Ethnicity and season are major determinants of serum 25-hydroxyvitamin D in New Zealand children

T.J. Green, J.E. Rockell, C.M. Skeaff, S.J. Whiting, R. W. Taylor, S. M. Williams, W.R. Parnell, R. Scragg, N. Wilson, D. Schaaf, E.D. Fitzgerald

Department of Human Nutrition, LINZ Activity and Health Research Unit and Preventive and Social Medicine, University of Otago, Dunedin, New Zealand; College of Pharmacy and Nutrition, University of Saskatchewan, Saskatoon, Saskatchewan, Canada; School of Population Health, Faculty of Medicine and Health Science, University of Auckland, New Zealand; School of Mäori Studies, Massey University, Palmerston North, New Zealand.

Abstract. The aim of this study was to determine 25hydroxyvitamin vitamin D concentrations and their determinants in a national sample of New Zealand children aged 5-14 y. The 2002 National Children's Nutrition Survey was designed to survey New Zealand school aged children, including over-sampling of Mäori and Pacific children to allow ethnic-specific analyses. Serum 25hydroxyvitamin D concentration [mean (99% CI) nmol/L] in Mäori children (n=456) was 43 (38, 49), in Pacific (n=646) 36 (31, 42), and in New Zealand European and Others (NZEO) (n=483) 53 (47, 59). Among Mäori, Pacific and NZEO respectively, prevalence (%, 99% CI) of serum 25-hydroxyvitamin D deficiency (<17.5 nmol/L) was 5% (2, 12), 8% (5, 14), and 3% (1, 7). Prevalence of insufficiency (<37.5 nmol/L) was 41% (30, 54), 60% (43, 74) and 26% (16, 37), respectively. Multiple regression analysis found 25-hydroxyvitamin D concentrations were lower in winter than summer [adjusted mean difference (99% CI) nmol/L; 15 (8, 22)], lower in females than males [5 (1, 10)], and lower in obese than those of 'normal' weight [8 (3,13)]. Relative to NZEO, 25-hydroxyvitamin D concentrations were lower in Mäori [9 (3, 15)] and Pacific children [16 (10, 22)]. Ethnicity and season are major determinants of serum 25-hydroxyvitamin D. There is a high prevalence of vitamin D insufficiency in New Zealand children.

Introduction

Vitamin D plays an essential role in calcium and phosphorus homeostasis. The most serious clinical consequence of vitamin D deficiency in children is rickets, (1). Though still uncommon in western countries, it would appear from the increasing number of case reports being published that rickets is re-emerging as a public health problem (2). Lesser degrees of vitamin D deficiency, referred to as insufficiency, have been associated with lower bone mineral density (3) and bone accretion rates in children (4), as well as elevated serum parathyroid hormone concentrations (3); these effects are consistent with secondary hyperparathyroidism. Recent discoveries indicate that vitamin D has functions unrelated to calcium, specifically in cell differentiation and in the immune system. These biological effects add plausibility to reports that low vitamin D status is associated with increased risk of childhood-onset Type 1 diabetes and increased risk of some types of cancer in adult. We present the serum 25hydroxyvitamin D results from the Children's Nutrition Survey (CNS02), a large national survey of New Zealand school-aged children in 2002. New Zealand lies geographically from ~35 °S to ~47 °S and has a food supply with minimal vitamin D fortification. Furthermore, there are three main ethnic groups with varying skin colour: Mäori, Pacific, and New Zealand European. Moreover, the survey provides a unique opportunity to explore the independent effects of season, age, ethnicity, and obesity on serum 25-hydroxyvitamin D concentrations in children 5-14 y.

Subjects and Methods

The 2002 National Children's Nutrition Survey (CNS02) was a cross-sectional survey of a national sample of New Zealand school children and adolescents aged 5 to 14 v. conducted during the 2002 school year. A school based sampling frame of children was used with over-sampling of Mäori and Pacific children to allow for ethnic specific analysis. The CNS02 aimed to recruit 3000 children with 1000 from each of three ethnic groups: Mäori, Pacific, and New Zealand Europeans and Others (NZEO). Children from selected schools were assigned to one of three ethnic groups with a different probability of selection for each ethnic group. Details of the survey methodology are described more fully elsewhere (5). The total number of children invited to participate was 4728, 3275 participated and 1927 provided blood samples. Blood was available for vitamin D analysis for 1659 participants. Of these 1585 children had all data available that was relevant to this study; an overall response rate of 33.5%. Blood sample collection was by trained phlebotomists. The children were given no instructions to fast. Serum 25-hydroxyvitamin D surplus determined on blood using radioimmunoassay (RIA) kit (DiaSorin Stillwater, MN). Statistical analyses were carried out using STATA 8.0, adjusting for the complex survey design. Because age, sex, ethnicity, season, geographical location and obesity have been reported to affect serum 25-hydroxyvitamin D concentrations we used multiple linear regression models to examine the independent relationships between each of these variables and serum 25-hydroxyvitamin D. We estimated adjusted means based on these models. Results were considered significant when p<0.01. April through September were defined as 'winter' months, and October to December, and March, defined as 'summer' months.

