

Influence of vitamin D supplementation on growth, body composition, pubertal development and spirometry in South African schoolchildren: a randomised controlled trial (ViDiKids)

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ABSTRACT

Objective To determine whether weekly oral vitamin D supplementation influences growth, body composition, pubertal development or spirometric outcomes in South African schoolchildren.

Design Phase 3 double-blind randomised placebo-controlled trial.

Setting Socioeconomically disadvantaged peri-urban district of Cape Town, South Africa.

Participants 1682 children of black African ancestry attending government primary schools and aged 6–11 years at baseline.

Interventions Oral vitamin D₃ (10 000 IU/week) versus placebo for 3 years.

Main outcome measures Height-for-age and body mass index-for-age, measured in all participants; Tanner scores for pubertal development, spirometric lung volumes and body composition, measured in a subset of 450 children who additionally took part in a nested substudy.

Results Mean serum 25-hydroxyvitamin D₃ concentration at 3-year follow-up was higher among children randomised to receive vitamin D versus placebo (104.3 vs 64.7 nmol/L, respectively; mean difference (MD) 39.7 nmol/L, 95% CI 37.6 to 41.9 nmol/L). No statistically significant differences in height-for-age z-score (adjusted MD (aMD) –0.08, 95% CI –0.19 to 0.03) or body mass index-for-age z-score (aMD –0.04, 95% CI –0.16 to 0.07) were seen between vitamin D versus placebo groups at follow-up. Among substudy participants, allocation to vitamin D versus placebo did not influence pubertal development scores, % predicted forced expiratory volume in 1 s (FEV1), % predicted forced vital capacity (FVC), % predicted FEV1/FVC, fat mass or fat-free mass.

Conclusions Weekly oral administration of 10 000 IU vitamin D₃ boosted vitamin D status but did not influence growth, body composition, pubertal development or spirometric outcomes in South African schoolchildren.

Trial registration numbers ClinicalTrials.gov NCT02880982, South African National Clinical Trials Register DOH-27-0916-5527.

WHAT IS ALREADY KNOWN ON THIS TOPIC

- ⇒ Observational studies have reported independent associations between vitamin D deficiency in childhood and slower linear growth, reduced lean mass, obesity and precocious puberty.
- ⇒ A phase 2 clinical trial conducted in Mongolia reported that a 6-month course of vitamin D supplementation increased height gain in 113 vitamin D deficient schoolchildren aged 12–15 years; however, these results were not confirmed by a recent phase 3 trial conducted in the same setting.
- ⇒ Randomised controlled trials to determine the effects of vitamin D supplementation on growth and development in schoolchildren have not been conducted in other settings.

WHAT THIS STUDY ADDS

- ⇒ This placebo-controlled phase 3 clinical trial, conducted in 1682 black African schoolchildren in Cape Town, South Africa, showed that a 3-year course of weekly vitamin D supplementation was effective in elevating circulating 25-hydroxyvitamin D concentrations.
- ⇒ However, this was not associated with any effect on linear growth, body composition, pubertal development or spirometric lung volumes.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

- ⇒ Our findings do not support the use of vitamin D supplementation as an intervention to influence child growth, body composition, pubertal development or spirometric lung volumes.

INTRODUCTION

Middle childhood and early adolescence represent key periods for growth and development that have an important influence



on stature and health outcomes in later adolescence and adulthood.¹ Stunting in childhood has long been recognised to associate with multiple adverse long-term health outcomes,² and is particularly common in lower-income countries.³ These settings have also witnessed an emerging epidemic of childhood obesity,⁴ which has in turn been associated with accelerated pubertal development.⁵ Interventions to alleviate these public health challenges are urgently needed.

Vitamin D is a fat-soluble micronutrient with pleiotropic effects on human health.⁶ Observational studies have reported that vitamin D deficiency, as evidenced by low circulating concentrations of its major metabolite 25-hydroxyvitamin D (25(OH)D), associates with slower linear growth,⁷ reduced lean mass,⁸ childhood obesity^{9–11} and precocious puberty,^{12–13} potentially reflecting the ability of vitamin D to stimulate production of insulin-like growth factor 1¹⁴ and regulate adipogenesis.¹⁵ A phase 2 randomised controlled trial (RCT) conducted in Mongolia reported that a 6-month course of vitamin D supplementation increased height gain in 113 vitamin D deficient schoolchildren aged 12–15 years,¹⁶ but these results were not confirmed by a recent phase 3 RCT conducted in the same setting.¹⁷ RCTs to determine the effects of vitamin D supplementation on growth and development in schoolchildren have not been conducted in other settings representing children at different risks of malnutrition, stunting and obesity. An opportunity to conduct such an investigation recently arose as part of the ViDiKids trial, a multicentre phase 3 RCT that investigated the effects of weekly oral administration of 10 000 IU vitamin D₃ for 3 years on the primary outcome of tuberculosis infection in a cohort of 1682 schoolchildren aged 6–11 years living in a socioeconomically disadvantaged peri-urban district of Cape Town, South Africa.¹⁸ Height-for-age z-scores and body mass index (BMI)-for-age z-scores were assessed in all participants (n=1682), and Tanner scores for pubertal development, spirometric lung volumes and body composition were assessed in a subset of 450 children who also took part in a nested substudy.

METHODS

Trial design, setting and sponsorship

We conducted a multicentre phase 3 double-blind individually randomised placebo-controlled trial of weekly oral vitamin D supplementation in 23 government schools in Cape Town, South Africa, as previously described.^{18–19} The primary outcome was the acquisition of tuberculosis infection, as evidenced by the conversion of a QuantiFERON-TB Gold Plus (QFT-Plus) assay result from negative at baseline to positive at 3-year follow-up. The current manuscript reports the effects of the intervention on prespecified secondary outcomes relating to growth in all study participants, and body composition, pubertal development and spirometry in a subset of participants who

additionally took part in a nested substudy. The trial was sponsored by Queen Mary University of London.

