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Epidemiology and risk factors of hypovitaminosis D in a cohort of internationally adopted children: a retrospective study

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Abstract

Background: Predictors of hypovitaminosis D were extensively studied in the adult population, leading to an approximately complete understanding of them, while there is a lack of studies in the pediatric population, especially in migrant and internationally adopted children.

In this retrospective study, we tried to identify the major laboratory predictors of hypovitaminosis D in a cohort of internationally adopted children.

Methods: Data were extracted from the database of the "Ethnopediatrics Outpatient Clinic" of the "A. Gemelli" Foundation University Hospital in Rome, Italy. Our study included 873 children evaluated from March 2007 to May 2016. Analysis of variance, chi square test, t test and multivariate logistic regression were performed, a "p" value < 0.05 was considered significant, with a confidence interval of 95%.

Results: We did not find any significant correlation between Vitamin D and Calcium, Phosphates or Magnesium levels within the population we examined. Moreover, parathyroid hormone is not a good predictor of Vitamin D Status.

Conclusions: Considering the strong influence Vitamin D status has not only on bone health, but also on general wellbeing, it is due to perform a vitamin D assessment as soon as possible, especially in internationally adopted children.

Keywords: Vitamin D deficiency, Internationally adopted children, Parathormone, Rickets

Background

Vitamin D (VD) is an essential nutrient with hormone-like activity, which regulates calcium and bone metabolism. Moreover, VD has been recently associated with immune status, autoimmune disorders, infectious diseases, decreased response to corticosteroids and psychiatric diseases [1-5].

These associations highlighted the need for a definition of VD optimal values, and this is particularly true for the pediatric population, because of the importance

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of VD for the bone health and the immune status during the tricky phase represented by the growth period.

Serum 25-hydroxy-vitamin D (25(OH)D) concentration is currently the marker of choice, as it is the major circulating form of VD and its value reflects both the amount produced in the skin after sun exposure and that assumed with food [6].

Most studies consider values < 10 ng/ml or < 20 ng/ml as deficiency and values > 20 ng/ml or > 30 ng/ml as sufficiency, with a gray zone between 10 and 20 ng/ml and 20–30 ng/ml, respectively, indicated as insufficiency [7–11]. When 25(OH)D serum concentration falls below 4 ng/ml, especially if associated with hypocalcaemia, hypophosphatemia and increased serum alkaline phosphatase (ALP), the risk of rickets and osteomalacia in children rises exponentially [12].



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Relatively high rates of subclinical vitamin D deficiency (VDD) have been reported in otherwise healthy infants, children and adolescents in several studies, especially from low-income countries [13-26].

Limited data on the prevalence of hypovitaminosis D among internationally adopted children, a population which could be considered "at risk" for geographical origin and a housing system not always subdued to an optimal surveillance, are available, and there is no study assessing parathyroid hormone (PTH) levels in these groups. The relationship between serum concentrations of 25(OH)D and PTH has been considered as a possible way to define VD status, since PTH increases when vitamin D and, consequently, calcium decrease. Studies performed in adults, and very few studies in pediatric subjects, have showed conflicting results about this inverse relationship [27].

Therefore, it would be of utmost importance to define VD levels under which PTH begins to increase, and if it could be considered a marker of bone health or immune status.

We hypothesized that the prevalence of biochemical rickets and hyperparathyroidism in a population of internationally adopted children was higher than in other children subpopulations. Hence, we evaluated the prevalence of hypovitaminosis D in internationally adopted children. Moreover, we tested if some of the population characteristics, such as age, BMI, sex, housing solution before adoption, season and month of observation, country of origin, could be considered risk factors for hypovitaminosis D in this particular cohort, aiming to find a model applicable to all our patients.

In order to define the prevalence of biochemical rickets, we considered the relationship existing between Vitamin D levels and serum calcium, serum phosphates and serum alkaline phosphatase levels. We also tried to define the relationship between VD status and PTH levels in internationally adopted children.

Methods

Characteristics of the study population

We retrospectively gathered data provided by medical records of the Ethnopediatrics outpatient clinic of the "A. Gemelli" University Hospital in Rome, Italy. Inclusion criteria of our study were: being an internationally adopted child (age 1–15) and having one determination of 25(OH)D values during the first observation in our outpatient clinic from March 2007 to May 2016. The only exclusion criteria we applied was the lack of 25(OH)D measurement during the first observation. At the end of the assessment of the medical records, 859 children were included in the study.

Missing data about physical characteristics, due to the refusal of the child to undergo physical evaluation at the time of the first visit, were treated as "missing", as well as missing data about clinical history of the child, due to a lacking anamnesis, were.

Retrieving the data: Biochemical analysis methods and normal values

Blood samples were collected by venipuncture in the morning and routinely analyzed at the Hormone Laboratory ("A. Gemelli" University Hospital).

Serum 25(OH)D was measured by ChemiLuminescence ImmunoAssay (CLIA, Siemens) with an intra- and inter-assay variation of 5%. This method does not allow to differentiate D_2 and D_3 forms. The lower detection limit of the assay was 7 ng/mL. We entered in the database every value indicated as <7 ng/mL within the test results as 6 ng/mL. Normal values are 31–100 ng/mL.

