

Vitamin D insufficiency among children with cancer in India

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INTRODUCTION

Vitamin D and its numerous implications on health have been a major field of research in the preceding decades, and the results from these studies have been found to be revelatory, prompting the need for routine Vitamin D level assays in both the general population and among children with cancer. Vitamin D levels have been implicated in the occurrence of a wide variety of diseases such as asthma,^[1-3] diabetes,^[4-6] heart disease,^[5,7] and cancer^[6] where it was once thought the ultimate manifestation of Vitamin D deficiency was rickets or osteomalacia.^[8-10] Although several studies have been done internationally on this topic,^[11-14] none has been carried out thus far in an Indian population. The aim of our study was to find the prevalence of Vitamin D insufficiency in children with cancer and once found to compare it with the prevalence of the same among the general population and also to compare Vitamin D levels based on characteristics such as age, sex, duration of therapy, and type of cancer.

MATERIALS AND METHODS

Study population and inclusion criteria

Vitamin D concentration in serum was measured from 51 cases and 51 controls over a span of 5 months from

ABSTRACT

Background: Vitamin D plays an important role in regulating various homeostatic mechanisms and has yet untapped potential in cancer prevention and prognosis. Only a few studies have been done worldwide in relating the Vitamin D levels in pediatric cancer patients to the general population but none so far in an Indian setting to the best of our knowledge. **Objective:** To compare the Vitamin D levels in a group of children with cancer to that of the general pediatric population and to note differences in the prevalence of Vitamin D insufficiency and make inferences arising from demographic and therapeutic variations. **Materials and Methods:** Vitamin D levels were found by immuno-chemilumino-metric assay in 102 children (51 cases and 51 controls) over a 6 months period. **Results:** In comparing the Vitamin D levels of children with cancer and controls from a healthy population we found an increased incidence of Vitamin D insufficiency in cancer children (80.39%) when compared to controls (50.98%) and a much lower mean Vitamin D value in cancer children (22.8 ng/ml) when compared to controls (33 ng/dl). It was also found that cancer children above 6 years had a greater chance for developing Vitamin D insufficiency ($P = 0.038$) as did children suffering from hematological malignancies ($P = 0.025$). **Conclusion:** Our study showed an increased prevalence of Vitamin D insufficiency in children with cancer and hence we suggest routine measurement of Vitamin D levels in children with cancer and subsequent supplementation.

Key words: Childhood cancer, India, Vitamin D

March 2013 to July 2013 in our center. Inclusion criteria for subjects were children aged 0-18 years previously or newly diagnosed with malignancy and on treatment whose Vitamin D levels were measured at least once during admission. Only the first measurement of 25-hydroxyvitamin D (25-OH D) was considered and children receiving supplementation were excluded from the study. Exclusion criteria also included children of therapy for a period >2 years, patients suffering from a disease that appeared due to Vitamin D deficiency and patients above 18 years of age at diagnosis. Approval for the study was obtained from the Institute's Ethics Committee.

Data collection

Demographic data collected from our subjects included age and sex. Age was recorded in years and categorized into two groups, those below 6 years and those above 6 years as seen in healthy populations.^[15-17] Data were also collected on two other criteria and patients divided into

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two categories based on the type of malignancy as either diagnosed with leukemia/lymphoma or solid tumors and also the duration of therapy.

Assessment of Vitamin D levels

25-OH Vitamin D levels were assessed by immuno-chemilumino-metric assay. Blood samples measuring at least 3 ml were stored in an ice box and sent to the laboratory within 1 h of collection of the sample. 25-OH Vitamin D levels were chosen over 1,25-OH Vitamin D levels since Vitamin D deficiency leads to increase in parathyroid hormone levels, which increase the activity of the enzyme 1 α -hydroxylase that in turn affects 1,25-OH Vitamin D levels.^[18,19] In addition 1,25-OH Vitamin D has a half-life of only 4 h as opposed to 25-OH Vitamin D with a half-life of 2-3 weeks.^[18,19] The cut-off value to define Vitamin D insufficiency was <30 ng/ml as decided by previous studies.^[16,20]

Statistical analysis

The main objective of our study was to establish the prevalence of Vitamin D insufficiency in children with cancer. The collected data were analyzed with SPSS for Windows, Version 16.0, SPSS Inc., Chicago, IL, USA. Percentage analysis was used for categorical variables and for continuous variables the mean and standard deviation was used. To find the significant difference between the bivariate samples in independent groups the unpaired *t*-test was used and to find the significance in categorical data Chi-square test was used. In all the above statistical tools 0.05 was set as the probability value below which results were considered significant.