Participants

Inclusion criteria for the main trial were enrolment in Grades 1–4 at a participating school; age 6–11 years at screening; and written informed assent/consent to participate in the main trial provided by children and their parent/legal guardian, respectively. Exclusion criteria for the main trial were a history of previous tuberculosis infection, active tuberculosis disease or any chronic illness other than asthma (including known or suspected HIV infection) prior to enrolment; use of any regular medication other than asthma medication; use of vitamin D supplements at a dose of more than 400 IU/day in the month before enrolment; plans to move away from study area within 3 years of enrolment; inability to swallow a placebo soft gel capsule with ease; and clinical evidence of rickets or a positive QFT-Plus assay result at screening. An additional inclusion criterion for the substudy was enrolment in Grade 4 at a participating school (ie, only children in Grade 4 were eligible for the substudy).

Enrolment

Full details of enrolment procedures are described in Supplementary Methods (online supplemental material). Eligible participants underwent measurement of weight using a Digital Floor Scale (Charder Medical, Taichung City, Taiwan), height using a portable stadiometer (HM200P, Charder Medical) and waist circumference using a measuring tape. Substudy participants also underwent spirometry according to European Respiratory Society and American Thoracic Society standards²⁰ using a portable spirometer (Carefusion, San Diego, California, USA) and measurement of body composition by dual-energy X-ray absorptiometry (DXA) as described below.

Randomisation and blinding

Full details of randomisation and blinding procedures have been described previously,^{18–19} and are presented in Supplementary Methods (online supplemental material). Briefly, eligible and assenting children whose parents consented to their participation in the trial were individually randomised to receive a weekly capsule containing vitamin D₃ or placebo for 3 years, with a one-to-one allocation ratio and randomisation stratified by the school of attendance. Treatment allocation was concealed from participants, care providers and all trial staff (including senior investigators and those assessing outcomes) until completion of the trial to maintain the double-blind.

Intervention

Study medication comprised a 3-year course of weekly soft gel capsules manufactured by the Tishcon Corporation (Westbury, New York, USA), containing either 0.25 mg (10 000 international units) cholecalciferol (vitamin D₃) in olive oil (intervention arm) or olive oil without any vitamin D₃ content (placebo arm). A weekly dose of

10000 IU (equivalent to 1429 IU/day) was selected in preference to a daily dose of 6000 IU (the Recommended Daily Allowance for this age group) because we were concerned that the latter dose would be inadequate to maintain serum 25(OH)D concentrations >50 nmol/L,²¹ and we felt that adherence to a directly-observed weekly dose would be superior to daily self-administration.²² Weekly supplementation has also been shown to be effective in elevating 25(OH)D concentrations into the physiological range in children by other investigators.²³ Active and placebo capsules had identical appearance and taste. Capsules were taken under direct observation of study staff during school term time. Further details of the administration of study medication are provided in Supplementary Methods (online supplemental material).

Follow-up assessments

At 1-year, 2-year and 3-year follow-up, height, weight and waist circumference were measured as at baseline. At 3-year follow-up, substudy participants were additionally invited to undergo repeat spirometry and DXA scanning and to complete a Tanner self-assessment questionnaire for pubertal development.²⁴

Outcomes

The primary outcome of the main trial, reported elsewhere,¹⁸ was the QFT-Plus result at the end of the study. The following secondary outcomes were assessed for all participants: height-for-age, BMI-for-age, waist circumference-for-age and waist-to-height ratio (all participants). Additional outcomes assessed in substudy participants only were: whole body fat mass, fat-free soft tissue mass, % predicted forced vital capacity (FVC), % predicted forced expiratory volume in 1 s (FEV1), % predicted FEV1/FVC, mean Tanner scores for pubic hair (men and women), the external genitalia (men only) or breast development (women only), the proportion of participants reaching menarche by the end of the trial (women only) and mean age at menarche (women who reached menarche by the end of the trial only).

DXA

Body composition (whole body fat mass and fat-free soft tissue mass) was assessed using a Hologic Discovery-W DXA scanner at the Sports Science Institute of South Africa, University of Cape Town. All scans were performed by a trained radiographer on one scanner (Hologic, Bedford, Massachusetts, USA) using standard procedures, and analysed using Apex software (V.13.4.1). Quality assurance checks were carried out prior to scanning and generated coefficients of variation <0.5%. DXA outcomes relating to the effects of the trial intervention on bone mineral content have been reported elsewhere.¹⁹

Laboratory assessments

Serum concentrations of 25(OH)D₃ were measured using liquid chromatography-tandem mass spectrometry as previously described.²⁵ Further details are presented

in Supplementary Methods (online supplemental material).

Sample size

The sample size for the main trial was calculated as the number needed to detect a 25% reduction in the proportion of children with a positive QFT-Plus assay result at a 3-year follow-up (primary outcome) with 80% power and 5% type 1 error, assuming a 3.5% annual risk of QFT-Plus conversion, 20% loss to follow-up and a 5% risk of an indeterminate QFT-Plus assay result at the end of the study.¹⁸ The sample size for the substudy was calculated as the number needed to detect an inter-arm difference of 0.35 SDs in mean bone mineral content at the whole body less head and lumbar spine with 88% power and 5% type 1 error, assuming 29% loss to follow-up at 3 years.¹⁹

Statistical analyses

Full details of statistical analyses are provided in Supplementary Methods (online supplemental material). Briefly, effects of treatment on age-adjusted and sex-adjusted z-scores for anthropometric outcomes were estimated by fitting allocation to vitamin D versus placebo as the sole fixed effect in a mixed effects linear regression model with a random effect for repeated assessments of each individual participant and a random effect of the school of attendance. Effects of treatment on DXA and spirometric outcomes were analysed using multilevel mixed models with adjustment for baseline values and a random effect for school. Tanner metrics were analysed in a similar fashion, but restricted to the applicable sex and without baseline adjustment. Prespecified subgroup analyses were conducted to determine whether the effect of vitamin D supplementation was modified by sex (male vs female), baseline deseasonalised 25(OH)D₃ concentration (<75 vs ≥75 nmol/L)²⁶ and calcium intake (< vs ≥ median value of 466 mg/day).¹⁹ These were performed by repeating efficacy analyses with the inclusion of an interaction term between allocation (to vitamin D vs placebo) and each posited effect-modifier with the presentation of the p value associated with this interaction term. An Independent Data Monitoring Committee reviewed accumulating serious adverse event data at 6-monthly intervals, and recommended continuation of the trial at each review. No interim efficacy analysis was performed.