Serum intact parathyroid hormone (PTH) was assayed by ElettroChemiLuminescence ImmunoAssay (ECLIA, Roche) with an intra- and interassay CVs of 5% (normal values 9-65 pg/mL). Serum calcium (Ca) and phosphates (P) were determined by colorimetric assay (COBAS, Siemens). Serum calcium normal values are 8.8-10.8 mg/dl for all ages, while serum P values are 4-7 mg/dL when the child is younger than 10 years, or 2.5-4.5 mg/dL when the child is older than 10 years. Intra- and interassay CVs were 2.7 and 2.9%, respectively. Alkaline phosphatase (ALP) was measured by kinetic before 2011, normal values < 750 IU/L for children younger than 10 years, <1000 IU/L for children older than 10 years. A kinetic photometric test IFCC (AMP) after 2011 (Siemens) was used instead after 2011, normal values were 40-300 IU/L for children aged <13-years-old, while for children older than 13 years it is necessary to differentiate between males (normal values 40-309 IU/L) and females (normal values 40-187 IU/L). CVs intra- and interassay was 4% respectively.

Missing data, due to unavailability of reagents at certain time in our laboratory, are treated as "missing".

Assessment of 25-OH-D status

We chose to apply the Endocrine Society clinical practice guidelines cut-offs [28, 29]. Therefore, VD status was defined as severe deficiency when 25(OH)D was lower than 10 ng/ml, moderate deficiency when 25(OH)D value was between 10 and 20 ng/ml, mild deficiency when 25(OH)D value was between 20 and 30 ng/ml, or sufficiency when 25(OH)D was above 30 ng/ml.

Biochemical rickets was defined according to the following criteria: VDD (all categories) associated with increased ALP levels, normal or decreased calcium levels and normal or decreased phosphate levels.

Statistical analyses

The population in study was divided into three climatic zones on the basis of their birth country: temperate zone (from circles to the latitude of 40° N/S), subtropical zone (from 40° N/S to the tropic of cancer/capricorn) and tropical zone (from Cancer's tropic to Capricorn's tropic).

Standing height was measured with a wall-mounted stadiometer, while standing body weight was measured with a mechanical scale. BMI was calculated using the Quetelet equation (Weight (kg)/Height (m)²). Height, weight, and BMI were expressed in centiles. BMI was categorized into age- and gender-specific percentiles for children and adolescents (WHO, 2006): i) BMI < 3° centile – severly underweight; ii) 3–10° centile – underweight; iii) BMI 11–85° centile – normal weight; iv) BMI 86–95° centile – overweight; v) BMI > 95° centile – obese.

Skin color was categorized into three groups: very fair/ fair, intermediate, olive/brown. The Von Luschan's chromatic scale was used to differentiate skin colors (0–13 on Von Luschan's scale was categorized as "very fair/ fair"; 14–20 as "intermediate"; 21–36 as "olive/brown").

Four groups were arranged on the basis of the season of the blood sampling: winter (December 21 – March 20), spring (March 21 – June 20), summer (June 21 – September 20) and autumn (September 21 – December 20).

Descriptive and inferential statistics were carried out using the Statistical Package of Social Sciences (Chicago, IL, USA) for Windows software program version 22.0. Frequencies and percentages (%) were used to describe categorical variables (gender, macro-area, climatic zones, skin color, season at blood sampling, housing solution before adoption, category of BMI, VD status, PTH status, ALP status, Ca status, P status, biochemical rickets), while mean and standard deviation (mean ± SD) were used to express continuous variables (age on arrival in Italy, age at blood draw, time between the arrival and the blood draw, duration of the institutionalization, height, weight, BMI, 25(OH)D, PTH, ALP, Ca, P). We determined the normal distribution of continuous variables with a Kolmogorov-Smirnof test: the result permitted us to use parametric tests.

Differences among the categorical variables were tested for statistical significance using the Chi-square test. A p value < 0.05 was considered significant, Confidence Interval (CI) was 95%. Differences among mean values of continuous variables were tested for statistical significance using the Analysis of Variance (ANOVA) test and T test. A p value < 0.05 was considered significant, with a CI = 95%.

Post hoc analysis was performed for χ^2 tests significant results, when a contingency table bigger than 2*2 was generated Bonferroni adjusted *p*-values were considered to determine which couple of categorical variables significantly differed.

A multivariate logistic regression was performed for exploring possible predictors of low 25(OH)D levels at a cut-off point of 30 ng/ml.

Variables to be entered in the multivariate model were chosen on the basis of the univariate analysis results (p < 0.25). To identify the best model a backward approach was developed, based on Likelihood Ratio test. The final model was assessed in terms of calibration and discrimination through Hosmer-Lemeshow and C-statistics.

ORs for proportions below these cut-offs and their confidence intervals are reported.

Results

Characteristics of the population, season at blood drawing and mean values of the analytes investigated (25(OH)D, PTH, ALP, Ca, P) are resumed in Table 1. Detailed informations about post-hoc analyses are included in Additional file 1.

Vitamin D mean value and status and characteristics of the population (Table 2)

We did not find any significant difference regarding differences between mean values in females and males (p = 0.594) regarding mean 25(OH)D values. On the other hand, we found a slightly significant influence of sex on Vitamin D status (p = 0.043). Post hoc analysis determined that sex was significantly related to VD status only when testing a severe versus a moderate VDD, with female children having an increased risk of developing severe VDD than male children (OR 0.55, 95% CI 0.36– 0.86).

We did not find any statistically significant difference between 25(OH)D mean values and BMI categories (p = 0.477). The difference is not statistically significant either when applied to Vitamin D status (p = 0.521).