Results

Mean serum 25-hydroxyvitamin D concentrations as well as prevalence rates of vitamin D deficiency (<17.5 nmol/L) and insufficiency (<37.5 nmol/L) by age, sex and ethnicity are presented in **Table 1**. New Zealand children had a mean concentration of 50 nmol/L, with mean

 $\begin{tabular}{ll} \textbf{Table 1} \\ \textbf{Serum 25-hydroxyvitamin D concentrations} & \textbf{and prevalence of deficiency and insufficiency} & \\ \hline \\ \textbf{in New Zealand children by age, sex and ethnicity} & \\ \hline \\ \hline \\ \end{matrix}^2 .$

		Serum 25-hydroxyvitamin D (nmol/L)			
Sex, age and ethnicity	n	Mean (99% CI)	%<17.5nmol/L	% <37.5 nmol/L	
			(99% CI)	(99% CI)	
New Zealand children					
All	1585	50 (45, 54)	4(2, 6)	32 (23, 42)	
Male	801	52 (47, 58)	3 (2, 6)	27 (18, 38)	
5-6	157	57 (49, 65)	1(0,4)	19 (10, 34)	
7-10	360	53 (48, 58)	2(1,6)	24 (16, 34)	
11-14	284	50 (41, 59)	5 (2, 10)	33 (18, 52)	
Female	784	47 (41, 53)	4(2,9)	36 (25, 48)	
5-6	137	48 (43, 54)	1(1,4)	29 (17, 44)	
7-10	362	51 (44, 58)	3 (1, 6)	31 (21, 45)	
11-14	285	42 (33, 51)	6 (3, 16)	43 (25, 63)	
M ori					
All	456	43 (38, 49)	5 (2, 12)	41 (30, 54)	
Male	232	47 (40, 53)	5 (2, 12)	37 (24, 52)	
5-6	52	48 (41, 55)	2(0,7)	26 (11, 50)	
7-10	116	47 (40, 54)	3 (1, 11)	38 (22, 56)	
11-14	64	45 (34, 57)	7 (3, 20)	40 (21, 63)	
Female	224	40 (35, 45)	6 (2, 16)	46 (33, 59)	
5-6	50	42 (36, 48)	2(1,8)	45 (26, 65)	
7-10	116	46 (40, 52)	4(1, 12)	33 (19, 50)	
11-14	58	35 (28, 42)	10 (3, 28)	58 (36, 77)	
Pacific					
All	646	36 (31, 42)	8 (5, 14)	60 (43, 74)	
Male	297	38 (33, 44)	7 (3, 15)	54 (36, 72)	
5-6	64	42 (34, 49)	3 (1, 10)	49 (29, 69)	
7-10	129	40 (34, 46)	5 (2, 15)	50 (30, 70)	
11-14	104	36 (30, 42)	12 (6, 23)	61 (40, 78)	
Female	349	34 (29, 40)	9 (5, 18)	46 (28, 64)	
5-6	47	39 (31, 47)	3 (1, 10)	47 (28, 67)	
7-10	145	34 (28, 40)	7 (3, 16)	66 (48, 81)	
11-14	157	32 (25, 39)	15 (7, 33)	71 (52, 85)	
NZEO					
All	483	53 (47, 59)	3(1,7)	26 (16, 37)	
Male	272	56 (49, 63)	2(1,6)	21 (11, 36)	
5-6	41	62 (52, 72)	1 (0, 3)	13 (5, 33)	
7-10	115	56 (50, 63)	1 (0, 5)	17 (10, 28)	
11-14	116	53 (41, 65)	3 (1, 10)	28 (11, 54)	
Female	211	50 (43, 57)	3 (1, 9)	30 (18, 45)	
5-6	40	52 (45, 59)	1(0,3)	20 (8, 43)	
7-10	101	54 (45, 64)	2(1,6)	27 (14, 44)	
11-14	70	45 (34, 56)	4 (1, 15)	36 (17, 60)	