Patient and public involvement

The Desmond Tutu HIV Centre's Community Advisory Group was consulted on the design and conduct of the ViDiKids trial.

RESULTS

Participants

A total of 2852 children were screened for eligibility from March 2017 to March 2019, of whom 2271 underwent QFT testing: 1682 (74.1%) QFT-negative children were randomly assigned to receive vitamin D₃ (829

participants) or placebo (853 participants) as previously described.¹⁸ 450/1682 (26.8%) participants in the main trial also participated in the substudy, of whom 228 versus

222 participants were allocated to the vitamin D versus placebo arms, respectively (figure 1). Table 1 presents baseline characteristics of children in the main trial and

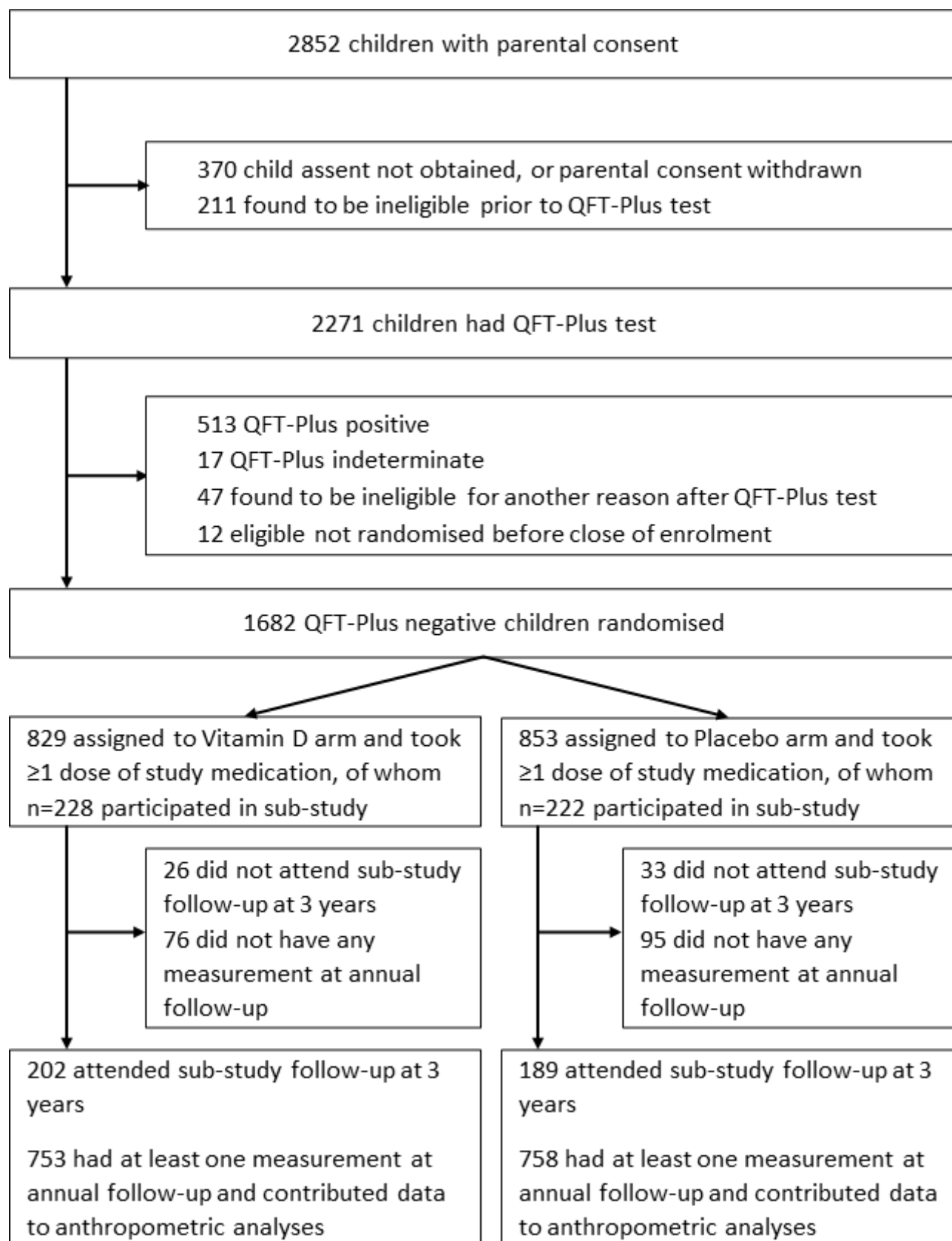


Figure 1 Participant flow diagram. QFT, QuantiFERON-TB.

Table 1 Participants' baseline characteristics by allocation: main study and substudy