A statistically significant difference was found for skin color (p < 0.001) regarding mean 25(OH)D values. Mean 25(OH)D value was 18.00 ng/mL (SD ± 10.78 ng/mL) for children included in the group "olive to brown"; 20.86 ng/mL (SD ± 10.12 ng/mL) for children included in the group "very fair to fair"; 23.24 ng/mL (SD ± 11.76 ng/mL) for children included in the group "intermediate". This difference remains statistically significant when considering Vitamin D Status (p < 0.001). However, post hoc analysis highlighted that statistically significant differences only exist with regards to intermediate skin color when compared to olive/brown skin color. As a matter of fact, the character "olive/ brown" skin resulted to be predictive of a worse VD status both when the severe VDD status is compared to mild VDD status (OR 0.27, 95% CI 0.151-0.489) and

Total	859	100%
Sex (n = 859)		
Male	483	56.2%
Female	376	43.8%
Macro-area of origin (n = 859)		
Europe and Russian federation	256	29.8%
Latin America	231	26.9%
Asia and Indian subcontinent	223	26.0%
Africa	149	17.3%
Climatic zone of origin (n = 859)		
Tropical	438	51.0%
Sub-tropical	166	19.3%
Temperate	255	29.7%
Housing solution before adoption ($n = 828$)		
Foster home	70	8.5%
Foster family	65	7.9%
Institute	693	80.6%
Mean duration of institutionalization ($n = 750$)	3.00 years	SD \pm 2.03 years
Complexion (n = 859)		
Very fair/Fair	290	33.8%
Intermediate	340	39.6%
Olive/Brown	229	26.7%
Mean height (n = 793)	109 cm	SD ± 21 cm
Mean weight (n = 797)	19.93 kg	SD ± 9.02 kg
Mean BMI (n = 699)	16.0	SD ± 2.28
BMI status (n = 699)		
Obese	59	8.4%
Overweight	67	9.6%
Normal	536	76.7%
Underweight	37	5.3%
Mean age at the arrival in Italy (n = 808)	5.31 years	$SD \pm 2.92$ years
Mean age at the blood draw (n = 859)	5.61 years	SD \pm 2.97 years
Mean time from the arrival to the blood draw (n = 808)	0.33 years	SD \pm 0.44 years
Season at blood draw (n = 859)		
Spring	191	22.2%
Summer	149	17.3%
Fall	259	30.2%
Winter	260	30.3%
Mean Vitamin D (n = 859)	21.04 ng/mL	SD ± 11.15 ng/ml
Vitamin D status (n = 859)		
Severe Vitamin D Deficiency (< 10 ng/mL)	107	12.5%
Moderate Vitamin D Deficiency (10–20 ng/mL)	348	40.5%
Mild Vitamin D Deficiency (20–30 ng/mL)	276	32.1%
Normal Vitamin D (> 30 ng/mL)	128	14.9%
Mean PTH (n = 822)	32.24 pg/mL	SD ± 15.56 ng/ml

BMI body mass index, PTH Parathyroid hormone, ALP Alkaline phosphatase, Ca Serum calcium, P Serum phosphates

when the severe VDD status is compared to normal VD status (OR 0.196, 95% CI 0.101-0.391).

More detailed informations about the results above illustrated can be found in Table 2 and in Additional file 1.

Time from the arrival to the first evaluation was not found to be statistically related with 25(OH)D mean values (p = 0.388) and Vitamin D Status (p = 0.912). Age at the arrival in Italy was significantly associated both with 25(OH)D mean values (p < 0.001) and Vitamin D status (p < 0.001). Age at the blood draw association with 25(OH)D mean values (p = 0.049) was marginally significant, but it had a highly statistically significative relationship with Vitamin D status (p < 0.001).

Table 3 shows mean time from the arrival to first evaluation, age at the arrival and age at first evaluation in relation to Vitamin D Status. Table 4 shows age at the arrival in Italy in years (mean ± SD) per Vitamin D Status, and its relations with sex, macro-area, season at blood draw, housing solution and BMI status.

Figure 1 shows mean age at the arrival, in years, and its relationship with Vitamin D Status, while fig. 2 shows

Table 2 Vitamin D) status and	characteristics	of the	population
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		Vitamin D Status	Vitamin D Status					
		Severe VDD (< 10 ng/mL)	Moderate VDD (10–20 ng/mL)	Mild VDD (20–30 ng/mL)	Normal VD (> 30 ng/mL)	p values		
Sex (n = 859)	Male	49 (10.1%)	211 (43.7%)	149 (30.9%)	74 (15.3%)	0.043		
	Female	58 (15.4%)	137 (36.4%)	127 (33.8%)	54 (14.4%)			
BMI category ($n = 699$)	Underweight	2 (5.4%)	17 (45.9%)	14 (37.8%)	4 (10.8%)	0.521		
	Normal weight	78 (14.5%)	235 (43.8%)	160 (29.9%)	63 (11.8%)			
	Overweight	11 (16.4%)	21 (31.3%)	26 (38.8%)	9 (13.4%)			
	Obese	7 (11.9%)	29 (49.1%)	18 (30.5%)	5 (8.5%)			
Skin color <i>(n = 859)</i>	Very fair/fair	36 (12.4%)	115 (39.7%)	98 (33.8%)	41 (14.1%)	< 0.001		
	Intermediate	26 (7.6%)	128 (37.7%)	121 (35.6%)	65 (19.1%)			
	Olive/Brown	45 (19.6%)	105 (45.9%)	57 (24.9%)	22 (9.6%)			