¹ Insufficiency <17.5 nmol/L; deficiency <37.5 nmol/L.

concentrations in subgroups ranging from 32 nmol/L in Pacific girls aged 11 to 14 y, to 62 nmol/L in NZEO boys aged 5 to 6 y. Four percent of New Zealand children were vitamin D deficient; nearly one third were insufficient. Prevalence of vitamin D deficiency ranged from 1% in NZEO boys and girls 5 to 6 v, to 15% in Pacific girls 11 to 14 y. Similarly, the prevalence of vitamin D insufficiency ranged from 13% in NZEO boys 5 to 6 y to more than 70% in Pacific girls 11 to 14 y. The independent effects of age, sex, ethnicity, latitude (North cf South Island), season ('summer' cf 'winter' months) and overweight/obesity on serum 25-hydroxyvitamin D concentrations are presented as adjusted means and prevalence in Table 2. Boys had higher 25-hydroxyvitamin D concentrations than females by [adjusted mean difference and 99% CI] 5 (1, 10) nmol/L. The effect of age category on serum 25hydroxyvitamin D was not significant. Ethnicity was a strong determinant of 25-hydroxyvitamin D concentration; Mäori children were 9 (3, 15) nmol/L and Pacific children 16 (10, 22) nmol/L lower than NZEO children. Obese children had an 8(3,13) nmol/L lower serum 25hydroxyvitamin D concentration than normal weight children. There was a marked independent effect of season on mean 25-hydroxyvitamin D concentrations, with a difference of 15 (8, 22) nmol/L between 'winter' and 'summer' (Table 4). Mean 25-hydroxyvitamin D concentrations adjusted for age, sex, ethnicity, latitude, and

TABLE 2
Adjusted mean serum 25-hydroxyvitamin D concentrations and prevalence of deficiency and insufficiency¹ in New Zealand children²³

		Serum 25-hydroxy	Adjusted Prevalence <17.5 nmol/L <37.5 nmol/L	
	_	vitamin D (nmol/L)		
	n	Adjusted Mean (99% CI)	% (99% CI)	% (99% CI)
Sex				
Male	801	52 (48, 56) ^a	2(1,4)	24 (17, 33)
Female	784	47 (43, 51) ^b	3 (1, 6)	33 (24, 43)
Age (yr)				
5-6	294	53 (47, 58)	1 (0, 3) ^a	21 (13, 32)
7-10	722	51 (47, 55)	2 (1, 5)	25 (18, 34)
11-14	569	47 (41, 52)	4 (2, 10) ^b	35 (23, 49)
Ethnicity				
M ori	456	44 (39, 48) ^a	4(1, 11)	38 (26, 52) ^a
Pacific People	646	37 (34, 40) ^b	7 (3, 13)	59 (49, 68) ^b
NZEO	483	53 (48, 57) ^c	2 (1, 4)	23 (16, 32) ^c
Region				
South Island	174	48 (44, 53)	2 (0, 8)	27 (15, 45)
North Island	1411	50 (45, 54)	3 (1, 5)	29 (21, 38)
Season				
Winter (April-Sept)	924	43 (39, 47) ^a	4(2,7)	42 (32, 53) ^a
Summer (Oct-Dec, Mar)	661	58 (53, 64) ^b	1 (0, 4)	14 (8, 23) ^b
Obesity ⁴				
Obesity	188	42 (37, 47) ^a	1 (1, 4)	46 (29, 64) ^a
Overweight	332	50 (45, 55) ^b	2(1,4)	28 (18, 41)
Normal Weight	1065	50 (46, 54) ^b	3 (1, 6)	27 (20, 35) ^b

Insufficiency <17.5nmol/L; deficiency <37.5nmol/L.

overweight/obesity, fell from a peak in March of 68 (57, 79) nmol/L to a nadir in August of 36 (32, 40) nmol/L

Conclusions

We report a high prevalence of insufficient vitamin D status in New Zealand school-age children; from 32% for all children to a high of 60% in Pacific children.

References

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All data adjusted for sample weighting.

Data rounded to whole numbers

² All data adjusted for sample weighting.

³ Adjusted for age, sex, ethnicity, season, region and obesity.

⁴ Normal, overweight or obese classification according to Cole 2000²⁴ Numbers not sharing a common superscript are significantly different, p<0.01 Data rounded to whole numbers.