	Main study (n=1682)			Substudy (n=450 subset)		
	Overall (n=1682)	Vitamin D arm (n=829)	Placebo arm (n=853)	Overall (n=450)	Vitamin D arm (n=228)	Placebo arm (n=222)
Mean age, years (SD)	8.9 (1.4)	8.9 (1.4)	8.8 (1.3)	10.1 (0.7)	10.2 (0.7)	10.0 (0.6)
Female sex, n (%)	880 (52.4)	437 (52.8)	443 (51.9)	234 (52.0)	116 (50.9)	118 (53.2)
Ethnic origin*						
Xhosa, n (%)	1615 (97.9)	788 (97.3)	827 (98.5)	424 (96.4)	214 (96.0)	210 (96.8)
Other, n (%)	35 (2.1)	22 (2.7)	13 (1.5)	16 (3.6)	9 (4.0)	7 (3.2)
Type of residence						
Brick, n (%)	867 (51.5)	423 (51.0)	444 (52.1)	230 (51.1)	121 (53.1)	109 (49.1)
Informal, n (%)	815 (48.5)	406 (49.0)	409 (47.9)	220 (48.9)	107 (46.9)	113 (50.9)
Parental education**†						
Primary school, n (%)	60 (3.6)	34 (4.1)	26 (3.1)	24 (5.3)	16 (7.0)	8 (3.6)
Secondary school or higher, n (%)	1618 (96.4)	792 (95.9)	826 (96.9)	426 (94.7)	212 (93.0)	214 (96.4)
Mean monthly household income, 1000 ZAR (SD)	1.88 (2.20)	1.81 (2.15)	1.95 (2.25)	1.57 (2.31)	1.52 (2.69)	1.63 (1.86)
Mean height-for-age z-score (SD)*	-0.55 (1.20)	-0.56 (1.26)	-0.54 (1.15)	-0.43 (0.96)	-0.43 (0.98)	-0.44 (0.95)
Height-for-age z-score category						
<-2.00 (stunted), n (%)	186 (11.1)	99 (11.9)	87 (10.2)	29 (6.4)	16 (7.0)	13 (5.9)
≥-2.00, n (%)	1496 (88.9)	730 (88.1)	766 (89.8)	421 (93.6)	212 (93.0)	209 (94.1)
Mean BMI-for-age z-score (SD)*	0.33 (1.03)	0.31 (1.03)	0.34 (1.03)	0.20 (0.97)	0.19 (0.92)	0.21 (1.02)
BMI-for-age z-score category						
<-2.00 (thin), n (%)	26 (1.5)	15 (1.8)	11 (1.3)	7 (1.6)	4 (1.8)	3 (1.4)
>-2.00 and ≤1.00, n (%)	1223 (72.7)	602 (72.6)	621 (72.8)	354 (78.7)	178 (78.1)	176 (79.3)
>1.00 and ≤2.00 (overweight), n (%)	340 (20.2)	165 (19.9)	175 (20.5)	72 (16.0)	42 (18.4)	30 (13.5)
>2.00 (obese), n (%)	93 (5.5)	47 (5.7)	46 (5.4)	17 (3.8)	4 (1.8)	13 (5.9)
Mean waist circumference-for-age z-score (SD)*	-0.13 (0.90)	-0.11 (0.90)	-0.15 (0.91)	-0.20 (0.98)	-0.20 (0.98)	-0.20 (0.98)
Mean waist-to-height ratio z-score (SD)*	0.03 (0.85)	0.05 (0.83)	0.00 (0.86)	-0.05 (0.86)	-0.05 (0.85)	-0.04 (0.88)
Mean whole body fat mass, kg (SD)	-	-	-	3.71 (2.26)	3.66 (2.13)	3.77 (2.40)
Mean whole body fat-free soft tissue mass, kg (SD)	-	-	-	9.27 (1.95)	9.39 (2.02)	9.15 (1.86)
% predicted FEV1 (SD)	-	-	-	83.2 (15.7)	83.8 (15.2)	82.7 (16.2)

Continued



Table 1 Continued

	Main study (n=1682)	Substudy (n=450 subset)
% predicted FVC (SD)	-	94.8 (18.7)
% predicted FEV1/FVC (SD)	-	91.4 (16.9)
Calcium intake*	<median, n (%) [‡] 817 (50.0)	434 (52.3)
	≥median, n (%) [‡] 817 (50.0)	193 (43.8)
Mean serum 25(OH)D ₃ concentration, nmol/L (SD) ^{*§}	71.2 (14.8)	70.0 (13.5)
Serum 25(OH)D ₃ concentration, category ^{*§}	<25 nmol/L, n (%) 1 (0.1)	1 (0.3)
	≥25 nmol/L and <50 nmol/L, n (%) 74 (5.4)	40 (5.8)
	≥50 nmol/L and <75 nmol/L, n (%) 787 (57.7)	393 (56.6)
	≥75 nmol/L, n (%) 502 (36.8)	214 (59.0)
		114 (61.0)
		66 (35.3)
		62 (35.2)

*Missing data (for bone substudy: ethnicity, n=5 vitamin D arm, n=5 placebo arm; for calcium intake, n=6 vitamin D arm, n=3 placebo arm; for serum 25(OH)D₃ concentration, n=66 vitamin D arm, n=62 placebo arm; for serum adjusted calcium and total ALP concentration, n=41 vitamin D arm, n=45 placebo arm; for PTH: n=42 vitamin D arm, n=46 placebo arm; for CTX: n=41 vitamin D arm, n=46 placebo arm; for P1NP: n=43 vitamin D arm, n=46 placebo arm; for fracture study: ethnicity, n=19 vitamin D arm, n=13 placebo arm; parental education, n=3 vitamin D arm, n=1 placebo arm; BMI-for-age z-score, n=2 vitamin D arm, n=0 placebo arm; height-for-age z-score, n=2 vitamin D arm, n=0 placebo arm; for calcium intake, n=25 Vitamin D arm, n=23 placebo arm; for serum 25(OH)D₃ concentration, n=159 vitamin D arm, n=159 placebo arm.

[‡]Highest level of education of at least one parent.

[§]Median calcium intake 466 mg/day.

[§]Deseasonalised values.

.ALP, alkaline phosphatase; BMC, bone mineral content; BMI, body mass index; CTX, C-terminal cross-linked telopeptide; N_i, propeptide; 25(OH)D₃, 25-hydroxyvitamin D₃; P1NP, serum procollagen type I; PTH, parathyroid hormone; ZAR, South African rand.

in the substudy, overall and by study arm. Mean age was higher among participants in the substudy versus all those in the main trial (10.1 vs 8.9 years, respectively), reflecting the fact that participation in the substudy was restricted to children enrolled in Grade 4. Prevalence of obesity and stunting were lower in the substudy versus the main trial (3.8% vs 5.5%, and 6.4% vs 11.1%, respectively). Baseline characteristics were otherwise well balanced for all participants in the main trial vs those who additionally participated in the substudy: 52.4% vs 52.0% were female and mean serum 25(OH)D₃ concentrations were 71.2 nmol/L vs 70.0 nmol/L. Within the main trial and the substudy, baseline characteristics of those randomised to vitamin D versus placebo were also well balanced. The median duration of follow-up was 3.16 years (IQR, 2.83–3.38 years) and was not different between the two study arms. For the main trial, mean serum 25(OH)D₃ concentrations at 3-year follow-up were higher among children randomised to receive vitamin D versus placebo (104.3 vs 64.7 nmol/L, respectively; mean difference 39.7 nmol/L, 95% CI for difference 37.6 to 41.9 nmol/L).