 χ^2 -test was used to determine statistically significant relationships among categorical variables

Bonferroni-adjusted P value was uses to determine if there were statistically significant differences among groups

VDD Vitamin D Deficiency; VD Vitamin D; BMI Body Mass Index

Total

Decreased

		10070
PTH status (n = 822)		
Increased (> 65 pg/mL)	24	2.8%
Normal (10–65 pg/mL)	795	96.7%
Decreased (< 10 pg/mL)	3	0.4%
Mean ALP ($n = 831$)	385.01 IU/L	SD ± 299.27 IU/L
ALP status ($n = 831$)		
Increased	201	24.2%
Normal	630	75.8%
Mean Ca (n = 824)	9.94 mg/dL	SD \pm 0.45 mg/dL
Ca status (n = 824)		
Increased	18	2.2%
Normal	803	97.5%
Decreased	3	0.4%
Mean P ($n = 813$)	4.86	mg/dL
P status (n = 813)		
Increased	46	5.7%
Normal	731	89.9%

859

36

Table 1 Characteristics of the population included in the study (Continued)

100%

4.4%

Table 3 Vitamin D status and mean time from the arrival to 1st evaluation, age at the arrival and age at 1st evaluation

	Vitamin D Status					
	Severe VDD (< 10 ng/mL)	Moderate VDD (10–20 ng/mL)	Mild VDD (20–30 ng/mL)	Normal VD (> 30 ng/mL)	p values	
Time from the arrival in Italy to 1st evaluation ($n = 808$) mean \pm SD (years)	0.32 ± 0.47	0.34 ± 0.41	0.32 ± 0.47	0.32 ± 0.47	0.912	
Age at the arrival in Italy ($n = 808$) mean \pm SD (years)	6.31 ± 2.84	5.71 ± 2.77	5.09 ± 2.79	3.86 ± 3.07	< 0.001	
Age at 1st evaluation ($n = 859$) mean \pm SD (years)	6.69 ± 2.85	6.03 ± 2.79	5.36 ± 2.77	4.11 ± 3.15	< 0.001	

ANOVA test was used to determine statistical significance

mean 25(OH)D values for children aged < 1 year and their relationship with macro-area of origin.

Vitamin D mean value and status, geographical origin and social environment (Table 5)

Macro-area of origin was found to have a statistically significant relationship with 25(OH)D mean values (p = 0.001) and Vitamin D status (p < 0.001). Mean 25(OH)D value in children coming from Africa was 19.04 ng/mL (SD ± 11.67 ng/mL); in children coming from Latin America was 20.18 (SD ± 7.56 ng/mL); in children coming from Europe and Russian Federation was 21.00 ng/mL (SD ± 10.27 ng/mL); in children coming from Asia and Indian subcontinent was 23.31 ng/mL (SD ± 14.17 ng/mL).

Post hoc analysis highlighted that statistically significant differences existed within children coming from Asia and children coming from Africa. As a matter of fact, children coming from Africa tent to have more "intermediate" values than children coming from Asia. The result was statistically significant when comparing severe VDD status with moderate VDD status and moderate VDD status with normal VD status, with different outcomes. Children coming from Africa tend to have higher values of VD when comparing the severe and the moderate VDD status (OR 3.101, 95% CI 1.659–5.927), but lower VD values when comparing the moderate VDD status with the normal VD status (OR 0.298, 95% CI 0.171–0.531).

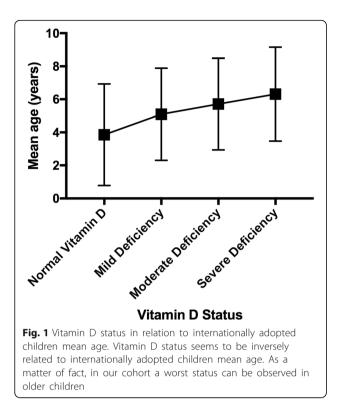
Details about the results above described can be found in Table 5 and Additional file 1.

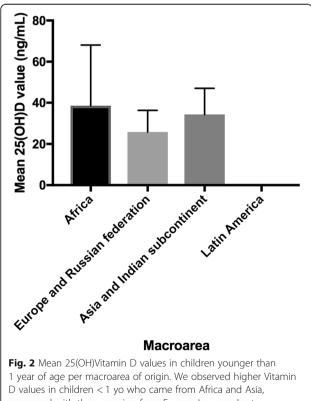
Climatic-zones of origin were also found to have a significant influence over 25(OH)D mean values (p < 0.001) and Vitamin D Status (p = 0.002). Mean 25(OH)D value in children coming from a tropical area was 22.27 ng/mL (SD ± 12.24 ng/mL), in children coming from a subtropical area was 17.83 ng/mL (SD ± 8.51 ng/mL), while in children coming from a temperate zone was 21.00 ng/mL (SD ± 10.28 ng/mL).