RESULTS

The Vitamin D levels of 51 patients with malignancy and 51 controls were collected using the method described above. The different characteristics on which the patients were segregated have been described in Table 1. The median age of our patient group was 9.2 years. The majority of patients had either leukemia or lymphoma, but patients with solid tumors were also assessed. All of our patients were on therapy at the time of measurement but were segregated into two groups based on the duration of therapy as less than or more than 1 year of therapy.

Vitamin D insufficiency in children with cancer

In our study, we observed that the prevalence of subjects with Vitamin D insufficiency was 80.39% whereas the prevalence of Vitamin D insufficiency in healthy controls was 50.98%. When segregated into two groups based on age it was found that the prevalence in children <6 years of age was 73% and in children above 6 years of age was 88%.

When segregated into groups based on level of Vitamin D deficiency, 5.8% of subjects were considered severely

deficient, i.e., having 25-OH D <10 ng/ml, 45.1% of subjects were considered deficient, i.e., having 25-OH D >10 but <20 ng/ml, 29.4% of subjects were considered insufficient, i.e., having 25-OH D >20 but <30 ng/ml and 19.6% of subjects had adequate levels of Vitamin D, i.e., >30 ng/ml. It was also found that the prevalence was greater in girls at 90% as compared to boys with 74%. The prevalence of leukemia/lymphoma cases was 82% and in children with solid tumors was 75%. Patients were also divided into two groups based on the duration of therapy at the time of measurement as either receiving therapy for less than a year or more than 1. This is shown in Table 1.

The comparison of Vitamin D levels in healthy subjects versus cancer patients is shown in Figure 1 and the distribution of Vitamin D levels among cases is shown in Table 2.

DISCUSSION

The problem of Vitamin D insufficiency and deficiency is widespread in India despite being closer to the

Table 1: Characteristics of the study population by 25-hydroxyvitamin D levels (n = 51)

Characteristics	Total n (%)	<30 ng/ml n (%)	>30 ng/ml n (%)	P
Age (years)				
<6	26 (51)	20 (77)	6 (23)	0.038
>6	25 (49)	22 (88)	3 (12)	
Sex				
Female	20 (39)	18 (90)	2 (10)	0.458
Male	31 (61)	23 (74)	8 (26)	
Type of cancer				
Leukemia/lymphoma	39 (76)	32 (82)	7 (18)	0.025
Solid tumors	12 (24)	9 (75)	3 (25)	
Duration of therapy at the time of sample collection				
<1 year	11 (22)	7 (64)	4 (36)	0.31
>1 year	40 (78)	34 (85)	6 (15)	

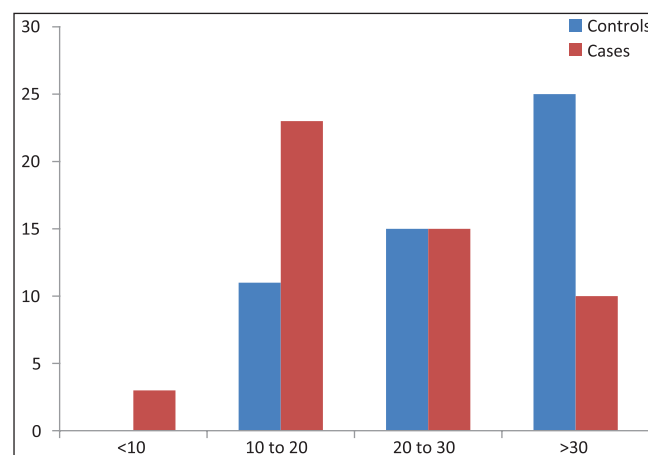


Figure 1: Comparison of Vitamin D levels in healthy subjects versus cancer patients (values in ng/ml) (original)

Table 2: Distribution of 25-hydroxyvitamin D levels in the study population (n = 51)

25-OH D levels in ng/ml	n (%)
>30	3 (5.8)
20-29	23 (45.1)
10-19	15 (29.4)
<10	10 (19.6)