Growth outcomes

Among participants in the main trial, allocation to vitamin D versus placebo did not significantly influence mean height-for-age z-scores at annual follow-up, either overall or within subgroups defined by sex, calcium intake or baseline 25(OH)D concentration (table 2: p values for interaction >0.05). Similarly, no statistically significant effect of the intervention was seen on mean BMI-for-age z-scores (table 3), waist circumference-for-age z-scores (online supplemental table S1) or waist-to-height ratio z-scores (online supplemental table S2), either overall or by subgroup (p values for interaction >0.05). Among substudy participants, allocation to vitamin D versus placebo did not significantly influence fat mass or fat-free soft tissue mass (online supplemental table S3) or % predicted FEV1 or FEV1/FVC (online supplemental table S4), either overall or by subgroup. For the outcome of % predicted FVC, allocation to vitamin D versus placebo did not have a statistically significant effect overall, or within subgroups defined by calcium intake or baseline 25(OH)D concentration; however, the p value for interaction associated with the subgroup analysis by sex (p=0.049) raised the possibility that this factor might modify the effect of vitamin D on % predicted FVC (online supplemental table S4).

Developmental outcomes

Among substudy participants, no statistically significant inter-arm differences were seen in mean Tanner scores for pubic hair (men and women), external genitalia (men only), proportion menstruating (women only), mean age at menarche (women only) or breast development (women only), either overall or within subgroups defined by calcium intake or baseline 25(OH)D concentration (table 4).

Adverse events

Incidence of adverse events by the trial arm has been reported elsewhere.¹⁸ No serious events arising in the trial were adjudged related to the administration of vitamin D or placebo.

DISCUSSION

We report findings from the first RCT to investigate the effects of vitamin D supplementation on growth and developmental indices in schoolchildren of black African ancestry. Administration of oral vitamin D supplementation at a dose of 10000 IU per week for 3 years was effective in boosting vitamin D status, but it did not have statistically significant effects on linear growth, BMI, body composition, spirometric lung volumes or self-assessed pubertal development, either overall or in subgroups defined by sex, calcium intake or baseline vitamin D status.

Null findings from the current study contrast with those from observational studies reporting independent associations between vitamin D deficiency and reduced lean mass,⁸ slower linear growth,⁷ childhood obesity^{9–11} and precocious puberty.¹² However, they are consistent with those of the only other phase 3 RCT to investigate the effects of vitamin D on growth and development in school-age children that we are aware of, which reported no effect of weekly vitamin D supplementation on growth or development among Mongolian schoolchildren aged 6–13 years at baseline.¹⁷ Contrasting findings from observational studies vs clinical trials may reflect the fact that the former are more susceptible to the effects of confounding or reverse causality. Although we observed limited evidence (P for interaction 0.049) to support the hypothesis that the effects of vitamin D on FVC might be modified by sex, this finding may have arisen because of type I error, given the multiplicity of analyses conducted.

Our study has several strengths. The intervention was sustained (3 years), allowing ample time for any effects of vitamin D supplementation on outcomes of interest to manifest. Moreover, the intervention was effective in elevating serum 25(OH)D concentrations, reflecting adequacy of the dosing regimen employed as well as good adherence resulting from directly observed administration of weekly supplements in schools during term time. Our large sample size and low rates of loss to follow-up maximised the power to detect modest effects of the intervention, particularly for outcomes that were assessed in the main trial population. Furthermore, we assessed a broad range of anthropometric and developmental outcomes, that included the use of DXA, the gold standard investigation for the assessment of body composition.

Our study also has some limitations. The baseline prevalence of vitamin D deficiency was low, perhaps reflecting plentiful exposure to sunshine in the study setting.²⁵ Dietary factors are less likely to contribute to this phenomenon, as frequency of intake of oily fish and



Table 2 Mean height-for-age z-scores at annual follow-up by allocation, main study participants: overall and by subgroup

	Follow-up time point	Vitamin D arm: mean value (SD) (n)	Placebo arm: mean value (SD) (n)	Adjusted mean difference (95% CI)*	P for time point*	Overall P * interaction*	P for interaction*
Overall	1 year	-0.22 (1.12) (664)	-0.15 (1.07) (661)	-0.06 (-0.17 to 0.06)	0.33	0.22	-
	2 years	-0.14 (1.05) (613)	-0.06 (1.01) (606)	-0.04 (-0.16 to 0.07)	0.46		
	3 years	-0.26 (1.04) (670)	-0.14 (1.05) (691)	-0.08 (-0.19 to 0.03)	0.17		
By sex	Boys					0.42	0.94
	1 year	-0.31 (1.14) (304)	-0.27 (1.06) (306)	-0.02 (-0.17 to 0.14)	0.85		
	2 years	-0.21 (1.11) (285)	-0.19 (0.95) (283)	0.04 (-0.13 to 0.20)	0.67		
Girls	3 years	-0.34 (1.07) (310)	-0.28 (1.00) (326)	-0.03 (-0.19 to 0.13)	0.69	0.34	
	1 year	-0.15 (1.09) (360)	-0.04 (1.07) (355)	-0.10 (-0.25 to 0.06)	0.21		
	2 years	-0.08 (0.99) (328)	0.05 (1.06) (323)	-0.12 (-0.28 to 0.04)	0.14		
By calcium intake [†]	< median					0.02	0.07
	1 year	-0.16 (1.05) (314)	-0.10 (1.02) (331)	-0.04 (-0.18 to 0.11)	0.64		
	2 years	-0.17 (0.96) (288)	-0.05 (0.96) (310)	-0.04 (-0.19 to 0.11)	0.64		
≥ median	3 years	-0.28 (0.99) (312)	-0.15 (0.99) (362)	-0.08 (-0.23 to 0.07)	0.29	0.72	
	1 year	-0.25 (1.16) (327)	-0.18 (1.13) (314)	-0.05 (-0.21 to 0.12)	0.60		
	2 years	-0.07 (1.12) (307)	-0.06 (1.06) (279)	-0.03 (-0.20 to 0.14)	0.76		
By baseline 25(OH) D concentration [‡]	< 75 nmol/L					0.53	0.50
	1 year	-0.23 (1.13) (349)	-0.03 (1.11) (332)	-0.17 (-0.33 to -0.01)	0.04		
	2 years	-0.14 (1.11) (325)	0.05 (1.06) (323)	-0.12 (-0.28 to 0.04)	0.15		
≥ 75 nmol/L	3 years	-0.25 (1.10) (340)	-0.06 (1.07) (359)	-0.15 (-0.31 to 0.01)	0.07	0.15	
	1 year	-0.22 (1.15) (187)	-0.25 (0.98) (201)	0.02 (-0.17 to 0.22)	0.82		
	2 years	-0.17 (1.00) (171)	-0.08 (0.96) (179)	-0.07 (-0.26 to 0.13)	0.50		
3 years	-0.30 (1.00) (196)	-0.18 (0.99) (206)	-0.06 (-0.25 to 0.13)	0.53			