Table 4 Age at the arrival in Italy in years (Mean \pm standard deviation) per Vitamin D status

		Vitamin D status			
		Severe VDD	Moderate VDD	Mild VDD	Normal
Sex	Male	6.15 ± 2.77	5.50 ± 2.75	4.91 ± 2.59	3.74 ± 2.75
	Female	6.44 ± 2.92	6.06 ± 2.78	5.32 ± 3.01	4.02 ± 3.49
Macro-area	Africa	5.18 ± 2.48	4.85 ± 2.19	4.24 ± 2.51	3.16 ± 2.64
	Europe	6.16 ± 2.84	5.80 ± 2.71	5.28 ± 2.43	4.88 ± 3.40
	Asia	6.32 ± 2.99	4.14 ± 2.15	3.25 ± 2.39	1.82 ± 1.29
	Latin America	8.08 ± 2.37	7.16 ± 2.80	6.99 ± 2.37	6.31 ± 2.68
Season at blood draw	Spring	6.46 ± 2.83	6.09 ± 2.85	3.87 ± 2.49	2.44 ± 1.95
	Summer	5.77 ± 3.22	5.24 ± 2.61	4.62 ± 2.59	4.55 ± 3.73
	Fall	5.70 ± 2.96	5.57 ± 2.61	5.59 ± 2.72	3.85 ± 2.59
	Winter	6.46 ± 2.85	5.70 ± 2.90	5.59 ± 2.99	2.68 ± 3.86
Housing solution	Foster home	7.37 ± 3.55	6.73 ± 2.86	6.62 ± 2.67	4.64 ± 2.78
	Foster family	8.86 ± 2.96	6.81 ± 3.18	5.40 ± 2.43	6.79 ± 4.26
	Orphanage	6.09 ± 2.85	5.73 ± 2.79	4.88 ± 2.78	3.57 ± 3.07
BMI Status	Underweight	2.87 ± 0.49	5.07 ± 2.00	5.17 ± 2.03	5.76 ± 5.99
	Normal weight	6.44 ± 2.56	6.08 ± 2.63	5.66 ± 2.39	5.42 ± 2.76
	Overweight	7.75 ± 3.01	6.65 ± 2.64	5.99 ± 2.41	4.18 ± 2.07
	Obese	6.71 ± 3.44	6.28 ± 2.26	7.19 ± 2.81	6.56 ± 2.12

VDD Vitamin D Deficiency, BMI Body Mass Index





compared with those coming from Europe. In our cohort, no children were internationally adopted from Latin America when younger than 1 year of age

Post hoc analyses highlighted that statistically significant differences existed only for those children coming from subtropical areas when compared to children coming from tropical areas. As a matter of fact, children coming from a tropical area tend to have higher VD values than children coming from a subtropical area when comparing the severe VDD status with normal VD status (OR 5.329, 95% CI 2.512-11.41).

Detailed informations about the results above described can be found in Table 5 and Additional file 1.

Even housing solution had a significant relationship with 25(OH)D mean values (p = 0.016) and vitamin D status (p = 0.008). Mean 25(OH)D values in children who were guest of a foster house were 24.21 ng/ml (SD ± 16.13 ng/mL), in children housed in a foster family were 18.79 ng/mL (SD ± 6.72 ng/mL), while in children who were guest of an orphanage were 20.95 ng/mL (SD $\pm 10.92 \text{ ng/mL}$).

Post hoc analyses highlighted how actually the only significant difference can be found comparing children coming from foster homes with children coming from foster families. Children coming from foster families seem to be more at risk of a worst VD status, especially when comparing moderate VDD status with normal VD status (OR 0.137, 95% CI 0.049-0.453).

Detailed data about the results above described can be found in Table 5 and Additional file 1.

Vitamin D mean values and status and their relations with season of blood sampling

Season at blood sampling was significantly related both with 25(OH)D mean values (p < 0.001) and Vitamin D status (p < 0.001). Mean 25(OH)D value in children having their blood drawn in spring was 18.85 ng/mL (SD \pm 10.39 ng/mL), in children undergoing blood draw in summer was 26.82 ng/mL (SD ± 11.08 ng/mL), in children who were firstly evaluated during fall was 22.28 ng/ mL (SD ± 10.10 ng/mL) and in children having their first examination during winter was 18.10 ng/mL (SD ± 11.32 ng/mL).

Post hoc analyses highlighted how during summer the risk of VDD is substanstially decreased, compared to all the other seasons. Interestingly, a statistically significant difference can be found between children observed during spring and children observed during fall, when severe VDD status is compared to mild VDD status. Children observed during fall tend to have a better VD status than children observed during spring (OR 3.319, 95% CI 1.656-6.715).

Table 5 shows Vitamin D status in relation to macroarea, climatic zone of origin, housing solution and season at blood sampling. Detailed data about post hoc analyses can be found in Additional file 1.

Table 5	Vitamin D	status,	geographica	l origin anc	l socia	environment
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		Vitamin D Status				
		Severe VDD (< 10 ng/mL)	Moderate VDD (10–20 ng/mL)	Mild VDD (20–30 ng/mL)	Normal VD (> 30 ng/mL)	p values
Macro-area (n = 859)	Europe	24 (16.1%)	68 (45.6%)	39 (26.2%)	18 (12.1%)	< 0.001
	Latin America	31 (12.1%)	103 (40.2%)	85 (33.2%)	37 (14.5%)	
	Asia	34 (14.2%)	67 (30.1%)	73 (32.7%)	49 (22.0%)	
	Africa	18 (7.8%)	110 (47.6%)	79 (34.2%)	24 (10.4%)	
Climatic zone (n = 859)	Temperate	31 (12.2%)	102 (40.0%)	85 (33.3%)	37 (14.5%)	0.002
	Subtropical	34 (20.5%)	68 (41.0%)	52 (31.3%)	12 (7.2%)	
	Tropical	42 (9.7%)	178 (40.6%)	139 (31.7%)	79 (18.0%)	
Housing solution ($n = 828$)	Foster home	8 (11.4%)	24 (34.4%)	19 (27.1%)	19 (27.1%)	0.008
	Foster family	6 (9.2%)	37 (56.9%)	18 (27.7%)	4 (6.2%)	
	Orphanage	90 (13.0%)	275 (39.7%)	227 (32.8%)	101 (14.5%)	
Season of first blood draw (n = 859)	Spring	29 (15.2%)	94 (49.2%)	50 (26.2%)	18 (9.4%)	< 0.001
	Summer	3 (2.8%)	41 (38.3%)	56 (52.3%)	49 (45.8%)	
	Fall	18 (6.9%)	101 (39.0%)	103 (39.8%)	37 (14.3%)	
	Winter	57 (21.9%)	112 (43.1%)	67 (25.8%)	24 (9.2%)	