25-OH D – 25-hydroxyvitamin D

equator and the general population having adequate exposure to sunlight. This is perhaps due to a higher percentage of dark skinned people who have an insufficient penetration of ultraviolet radiation needed for Vitamin D synthesis.^[20,21]

Pregnant and lactating women have been found to be Vitamin D deficient despite proper supplementation with 73% of women and 80% of children (at birth) being Vitamin D deficient.^[22-24]

Vitamin D is an essential component of bone and mineral metabolism. Vitamin D is obtained from three sources, sunlight, diet, and dietary supplements.^[9,10,18,19] Solar ultraviolet B radiation of wavelength 290-315 nm penetrates skin and converts 7-dehydrocholesterol to pre-Vitamin D₃, which is rapidly converted to Vitamin D₃.^[9] Asian children require 3 times the recommended amount of sunlight exposure to maintain the Vitamin D levels in view of their dark skin color.^[20,21] The major dietary sources of Vitamin D are eggs, oily fish, dairy products, and meat. Fortified foods are also an important source of Vitamin D.^[25] Fortification of milk has been found to be an effective, safe, and acceptable method,^[26] however, in a setting like India, where the per capita milk consumption is very low fortifying milk might not be useful but can be done in salt or cereals. A need for a national food fortification program for Vitamin D in India has been highlighted in a review by Babu and Calvo.^[27] It is postulated that pediatric oncology patients are at a risk of developing Vitamin D deficiency in their treatment course due to poor nutrition, lack of sun exposure, and owing to treatment too.^[11,14]

Since adequate sunlight exposure at solar noon is difficult to achieve because cancer children spend most of their times indoor, routine oral supplementation may help in preventing Vitamin D deficiency in this group.

Serum levels of Vitamin D are directly related to bone mineral density with optimal density achieved at 40 ng/ml or more^[28] Vitamin D deficiency causes growth retardation and skeletal deformities in children and osteomalacia and osteoporosis in adults. Vitamin D₃ is converted by 2 hydroxylation steps to the active form. In the liver, Vitamin D₃ is converted to 25(OH) D, the major circulating form and in the kidney, 25(OH) D is

converted to 1,25-dihydroxyvitamin D (1,25[OH]2D), the biologically active metabolite. Further, 1,25(OH)2D acts by binding to the nuclear Vitamin D receptor (VDR) within cells. This VDR is widely expressed throughout the body. The VDR is found in the endocrine glands such as pituitary, pancreas, parathyroid, gonads, and placenta and in cardiovascular tissues such as endothelial cells, vascular smooth muscle cells, and cardiomyocytes also.^[7,29,30] The VDR has also been found in hemato-lymphopoietic cells and also has been shown to regulate cell differentiation and the production of interleukins and cytokines.^[21,31] The wide variety of nonskeletal diseases in which Vitamin D plays a role include cardiovascular diseases,^[7] obesity,^[32] metabolic syndrome,^[5] insulin resistance,^[4] infection,^[33] allergy,^[1,2] some forms of cancers,^[6,34,35] and autoimmune disease.^[1]

It has been found that 1,25(OH)2D decreases cellular proliferation of both normal cells and cancer cells and induces their terminal differentiation.^[9,10,36] People living at higher latitudes are at an increased risk of Hodgkin's lymphoma as well as colon, pancreatic, prostate, ovarian, breast, and other cancers and are more likely to die from these cancers, as compared to people living at lower latitudes.^[6,34,35]

Both prospective and retrospective epidemiologic studies indicate that levels of 25-OH D below 20 ng/ml are associated with a 30-50% increased risk of prostate, colon, and breast cancer, and melanoma along with increased mortality from these cancers.^[35-37]

The likely explanation is that colon, prostate, breast, and other tissues express 25-OH D -1 α -hydroxylase and produce 1,25(OH)2D locally to control genes that help to prevent cancer by keeping cellular proliferation and differentiation in check.^[9,10,38] Children and young adults who are exposed to the most sunlight have a 40% reduced risk of non-Hodgkin's lymphoma^[39] and a reduced risk of death from malignant melanoma, as compared to those who have the least exposure to sunlight.^[37] It has been postulated that if a cell becomes malignant, 1,25(OH)2D can induce apoptosis and prevent angiogenesis, thereby reducing the potential for the malignant cell to survive.^[10] After 1,25(OH)2D completes these tasks, it initiates its own destruction by stimulating the *CYP24* gene to produce the inactive calcitric acid. This guarantees that 1,25(OH)2D does not enter the circulation to influence calcium metabolism.^[9,10]