*Effect estimates and p values from mixed effects linear regression models including a fixed effect for allocation to vitamin D versus placebo, a random effect for repeated assessments of each individual participant and a random effect of school of attendance.

[†]Median calcium intake 466 mg/day.

[‡]De-seasonalised values.

.n, number; 25(OH)D, 25-hydroxyvitamin D;

Table 3 Mean BMI-for-age z-scores at annual follow-up by allocation, main study participants: overall and by subgroup

	Vitamin D arm: mean value (SD)(n)	Placebo arm: mean value (SD) (n)	Adjusted mean difference (95% CI)	P for time point	Overall P	P for interaction
Overall						
	1 year	0.16 (1.03) (664)	0.17 (1.02) (661)	-0.02 (-0.14 to 0.09)	0.68	0.86
	2 years	0.15 (1.41) (613)	0.25 (1.21) (606)	-0.09 (-0.20 to 0.03)	0.14	
	3 years	0.17 (1.07) (669)	0.21 (1.17) (691)	-0.04 (-0.16 to 0.07)	0.45	
By sex						
	1 year	0.08 (0.99) (304)	-0.00 (0.99) (306)	0.09 (-0.08 to 0.27)	0.30	0.59
	2 years	0.00 (1.68) (285)	-0.04 (1.30) (283)	0.04 (-0.14 to 0.22)	0.64	
	3 years	-0.02 (1.04) (309)	-0.12 (1.20) (326)	0.09 (-0.09 to 0.26)	0.33	
	1 year	0.23 (1.06) (360)	0.32 (1.02) (355)	-0.14 (-0.28 to 0.00)	0.06	0.26
	2 years	0.28 (1.10) (328)	0.50 (1.06) (323)	-0.21 (-0.36 to -0.06)	0.00	
	3 years	0.33 (1.06) (360)	0.50 (1.07) (365)	-0.17 (-0.32 to -0.03)	0.02	
By calcium intake [†]						
	1 year	0.15 (1.01) (314)	0.15 (1.03) (331)	-0.01 (-0.17 to 0.14)	0.86	0.16
	2 years	0.18 (1.08) (288)	0.21 (1.34) (310)	-0.03 (-0.19 to 0.13)	0.75	
	3 years	0.21 (1.07) (312)	0.20 (1.14) (362)	0.01 (-0.14 to 0.17)	0.87	
	1 year	0.18 (1.04) (327)	0.22 (1.01) (314)	-0.04 (-0.21 to 0.13)	0.61	0.27
	2 years	0.13 (1.67) (307)	0.31 (1.07) (279)	-0.15 (-0.32 to 0.03)	0.10	
	3 years	0.15 (1.07) (336)	0.22 (1.23) (311)	-0.09 (-0.26 to 0.08)	0.31	
By baseline 25(OH) D concentration [‡]						
	1 year	0.30 (1.03) (349)	0.28 (0.98) (332)	0.03 (-0.14 to 0.19)	0.76	0.31
	2 years	0.22 (1.64) (325)	0.31 (1.33) (323)	-0.07 (-0.24 to 0.10)	0.41	
	3 years	0.25 (1.07) (340)	0.34 (1.08) (359)	-0.09 (-0.25 to 0.08)	0.31	
	1 year	0.02 (0.97) (187)	0.09 (1.02) (201)	-0.06 (-0.25 to 0.13)	0.57	0.58
	2 years	0.01 (1.05) (171)	0.18 (1.00) (179)	-0.11 (-0.30 to 0.09)	0.28	
	3 years	0.09 (1.00) (196)	0.06 (1.17) (206)	0.02 (-0.17 to 0.21)	0.87	

*Effect estimates and p values from mixed effects linear regression models including a fixed effect for allocation to vitamin D versus placebo, a random effect for repeated assessments of each individual participant and a random effect of school of attendance.

[†]Median calcium intake 466 mg/day.

[‡]Deasonalised values.

[§]n, number; 25(OH)D, 25-hydroxyvitamin D;

Table 4 End-study pubertal development indices by allocation, substudy participants: overall and by subgroup