 χ^2 -test was used to determine statistically significant relationships among categorical variables

Vitamin D mean values and status and their relations with other factors of the bone metabolism

PTH status was not significantly related to 25(OH)D mean values (p = 0.609) and Vitamin D Status (p = 0.086).

ALP status was not significantly associated with 25(OH)D mean values (p = 0.593), but it was significantly associated with Vitamin D status (p = 0.001). Mean 25(OH)D value in children having a normal ALP was 20.99 ng/mL (SD ± 10.87 ng/mL), while in children having an increased ALP it was 21.48 ng/mL (SD ± 12.30 ng/mL).

Post hoc analysis revealed that this statistically significant difference is only real when comparing severe VDD status with mild VDD status, with children having an increased ALP being at more risk of having a worse VD status (OR 0.4562, 95% CI 0.2737–0.7773).

Detailed data about post hoc analyses can be found in Additional file 1.

Ca status was not significantly related to 25(OH)D mean values (p = 0.338) or Vitamin D status (p = 0.296).

P mean values were significantly associated to 25(OH)D mean values (p = 0.004) and to Vitamin D status (p = 0.012). P status was equally significantly related to 25(OH)D mean values (p = 0.008) and to Vitamin D status (p = 0.014). 25(OH)D mean value in children having a normal P was 21.22 ng/mL (SD ± 11.14 ng/mL), in children having a decreased P was 23.06 ng/mL (SD ± 12.61 ng/mL).

Table 6 shows Vitamin D status in relation to PTH status, ALP status, Ca status and P status.

Biochemical rickets

Only 811 children (94.4%) out of 859 had the complete blood tests panel for the definition of biochemical

rickets. The analysis performed permitted to identify 129 children affected by biochemical rickets as defined in the *"Subjects and Methods"* section. 28 children (21.7%) had severe VDD and an increased ALP, associated with normal or reduced Ca and P. 61 children (47.3%) had moderate VDD and an increased ALP, associated with normal or reduced Ca and P. 40 children () had mild VDD and an increased ALP, associated with normal or reduced Ca and P.

Table 7 resumes details about children affected by biochemical rickets.

Multivariate analysis

We entered the statistically significant variables (p < 0.25) in relation to Vitamin D status (sex, skin color, macroarea of origin, climatic zone of origin, season at first evaluation, housing solutions, ALP status, P status, Age at arrival in Italy, Age at first visit) in a multiple logistic regression model. At the end of the assessment, skin color (p =0.011), season at first blood draw (p < 0.001), housing solution (p < 0.001), ALP status (p = 0.025) and the age at the first blood draw (p < 0.001) had a statistically significant association with Vitamin D Status.

Table 8 shows details about the multiple logistic regression analysis final results. Figure 3 shows the risk of developing hypovitaminosis D during the different seasons when using spring as baseline.

Discussion

Migrant children are a population of great interest nowadays, and their health problems—infectious ones, but also not-infectious ones—are a source of interesting discussions

		Vitamin D Status				
		Severe VDD (< 10 ng/mL)	Moderate VDD (10–20 ng/mL)	Mild VDD (20–30 ng/mL)	Normal VD (> 30 ng/mL)	p values
PTH status (n = 822)	<i>Normal</i> (< 10 pg/mL)	94 (11.8%)	323 (40.6%)	262 (33.0%)	116 (14.6%)	0.086
	Decreased (10–65 pg/mL)	0 (0.0%)	0 (0.0%)	3 (100%)	0 (0.0%)	
	Increased (> 65 pg/mL)	3 (12.5%)	14 (58.3%)	3 (12.5%)	4 (16.7%)	
ALP status (n = 831)	Normal ^a	69 (10.9%)	259 (41.1%)	218 (34.7%)	84 (13.3%)	0.001
	Increased ^a	34 (16.9%)	76 (37.8%)	49 (24.4%)	42 (20.9%)	
Ca status (n = 824)	<i>Normal</i> (8.8–10.8 mg/dL)	100 (12.5%)	324 (40.3%)	261 (32.5%)	84 (13.3%)	0.296
	Decreased (< 8.8 mg/dL)	0 (0.0%)	3 (100%)	0 (0.0%)	0 (0.0%)	
	Increased (> 10.8 mg/dL)	1 (5.6%)	5 (27.8%)	8 (44.4%)	4 (22.2%)	
P status (n = 813)	Normal ^b	84 (11.5%)	293 (40.1%)	249 (34.1%)	105 (14.3%)	0.014
	Decreased ^b	4 (11.1%)	16 (44.5%)	7 (19.4%)	9 (25.0%)	
	Increased ^b	12 (26.1%)	20 (43.5%)	11 (23.9%)	3 (6.5%)	

Table 6 Vitamin D status and its relations with PTH status, ALP status, Ca status and P status

χ²-test was used to determine statistically significant relationships among categorical variables

^a Normal values, depending on the date of the blood draw and the age/sex of the patient, are accurately described within the section "Subjects and Methods"; ^b Normal values, depending on the age of the patient, are accurately described within the section "Subjects and Methods"

for pediatricians all over the world. Internationally adopted children are a controlled sub-population of the migrant one, thus it could offer an insight on a larger population – though with a lot of differences regarding travelling, housing solutions and dietary habits and deprivations.