This is a possible explanation for why increased sun exposure and higher circulating levels of 25-OH D are associated with a decreased risk of cancers.^[21,38,39]

25(OH)D is the major circulating form of Vitamin D with a half-life of 2-3 weeks, which is the best available indicator

of Vitamin D status. Although 1,25(OH)D (calcitriol) is the active form, it has a half-life of only 4 h which is not a good indicator of Vitamin D stores as Vitamin D deficiency can cause PTH elevation that induces increased 1 α -hydroxylase activity that results in normal or increased levels of 1,25(OH)D. It also circulates at a concentration that is 100-1000-fold <25(OH)D.^[40]

Data on the prevalence of Vitamin D insufficiency in children are sparse or incomplete in most countries. However, it is an important public health problem in both developed and developing countries, with an estimated prevalence of 29-100% in children and adolescents.^[9,41]

There are only a few reports so far that have studied the prevalence of 25-OH D status in cancer survivors.^[11-13] A prospective cross-sectional study done in the United Kingdom by Sinha *et al.*^[13] comparing 61 cancer survivors aged between 1 and 18 years with 60 controls, found a higher prevalence of 25-OH D deficiency (<25 ng/ml) among survivors compared to controls, (21.3% in the cancer survivors vs. 3.3% in the controls). The risk factors, which predicted 25-OH D deficiency included seasonality, ethnicity, older age, and a cancer diagnosis.^[13]

Simmons *et al.* studied 78 leukemia survivors, who either received chemotherapy or hematopoietic stem cell transplant and found a prevalence of 25-OH D deficiency (<15 ng/ml) of 35%. The risk factors found were decreased exposure to sunlight, older age, and lack of 25-OH D supplementation.^[12]

Robien *et al.* in their study in Minnesota looked at 95 cancer survivors aging between 2.7 and 72 years with a diagnosis of leukemia or lymphoma and found the prevalence of 25-OH D deficiency (<50 ng/ml) to be 10%. Risk factors for 25-OH D deficiency included lack of 25-OH D supplementation and use of steroids.^[42]

Even in an Indian context, Vitamin D deficiency has been reported to be present in the majority of children in spite of adequate exposure to sunlight. The prevalence of Vitamin D deficiency is 50-90% in the Indian subcontinent and is attributed to low dietary calcium along with skin color and changing lifestyle.^[43,44]

The USA Endocrine Society Guideline recommends screening for Vitamin D deficiency in any population at risk. This includes patients with dark skin who live at a higher altitude and infants born to Vitamin D deficient mothers, infants with poor growth, chronic kidney disease, hepatic failure, malabsorption syndromes, suspected rickets, osteoporosis, inflammatory bowel disease, hyperparathyroidism; those on drug therapy such as anticonvulsants, glucocorticoids, antiretroviral therapy, antifungals, obese children; granuloma-forming disorders

such as sarcoidosis, tuberculosis, and histoplasmosis.^[45] Currently, they do not recommend cancer survivors or children on chemotherapy for screening, which needs to be considered and revised.

Recommended daily intake of Vitamin D for children, according to the Institute of Medicine is 200 IU a day but experts agree that in the absence of necessary exposure to sunlight, this amount may be increased to 800-1000 IU a day.^[46]

The prevalence of Vitamin D insufficiency in our study group (80.39%) was found to be higher than in results obtained from studies done before such as the one done by Sinha *et al.* where the prevalence of Vitamin D levels below 50 ng/ml in children with cancer was 62%.^[13] Despite the study being conducted in a region of higher latitude, the increased prevalence in our subjects may be due to poorer nutritional habits and skin color.

We found out that there was a statistically significant ($P = 0.038$) increase in the prevalence of Vitamin D insufficiency in children above 6 years (88%) as compared to children below 6 years (77%) that has already been established in healthy populations.^[15-17] It was also noted that the frequency of severe deficiency was higher in our group when compared to studies done on healthy populations.^[15-17,47] This is hardly surprising as cancer patients are exposed to less sunlight since they are discouraged from playing outside for sanitary reasons have a poorer nutritional status and are on therapy that is potentially bone weakening.