		Vitamin D arm: mean value (SD) (n)	Placebo arm: mean value (SD) (n)	Adjusted mean difference (95% CI)	P value	P for interaction	
Males	Overall	3.15 (0.91) (96)	2.81 (0.99) (90)	0.25 (-0.02 to 0.51)	0.07	-	
	Pubic hair, mean Tanner score (SD) (n)						
	Calcium intake [†]	<median	3.13 (0.93) (46)	2.73 (1.14) (52)	0.29 (-0.10 to 0.67)	0.14	0.39
		≥median	3.15 (0.90) (48)	2.92 (0.75) (38)	0.16 (-0.19 to 0.52)	0.37	
	Baseline 25(OH)D concentration [‡]	<75 nmol/L	3.11 (0.92) (44)	2.88 (0.92) (42)	0.07 (-0.28 to 0.41)	0.70	0.39
		≥75 nmol/L	3.18 (0.94) (34)	2.70 (1.18) (30)	0.43 (-0.07 to 0.93)	0.09	
	Overall	3.19 (0.91) (96)	2.90 (0.99) (90)	0.17 (-0.09 to 0.43)	0.19	-	
	External genitalia, mean Tanner score (SD) (n)						
	Calcium intake [†]	<median	3.28 (0.89) (46)	2.81 (1.10) (52)	0.35 (-0.01 to 0.71)	0.06	0.07
		≥median	3.10 (0.93) (48)	3.03 (0.82) (38)	-0.04 (-0.40 to 0.33)	0.84	
Females	Overall	3.27 (0.82) (44)	2.90 (0.91) (42)	0.22 (-0.10 to 0.54)	0.18	0.81	
	Pubic hair, mean Tanner score (SD) (n)						
	Calcium intake [†]	<75 nmol/L	3.15 (1.02) (34)	2.87 (1.17) (30)	0.30 (-0.17 to 0.77)	0.21	
		≥75 nmol/L	3.10 (0.98) (105)	3.11 (0.98) (99)	-0.09 (-0.35 to 0.17)	0.50	-
	Baseline 25(OH)D concentration [‡]	<75 nmol/L	3.15 (0.98) (47)	3.15 (1.01) (46)	0.01 (-0.38 to 0.40)	0.96	0.60
		≥75 nmol/L	3.11 (0.92) (54)	3.10 (0.97) (50)	-0.23 (-0.56 to 0.10)	0.17	
	Overall	3.14 (0.90) (59)	3.13 (0.88) (55)	-0.03 (-0.35 to 0.28)	0.83	0.94	
	External genitalia, mean Tanner score (SD) (n)						
	Calcium intake [†]	<75 nmol/L	2.96 (1.22) (26)	3.04 (1.08) (24)	-0.14 (-0.74 to 0.46)	0.65	
		≥75 nmol/L	3.15 (0.87) (105)	3.10 (0.91) (99)	-0.03 (-0.26 to 0.20)	0.80	-
Breast development, mean Tanner score (SD) (n)	Overall	3.15 (0.91) (47)	3.22 (0.92) (46)	-0.06 (-0.41 to 0.29)	0.74	0.93	
	Calcium intake [†]	<median	3.20 (0.81) (54)	3.04 (0.90) (50)	-0.05 (-0.36 to 0.26)	0.76	
		≥75 nmol/L	3.27 (0.85) (59)	3.07 (0.86) (55)	0.15 (-0.16 to 0.45)	0.34	0.48
	Baseline 25(OH)D concentration [‡]	<75 nmol/L	3.08 (0.93) (26)	3.08 (0.97) (24)	-0.08 (-0.55 to 0.38)	0.73	
		≥75 nmol/L	67/105 (63.81)	60/99 (60.61)	0.95 (0.51 to 1.77)	0.86	--
	Overall	28/47 (59.57)	26/46 (56.52)	1.33 (0.50 to 3.56)	0.57	0.24	
	Proportion menstruating	<median	38/54 (70.37)	34/50 (68.00)	0.56 (0.21 to 1.52)	0.26	
		≥75 nmol/L	43/59 (72.88)	35/55 (63.64)	1.39 (0.59 to 3.28)	0.45	0.59
	Baseline 25(OH)D concentration [‡]	<75 nmol/L	16/26 (61.54)	15/24 (62.50)	0.90 (0.26 to 3.15)	0.87	
		≥75 nmol/L	11.94 (0.93) (66)	11.86 (0.66) (58)	-0.03 (-0.31 to 0.25)	0.84	-
Mean age at menarche, years (SD) (n)	Overall	11.89 (0.83) (28)	11.71 (0.69) (24)	0.12 (-0.26 to 0.49)	0.54	0.32	
	Calcium intake [†]	<median	11.97 (1.01) (37)	11.97 (0.63) (34)	-0.11 (-0.51 to 0.29)	0.58	
		≥75 nmol/L	12.00 (0.88) (42)	11.88 (0.69) (34)	0.04 (-0.31 to 0.39)	0.82	0.64
	Baseline 25(OH)D concentration [‡]	<75 nmol/L	11.88 (1.15) (16)	11.93 (0.62) (14)	-0.14 (-0.73 to 0.44)	0.63	

*Effect estimates and p values from multilevel mixed models including a random effect for school of attendance.

[†]Median calcium intake 466 mg/day.

[‡]De-seasonalised values.

.n., number; 25(OH)D, 25-hydroxyvitamin D.

other foods containing vitamin D did not associate with baseline vitamin D status in the study population.²⁵ Our results cannot therefore be generalised to settings where vitamin D deficiency is common. However, we highlight that results from our trial in Mongolia (where the baseline prevalence of vitamin D deficiency was much higher than we observed in the current study) were also null for outcomes of linear growth and body mass index.¹⁷ Accordingly, null results from the current trial cannot be explained simply by the relatively high baseline vitamin D status of participants in Cape Town. A second potential limitation relates to the fact that pubertal status was assessed by participants themselves, rather than by health professionals, whose judgement may be more objective. However, the double-blind placebo-controlled trial design of our study will have distributed any resultant imprecision equally between study arms, ensuring that bias was not introduced. Moreover, pubertal self-assessment has previously been done by adolescent participants in another South African cohort study, and shown to be reliable.²⁷

In conclusion, we report that oral vitamin D supplementation at a dose of 10 000 IU/week for 3 years was effective in elevating serum 25(OH)D concentrations in schoolchildren of black African ancestry living in Cape Town, South Africa. However, this was not associated with any effects on linear growth, body habitus, spirometric outcomes or pubertal development. Taken together with null results from a trial of weekly vitamin D supplementation with similar outcomes conducted in Mongolia,¹⁷ our findings do not support the use of vitamin D supplements to influence child growth or development in populations where rickets have been excluded. Further research could be conducted to re-evaluate growth and developmental outcomes in the future, to determine whether any adverse effects might be associated with withdrawing vitamin D supplementation from individuals randomised to the intervention arm of this study, who may have become habituated to it during their participation in the trial.

