with early-stage CLL, which is a fairly high rate (18). It is argued that vitamin D levels might be a factor affecting the process of starting the treatment and maintaining normal vitamin D levels might retard disease progression in patients with CLL (18). Another study demonstrated that high vitamin D levels decreased CLL risk while vitamin D deficiency was accompanied by a progressive increase in the incidence of CLL (19).

Vitamin D deficiency was detected in 78% of patients with earlystage CLL, which is parallel to previous findings. The difference in data regarding vitamin D deficiency might be related to environmental factors, ethnic backgrounds, and dietary habits of societies.

In our study, vitamin D levels of 26/33 (78%) of patients with CLL were statistically significantly lower compared with those of healthy controls (P = 0.003).

Vitamin D is a multifunctional hormone (20). Several studies indicate that it is effective against cell proliferation, differentiation,

and apoptosis (21). B-CLL cells express vitamin D receptor more than normal B and T lymphocytes. The use of a pharmacological dose of vitamin D induced the apoptosis of CLL cells in vitro via p53independent mechanism (22). Vitamin D inhibited the proliferation of lymphocytes and induces differentiation in vitro (23,24). A study on 229 patients with CLL demonstrated no correlation between vitamin D deficiency, absolute number of lymphocytes, and advanced-stage disease (14).

The present study showed no correlation between RAI stage, presence of anemia, splenomegaly, hepatomegaly, presence of B symptoms, and vitamin D deficiency.

Although the correlation between vitamin D levels and solid tumor risk is well known, the relation between hematological malignancy and vitamin D levels needs further investigation. Previous studies aiming to explain the potential effect of vitamin replacement on cancer prevention did not always yield positive results (25). A study on 34 patients with low-grade NHL reported tumor regression in 24% of the patients treated with 1,25(OH)D' synthetic analog (four full remissions and four partial remissions) (26). The vitamin D levels of patients should be measured and monitored regularly to maintain an optimal level (27). A meta-analysis conducted in China reported that low serum vitamin D levels in hematological malignancy might be related to bad prognosis (21). Another study argued that vitamin D deficiency led to disease progression and negative response to treatment in patients with CLL (28). A 13-month remission was unexpectedly observed in a patient with CLL having vitamin D deficiency after administering cholecalciferol (29).

## CONCLUSIONS

A limitation of this study was that it was performed on a limited number of patients with early-stage CLL. However, this was the first study on vitamin D levels in patients with early-stage CLL in Turkey. The vitamin D deficiency rate was found to be very high in Turkish patients. The measurement of vitamin D levels in patients with early-stage CLL and adequate replacement might help in controlling disease progression and improving response to treatment. Therefore, vitamin D levels can be considered as a prognostic indicator of the disease independent of its clinical stage. Larger-scale, randomized, controlled studies should be conducted to validate the findings.

## REFERENCES

- Xu W, Li JY, Miao KR, et al. The negative prognostic significance of positive direct antiglobulin test in Chinese patients with chronic lymphocytic leukemia. Leuk Lymphoma 2009;50:1482-7.
- Rozman C, Montserrat E. Chronic lymphocytic leukemia. N Engl J Med 1995;1333:1052-57.
- 3. Zwiebel JA, Cheson BD. Chronic lymphocytic leukemia: staging and prognostic factors. Semin Oncol 1998;25:42-59.
- 4. Mauro FR, Foa R, Giannarelli D, et al. Clinical characteristics and outcome of young chronic lymphocytic leukemia patients: a single institution study of 204 cases. Blood 1999;94:448-54.
- Zeeshan R, Sultan S, Irfan SM, Kakar J, Hameed MA. Clinicohematological profile of patients with B-chronic lymphoid leukemia in Pakistan. Asian Pac J Cancer Prev 2015;16:793-6.
- Mozaheb Z, Hasanzadeh N, Abadi MH, Aghaee MA. Chronic lymphocytic leukemia and prognostic factors. Asian Pac J Cancer Prev 2012;13:3009-13.
- Zeeshan R, Irfan SM, Sultan S, Bhimani S. ZAP-70 protein expression in B-cell Chronic Lymphoid Leukemia: A single center experience from Pakistan. Asian Pac J Cancer Prev 2015;16:1587-90.
- Bikle D. Nonclassic actions of vitamin D. J Clin Endocrinol Metab. 2009;94(1):26-34.
- Yin L, Grandi N, Raum E, et al. Meta-analysis: longitudinal studies of serum vitamin D and colorectal cancer risk. Aliment Pharmacol Ther 2009;30:113-25.
- Chen P, Hu P, Xie D, et al. Meta-analysis of vitamin D, calcium and the prevention of breast cancer. Breast Cancer Res Treat 2010;121: 469-77.
- 11. Garland CF, Gorham ED, Mohr SB, Garland FC. Vitamin D for cancer prevention: global perspective. Ann Epidemiol 2009;19:468-83.
- Lappe JM, Travers-Gustafson D, Davies KM, Recker RR, Heaney RP. Vitamin D and calcium supplementation reduces cancer risk: results of a randomized trial. Am J Clin Nutr 2007;85(6):1586-1591.
- Aref S, Ibrahim L, Azmy E. Prognostic impact of serum 25-hydroxivitamin D (25(OH)D) concentrations in patients with lymphoid malignancies. Hematol 2013;18:20-5.
- 14. Shanafelt TD, Drake MT, Maurer MJ, et al. Vitamin D insufficiency and prognosis in chronic lymphocytic leukemia. Blood 2011;117:1492-8.
- Kermani IA, Dehdilani M, Dolatkhah R. Chronic lymphocytic leukemia in the recent 10 years and treatment effects of Fludarabin. Asian Pac J Cancer Prev 2007;8(3):367-71.