Vitamin D is an important pro-hormone, classically associated with bone health, but recently found to be involved in immune system and neuro-psychological disorders, and cardiovascular risk. It can be seen how a D hypovitaminosis could put at risk the entire well-being of a child [1–5].

Our study, carried on for nine years, tried to test some of the well-known predictors for the Vitamin D deficiency in the internationally adopted children population and the relationships existing between vitamin D and other factors implied in the bone health, such as parathyroid hormone, alkaline phosphatase, free calcium and phosphates.

From January 2007 to December 2015, 29,866 children arrived in Italy through an international adoption, from

Eastern Europe countries above all, followed by Latin America, Asia and Africa [30]. The cited report offers only details about 2014 and 2015, but it specifies that the percentages reported are similar to those previously observed. During this period, adopted male children were more than the female ones (58.3% vs 41.7%; ratio = 1.4:1), with a mean age at the arrival of 5.9 years (SD not specified).

The population we observed had a similar composition to that of the entire "internationally adopted children" Italian population: male to female ratio was 1.3:1, with a prevalence of children coming from Eastern Europe countries and a mean age at the arrival of 5.31 years (SD \pm 2.92 years).

We did not find a statistical difference of mean 25(OH)D values between sexes. However, a slight statistical significance was found with regards to the Vitamin D status, with a slight higher percentage of females affected by VDD than males. Female children are, as a matter of fact, more interested by severe VDD and mild

 Table 7 Biochemical rickets: number of children affected and 25(OH)D mean values

			P status		
			Normal ^b (25-OH-D mean value ± SD)	Decreased ^b (25-OH-D mean value ± SD)	Total
Vitamin D Status	Severe deficiency + increased ALP	Normal Ca ^c	26 (7.20 ± 1.46 ng/mL)	2 (6.65 ± 0.92 ng/mL)	129
	·····, ·····	Decreased Ca ^c	0	0	
	Moderate deficiency + increased ALP	Normal Ca ^c	59 (15.30 ± 2.93 ng/mL)	1 (15.00 ± NA ng/mL)	
		Decreased Ca ^c	1 (17.70 ± NA ng/mL)	0	
	Mild deficiency + increased ALP	Normal Ca ^c	40 (24.96 ± 2.89 ng/mL)	0	
		Decreased Ca ^c	0	0	

ALP Alkaline phosphatase; Ca Serum calcium; P Serum phosphate; 25-OH-D 25-hydroxylate-Vitamin D

^a Normal values, depending on the date of the blood draw and the age/sex of the patient, are accurately described within the section "Subjects and Methods";

^b Normal values, depending on the age of the patient, are accurately described within the section "Subjects and Methods"

^c Normal values can be found in Table 3 and within the section "Subjects and Methods"

Variable	Value	OR	95% CI	<i>p</i> value	
Skin color	Very fair / fair (ref)				
	Intermediate	1.014	0.590-1.743	0.96	
	Olive / Brown	2.387	1.261-4.519	0.08	
Season at 1st evaluation	Spring (ref)				
	Summer	0.207	0.104-0.410	< 0.001	
	Fall	0.726	0.362-1.455	0.366	
	Winter	1.000	0.485-2.063	1.000	
Housing solution	Foster home (ref)				
	Foster family	0.252	0.125-0.507	< 0.001	
	Orphanage	3.971	1.971-8.001	< 0.001	
ALP status	Normal ^a (ref)				
	Increased ^a	0.560	0.338-0.930	0.025	
Age at 1 st evaluation	Continuous variable	1.238	1.140-1.345	< 0.001	

Table 8 Multiple logistic regression results: showing predictors of developing Vitamin D Deficiency through Odds Ratios (ORs), 95% Confidence Intervals (CIs) and statistical significance of variables (*p* value)

ALP Alkaline phosphatase

^a Normal values are extensively described in the "Subjects and methods" section

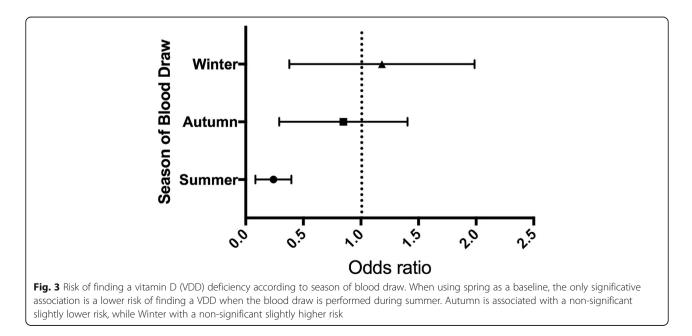
VDD than male children, while male children seem to be more affected by moderate VDD than females (Table 2 and Additional file 1).