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Contributors ARM conceived the study. KM, LM, AKC, JN, CC, NCH, RLH, RJW, L-GB and ARM contributed to study design and protocol development. KM led on trial implementation, with support from JS, DAJ, JN, L-GB and ARM. LM and AEM oversaw the performance of DXA scans. JT and WDF performed and supervised the conduct of biochemical assays. NW and AM drafted the statistical analysis plan. DAJ, KM, JS, NW and WM managed data. NW accessed, verified and analysed the data underlying the study. ARM and DAJ wrote the first draft of the trial report. All authors made substantive comments thereon and approved the final version for submission. ARM is guarantor for this report.

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REFERENCES

- 1 Eccles JS. The development of children ages 6 to 14. *Future Child* 1999;9:30–44.
- 2 Dewey KG, Begum K. Long-term consequences of Stunting in early life. *Matern Child Nutr* 2011;7 Suppl 3(Suppl 3):5–18.
- 3 Vaivada T, Akseer N, Akseer S, *et al.* Stunting in childhood: an overview of global burden, trends, determinants, and drivers of decline. *The American Journal of Clinical Nutrition* 2020;112(Suppl 2):777S–791S.
- 4 Di Cesare M, Sorić M, Bovet P, *et al.* The Epidemiological burden of obesity in childhood: a worldwide epidemic requiring urgent action. *BMC Med* 2019;17.
- 5 Eckert-Lind C, Busch AS, Petersen JH, *et al.* Worldwide secular trends in age at pubertal onset assessed by breast development among girls: A systematic review and meta-analysis. *JAMA Pediatr* 2020;174:e195881.
- 6 Bikle DD, *et al.* *Vitamin D: production, metabolism and mechanisms of action in.* South Dartmouth (MA, 2000.
- 7 Gilbert-Diamond D, Baylin A, Mora-Plazas M, *et al.* Vitamin D deficiency and Anthropometric indicators of Adiposity in school-age children: a prospective study. *Am J Clin Nutr* 2010;92:1446–51.
- 8 Foo LH, Zhang Q, Zhu K, *et al.* Relationship between vitamin D status, body composition and physical exercise of adolescent girls in Beijing. *Osteoporos Int* 2009;20:417–25.
- 9 Zakharova I, Klimov L, Kuryaninova V, *et al.* Vitamin D insufficiency in overweight and obese children and adolescents. *Front Endocrinol (Lausanne)* 2019;10.
- 10 Plesner JL, Dahl M, Fonvig CE, *et al.* Obesity is associated with vitamin D deficiency in Danish children and adolescents. *J Pediatr Endocrinol Metab* 2018;31:53–61.
- 11 Theodoratou E, Tzoulaki I, Zgaga L, *et al.* Vitamin D and multiple health outcomes: umbrella review of systematic reviews and meta-analyses of observational studies and randomised trials. *BMJ* 2014;348.
- 12 Lee HS, Kim YJ, Shim YS, *et al.* Associations between serum vitamin D levels and precocious puberty in girls. *Ann Pediatr Endocrinol Metab* 2014;19:91–5.
- 13 Liu S, Zhu X, Wang Y, *et al.* The association between vitamin D levels and precocious puberty: a meta-analysis. *J Pediatr Endocrinol Metab* 2020;33:427–9.
- 14 Gou Z, Li F, Qiao F, *et al.* Causal associations between insulin-like growth factor 1 and vitamin D levels: a two-sample Bidirectional Mendelian randomization study. *Front Nutr* 2023;10.
- 15 Nimitphong H, Park E, Lee MJ. Vitamin D regulation of Adipogenesis and Adipose tissue functions. *Nutr Res Pract* 2020;14:553–67.
- 16 Ganmaa D, Stuart JJ, Sumberzul N, *et al.* Vitamin D supplementation and growth in urban Mongol school children: results from two randomized clinical trials. *PLoS One* 2017;12:e0175237.
- 17 Ganmaa D, Bromage S, Khudyakov P, *et al.* Influence of vitamin D supplementation on growth, body composition, and pubertal development among school-aged children in an area with a high prevalence of vitamin D deficiency: A randomized clinical trial. *JAMA Pediatr* 2023;177:32–41.
- 18 Middelkoop K, Stewart J, Walker N, *et al.* Vitamin D supplementation to prevent tuberculosis infection in South African schoolchildren: Multicentre phase 3 double-blind randomised placebo-controlled trial (Vidikids). *Int J Infect Dis* 2023;134:63–70.
- 19 Middelkoop K, Micklesfield LK, Walker N, *et al.* Influence of vitamin D supplementation on bone mineral content, bone turnover markers, and fracture risk in South African schoolchildren: multicenter double-blind randomized placebo-controlled trial (Vidikids). *J Bone Miner Res* 2024.
- 20 Graham BL, Steenbruggen I, Miller MR, *et al.* Standardization of Spirometry 2019 update. an official American Thoracic society and European respiratory society technical statement. *Am J Respir Crit Care Med* 2019;200:e70–88.
- 21 Rajakumar K, Moore CG, Yabes J, *et al.* Estimations of dietary vitamin D requirements in black and white children. *Pediatr Res* 2016;80:14–20.
- 22 Iglay K, Cao X, Mavros P, *et al.* Systematic literature review and meta-analysis of medication adherence with once-weekly versus once-daily therapy. *Clin Ther* 2015;37:1813–21.
- 23 El-Hajj Fuleihan G, Nabulsi M, Tamim 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.
- 24 Chavarro JE, Watkins DJ, Afeiche MC, *et al.* Validity of self-assessed sexual maturation against physician assessments and hormone levels. *J Pediatr* 2017;186:172–8.
- 25 Middelkoop K, Walker N, Stewart J, *et al.* Prevalence and determinants of vitamin D deficiency in 1825 Cape town primary schoolchildren: A cross-sectional study. *Nutrients* 2022;14.
- 26 Ganmaa D, Khudyakov P, Buyanjargal U, *et al.* Prevalence and determinants of Quantiferon-diagnosed tuberculosis infection in 9810 Mongolian schoolchildren. *Clin Infect Dis* 2019;69:813–9.
- 27 Norris SA, Richter LM. Usefulness and reliability of Tanner pubertal self-rating to urban black adolescents in South Africa. *J of Research on Adolesc* 2005;15:609–24.