The prevalence was greater in girls at 90% as compared to boys with 74% but was found to be statistically insignificant, perhaps due to the small size of our sample.

The prevalence of leukemia/lymphoma cases (82%) was significantly ($P = 0.025$) higher than in children with solid tumors (75%) that went against the study conducted by Helou *et al.* where prevalence was higher in children suffering from solid tumors,^[48] perhaps due to the more intense chemotherapy that these patients receive and greater use of steroids in therapy.

When segregated based on the duration of therapy received there was no significant difference between children receiving therapy for less than a year compared to those receiving therapy for greater than a year and this may suggest that Vitamin D insufficiency may already set in during the beginning stages of therapy.

The limitations of our study were that we had a small sample size, pretreatment and posttreatment comparison of Vitamin D levels was not done which would give a more accurate analysis of effect of treatment on Vitamin D levels in cancer patients and the samples were all collected

in a span of 6 months, thereby failing to note the effect of seasonal variation on Vitamin D levels.

CONCLUSION

We have only just begun to understand the role of Vitamin D in homeostatic pathways. Where once it was thought rickets and osteomalacia were the apex of Vitamin D deficiency, we now realize that it is just one of many manifestations of Vitamin D deficiency/insufficiency. Our study shows evidence of increased incidence of Vitamin D insufficiency in children with cancer and as such suggest Vitamin D assays and subsequent supplementation be made a part of the initial tests run at diagnosis in children with cancer. It has also been proved conclusively that patients with certain types of cancer have an increased frequency of VDR genotype variants.^[29,30] A future area where the role of Vitamin D may be further studied is in determining VDR polymorphisms in patients with cancer and its role in determining Vitamin D levels and disease progression.

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Conflicts of interest

There are no conflicts of interest.