- 16. Holick MF. Vitamin D deficiency. N Engl J Med 2007;357(3):266-81.
- Parveen SI, Zeeshan R, Sultan S, Irfan SM.Serum 25-hydroxyvitamin D Insufficiency in B-Chronic Lymphoid Leukemia at the Time of Disease Presentation in Pakistan. Asian Pac J Cancer Prev 2015;16(14):5983-6.
- Molica S, Digiesi G, Antenucci A, et al. Vitamin D insufficiency predicts time to first treatment (TFT) in early chronic lymphocytic leukemia (CLL). Leuk Res 2012;36:443-7.
- Łuczyńska A, Kaaks R, Rohrmann S, et al. Plasma 25-hydroxyvitamin D concentration and lymphoma risk: results of the European Prospective Investigation into Cancer and Nutrition. Am J Clin Nutr 2013;98:827-38.
- Feldman D, Krishnan AV, Swami S, Giovannucci E, Feldman BJ. The role of vitamin D in reducing cancer risk and progression. Nat Rev Cancer 2014;14:342-57.
- Wang W, Li G, He X, et al. Serum 25-hydroxyvitamin d levels and prognosis in hematological malignancies: a systematic review and meta-analysis. Cell Physiol Biochem 2015;35:1999-2005.
- Pepper C, Thomas A, Hoy T, et al. The vitamin D3 analog EB1089 induces apoptosis via a p53-independent mechanism involving p38 MAP kinase activation and suppression of ERK activity in B-cell chronic lymphocytic leukemia cells in vitro. Blood 2003;101: 2454-60.
- Provvedini DM, Tsoukas CD, Deftos LJ, Manolagas SC. 1,25-dihydroxyvitamin D3 receptors in human leukocytes. Science. 1983;221(4616):1181-1183.
- Hickish T, Cunningham D, Colston K, et al. The effect of 1,25-dihydroxyvitamin D3 on lymphoma cell lines and expression of vitamin D receptor in lymphoma. Br J Cancer 1993;68(4):668-672.
- Greenwald P, Anderson D, Nelson SA, Taylor PR. Clinical trials of vitamin and mineral supplements for cancer prevention. Am J Clin Nutr 2007;85(1): 314S-317S
- Raina V, Cunningham D, Gilchrist N, Soukop M. Alfacalcidol is a nontoxic, effective treatment of follicular small-cleaved cell lymphoma. Br J Cancer 1991;63(3):463-465.
- Heaney RP. The Vitamin D requirement in health and disease. J Steroid Biochem Mol Biol 2005; 97(1-2):13-19.
- Thomas X, Chelghoum Y, Fanari N, Cannas G (2011). Serum 25-hydroxyvitamin D levels are associated with prognosis in hematological malignancies. Hematol 16, 278-83.
- 29. Arlet JB, Callens C, Hermine O, et al. Chronic lymphocytic leukaemia responsive to vitamin D administration. Br J Haematol 2012;156, 148-9.