REFERENCES

- Krieger MA, Manson JE, Costenbader KH. Does Vitamin D affect risk of developing autoimmune disease? A systematic review. *Semin Arthritis Rheum* 2011;40:512-31.e8.
- Camargo CA Jr., Rifas-Shiman SL, Litonjua AA, Rich-Edwards JW, Weiss ST, Gold DR, *et al.* Maternal intake of Vitamin D during pregnancy and risk of recurrent wheeze in children at 3 y of age. *Am J Clin Nutr* 2007;85:788-95.
- Brehm JM, Acosta-Pérez E, Klei L, Roeder K, Barmada M, Boutaoui N, *et al.* Vitamin D insufficiency and severe asthma exacerbations in Puerto Rican children. *Am J Respir Crit Care Med* 2012;186:140-6.
- Chiu KC, Chu A, Go VL, Saad MF. Hypovitaminosis D is associated with insulin resistance and beta cell dysfunction. *Am J Clin Nutr* 2004;79:820-5.
- Ganji V, Zhang X, Shaikh N, Tangpricha V. Serum 25-hydroxyvitamin D concentrations are associated with prevalence of metabolic syndrome and various cardiometabolic risk factors in US children and adolescents based on assay-adjusted serum 25-hydroxyvitamin D data from NHANES 2001-2006. *Am J Clin Nutr* 2011;94:225-33.
- Holick MF. Vitamin D: Importance in the prevention of cancers, type 1 diabetes, heart disease, and osteoporosis. *Am J Clin Nutr* 2004;79:362-71.
- Temmerman JC. Vitamin D and cardiovascular disease. *J Am Coll Nutr* 2011;30:167-70.
- El-Hajj Fuleihan G, Nabulsi M, Tamim H, Maalouf J, Salamoun M, Khalife H, *et al.* Effect of Vitamin D replacement on musculoskeletal parameters in school children: A randomized controlled trial. *J Clin Endocrinol Metab* 2006;91:405-12.
- Holick MF. Resurrection of Vitamin D deficiency and rickets. *J Clin Invest* 2006;116:2062-72.
- Holick MF, Garabedian M. Vitamin D: Photobiology, metabolism, mechanism of action, and clinical applications. In: Favus MJ, editor. *Primer on the Metabolic Bone Diseases and Disorders of Mineral Metabolism*. 6th ed. Washington, DC: American Society for Bone and Mineral Research; 2006. p. 129-37.
- Bilariki K, Anagnostou E, Masse V, Elie C, Grill J, Valteau-Couanet D, *et al.* Low bone mineral density and high incidences of fractures and Vitamin D deficiency in 52 pediatric cancer survivors. *Horm Res Paediatr* 2010;74:319-27.
- Simmons JH, Chow EJ, Koehler E, Esbenshade A, Smith LA, Sanders J, *et al.* Significant 25-hydroxyvitamin D deficiency in child and adolescent survivors of acute lymphoblastic leukemia: Treatment with chemotherapy compared with allogeneic stem cell transplant. *Pediatr Blood Cancer* 2011;56:1114-9.
- Sinha A, Avery P, Turner S, Bailey S, Cheetham T. Vitamin D status in paediatric patients with cancer. *Pediatr Blood Cancer* 2011;57:594-8.
- Haddy TB, Mosher RB, Reaman GH. Late effects in long-term survivors after treatment for childhood acute leukemia. *Clin Pediatr (Phila)* 2009;48:601-8.
- Huh SY, Gordon CM. Vitamin D deficiency in children and adolescents: Epidemiology, impact and treatment. *Rev Endocr Metab Disord* 2008;9:161-70.
- El-Hajj Fuleihan G, Nabulsi M, Choucair M, Salamoun M, Hajj Shahine C, Kizirian A, *et al.* Hypovitaminosis D in healthy schoolchildren. *Pediatrics* 2001;107:E53.
- Gordon CM, DePeter KC, Feldman HA, Grace E, Emans SJ. Prevalence of Vitamin D deficiency among healthy adolescents. *Arch Pediatr Adolesc Med* 2004;158:531-7.
- DeLuca HF. Overview of general physiologic features and functions of Vitamin D. *Am J Clin Nutr* 2004;80 (Suppl):1689S-96S.
- Bouillon R. Vitamin D: From photo-synthesis, metabolism, and action to clinical applications. In: DeGroot LJ, Jameson JL, editors. *Endocrinology*. Philadelphia: WB Saunders; 2001. p. 1009-28.
- Saintonge S, Bang H, Gerber LM. Implications of a new definition of Vitamin D deficiency in a multiracial us adolescent population: The national health and nutrition examination survey III. *Pediatrics* 2009;123:797-803.
- Holick MF. Vitamin D deficiency. *N Engl J Med* 2007;357:266-81.
- Hollis BW, Wagner CL. Assessment of dietary Vitamin D requirements during pregnancy and lactation. *Am J Clin Nutr* 2004;79:717-26.
- Lee JM, Smith JR, Philipp BL, Chen TC, Mathieu J, Holick MF. Vitamin D deficiency in a healthy group of mothers and newborn infants. *Clin Pediatr (Phila)* 2007;46:42-4.
- Bodnar LM, Simhan HN, Powers RW, Frank MP, Cooperstein E, Roberts JM. High prevalence of Vitamin D insufficiency in black and white pregnant women residing in the Northern United States and their neonates. *J Nutr* 2007;137:447-52.
- Lamberg-Allardt C. Vitamin D in foods and as supplements. *Prog Biophys Mol Biol* 2006;92:33-8.
- Natri AM, Salo P, Vikstedt T, Palssa A, Huttunen M, Kärkkäinen MU, *et al.* Bread fortified with cholecalciferol increases the serum 25-hydroxyvitamin D concentration in women as effectively as a cholecalciferol supplement. *J Nutr* 2006;136:123-7.
- Babu US, Calvo MS. Modern India and the Vitamin D dilemma: Evidence for the need of a national food fortification program. *Mol Nutr Food Res* 2010;54:1134-47.
- Bischoff-Ferrari HA, Giovannucci E, Willett WC, Dietrich T, Dawson-Hughes B. Estimation of optimal serum concentrations of 25-hydroxyvitamin D for multiple health outcomes. *Am J Clin Nutr* 2006;84:18-28.
- Ruza E, Sotillo E, Sierrasesúmaga L, Azcona C, Patiño-García A. Analysis of polymorphisms of the Vitamin D receptor, estrogen receptor, and collagen alpha1 genes and their relationship with height in children with bone cancer. *J Pediatr Hematol Oncol* 2003;25:780-6.

30. te Winkel ML, van Beek RD, de Muinck Keizer-Schrama SM, Uitterlinden AG, Hop WC, Pieters R, *et al.* Pharmacogenetic risk factors for altered bone mineral density and body composition in pediatric acute lymphoblastic leukemia. *Haematologica* 2010;95:752-9.
31. Verstuyf A, Carmeliet G, Bouillon R, Mathieu C. Vitamin D: A pleiotropic hormone. *Kidney Int* 2010;78:140-5.
32. Harel Z, Flanagan P, Forcier M, Harel D. Low Vitamin D status among obese adolescents: Prevalence and response to treatment. *J Adolesc Health* 2011;48:448-52.
33. Madden K, Feldman HA, Smith EM, Gordon CM, Keisling SM, Sullivan RM, *et al.* Vitamin D deficiency in critically ill children. *Pediatrics* 2012;130:421-8.
34. Jenab M, Bueno-de-Mesquita HB, Ferrari P, van Duijnhoven FJ, Norat T, Pischon T, *et al.* Association between pre-diagnostic circulating Vitamin D concentration and risk of colorectal cancer in European populations: A nested case-control study. *BMJ* 2010;340:b5500.
35. Giovannucci E, Liu Y, Rimm EB, Hollis BW, Fuchs CS, Stampfer MJ, *et al.* Prospective study of predictors of Vitamin D status and cancer incidence and mortality in men. *J Natl Cancer Inst* 2006;98:451-9.
36. Garland CF, Garland FC, Gorham ED, Lipkin M, Newmark H, Mohr SB, *et al.* The role of Vitamin D in cancer prevention. *Am J Public Health* 2006;96:252-61.
37. Berwick M, Armstrong BK, Ben-Porat L, Fine J, Krickler A, Eberle C, *et al.* Sun exposure and mortality from melanoma. *J Natl Cancer Inst* 2005;97:195-9.
38. Gorham ED, Garland CF, Garland FC, Grant WB, Mohr SB, Lipkin M, *et al.* Vitamin D and prevention of colorectal cancer. *J Steroid Biochem Mol Biol* 2005;97:179-94.
39. Chang ET, Smedby KE, Hjalgrim H, Porwit-MacDonald A, Roos G, Glimelius B, *et al.* Family history of hematopoietic malignancy and risk of lymphoma. *J Natl Cancer Inst* 2005;97:1466-74.
40. Holick MF. Vitamin D status: Measurement, interpretation, and clinical application. *Ann Epidemiol* 2009;19:73-8.
41. Shin YH, Kim KE, Lee C, Shin HJ, Kang MS, Lee HR, *et al.* High prevalence of Vitamin D insufficiency or deficiency in young adolescents in Korea. *Eur J Pediatr* 2012;171:1475-80.
42. Robien K, Strayer LG, Majhail N, Lazovich D, Baker KS, Smith AR, *et al.* Vitamin D status among long-term survivors of hematopoietic cell transplantation. *Bone Marrow Transplant* 2011;46:1472-9.
43. Balasubramanian S, Dhanalakshmi K, Amperayani S. Vitamin D deficiency in childhood — A review of current guidelines on diagnosis and management. *Indian Pediatr* 2013;50:669-75.
44. Harinarayan CV, Joshi SR. Vitamin D status in India – its implications and remedial measures. *J Assoc Physicians India* 2009;57:40-8.
45. Holick MF, Binkley NC, Bischoff-Ferrari HA, Gordon CM, Hanley DA, Heaney RP, *et al.* Evaluation, treatment, and prevention of Vitamin D deficiency: An Endocrine Society clinical practice guideline. *J Clin Endocrinol Metab* 2011;96:1911-30.
46. Heaney RP, Davies KM, Chen TC, Holick MF, Barger-Lux MJ. Human serum 25-hydroxycholecalciferol response to extended oral dosing with cholecalciferol. *Am J Clin Nutr* 2003;77:204-10.
47. Kumar J, Muntner P, Kaskel FJ, Hailpern SM, Melamed ML. Prevalence and associations of 25-hydroxyvitamin D deficiency in US children: NHANES 2001-2004. *Pediatrics* 2009;124:e362-70.
48. Helou M, Ning Y, Yang S, Irvine P, Bachmann LM, Godder K, *et al.* Vitamin D deficiency in children with cancer. *J Pediatr Hematol Oncol* 2014;36:212-7.

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