A reason for this evidence could be a higher BMI in female children than males, a justification often found by other studies, highlighting how a higher BMI is directly related to the risk of developing VDD, as VD is a lipophilic vitamin and a higher fat mass increase its distribution volume [31-35].

Our results, though, are opposite to those showed by other studies in which VDD was found to be more prevalent among males than in females, or it showed no significant relationship with sex at all [11, 36-39].

In our population, prevalence of VDD is significantly related to the age at the arrival in Italy and to the age at first evaluation. Figure 1 shows the mean age per group of Vitamin D status, highlighting that in our population a higher mean age is related to a worst Vitamin D status.

The negative association of age and serum 25(OH)D in children has already been described in literature. According with Mansbach et al., this might be because of



the difference in duration of sun exposure, while Gustafson et al. sustain that it is caused by the rapid growth spurt, or "catch-up growth" the children reach out in the first year after adoption [7, 40].

Catch-up growth is a process similar to the rapid growth experienced by infants and toddlers, when use of VD supplement is customary for the prevention of rickets [41]. Therefore, during this period internationally adopted need larger amounts of dietary or supplemental iron and VD than those a child of the same age would normally need.

Interestingly, in our population high 25(OH)D concentrations were observed in children aged < 1 year regardless of the area of origin (Fig. 1). This is most likely attributable to the use of fortified formula milk during the period spent in an institution and by pediatricians' increasing awareness of the hypovitaminosis D issue once they arrive in Italy. As reported by other studies, this relation might lead to nullify the expected effects of an adoptee's birth country on VD level [35, 40].

Our study showed a strong association between skin color and Vitamin D Status, that only remained significant for the value "olive-brown skin" after the multivariate analysis (logistic regression). Our data showed that children with olive to brown skin had a statistically significant 2-fold higher risk of having hypovitaminosis D than children with fair skin (OR 2.387, Table 8). These results agree with those reported by an Italian study by Franchi et al. evaluating 1.374 children living in the north of Italy. The study observed that ethnicity was a strong predictor of 25(OH)D levels, reporting a high prevalence of hypovitaminosis D in African, North African, and Indian children living in Italy (74.8, 81.2, 89.7, and 76.0%, respectively) [25]. Other studies reported the same results. A severe 25(OH)D deficiency associated to a proximal myopathy was observed in recent immigrants from Palestine, Pakistan and India to Northern Europe develop due to the limited effect of sunshine and a low dietary VD intake [37]. Hintzpeter [35] showed that Turkish and Arab-Islamic participants of both sexes as well as Asian and African girls were found to be at high risk for VDD, with the highest OR for African girls (7.8) [35]. Similarly, data from NHANES III showed that adolescents of African origin had an increased risk of VDD (OR 8.6) compared with white adolescents [15].

These consistent observations can lead us to conclude that a spontaneous recovery of VD status is unlikely in an adopted child arrived in Italy with VDD, independently from their ethnicity.

In our study *macro area* of origin was significantly associated with VD status (p < 0.001). Basing our observation on percentages alone, VDD seems to have been more common in Africa than in Latin America or Europe and Russian Federation. If statistically significant,

our results would differ from those showed by Chiappini et al., whose study highlighted the highest prevalence of VDD in children coming from the Russian federation [26]. However, post hoc analyses highlighted that the real statistically significant differences are only between children coming from Asia and children coming from Africa. See Additional file 1 and the "results" section for a more detailed description of this difference. Thus, it is not possible to conclude that VDD is more common in children coming from Africa than in children coming from Latin America or Europe and Russian Federation.

As previously said with regards to skin color, it is thought that occasional exposure to sunlight provides most of the VD requirement of the human population [42]. However, skin synthesis of VD may not compensate for the low nutritional intake in Europe, because of the relationship occurring between its latitude and the rake angle of solar radiations. As a matter of fact, Webb et al. [43] demonstrated that the photosynthesis of pre-VD is nearly impossible during winter months at latitudes similar to those encasing Europe. A high prevalence of hypovitaminosis D combined with high PTH levels and lower bone mineral density has recently been reported in Finnish girls and Italian children and adolescents, suggesting that Italian children are at risk for this condition, as well as other European populations [44-46]. Moreover, another Italian study by Vierucci et al., showed that no VD is produced in central Italy during late fall, winter and the beginning of spring because of sun exposure [38, 39].

The prevalence of VDD, even at southern latitudes, was remarkable, supporting other reports suggesting that living in sunny areas of the world does not necessarily protect children from VDD [39, 43, 47–50].

Our work highlighted that the risk of developing hypovitaminosis D was significantly higher in children coming from foster families and orphanages, compared to those coming from foster families. This relationship was still statistically significant after multivariate analysis.

It is a well-demonstrated knownledge that children in foster care experience high rates of child abuse, emotional deprivation, and physical neglect than children living with their families [51–53]. Moreover, it is plausible to think that children in orphanages likely do not spend much time outdoors and that they lack adequate sun exposure. Another reason for the association with VDD could be that as children grow up in institutional care, they shift from a diet of vitamin D–fortified formula milk to cooked food, which may not be fortified with vitamin D.

According to other studies, we observed how the season of blood sampling could be considered a predictor of the Vitamin D Status [26, 54]. As a matter of fact, it is only natural to find out that a pro-hormone which synthesis is related to sun-exposure has higher levels during summer than during winter. It has been shown that even minimal exposure of skin to sunlight, is adequate to reach VD sufficiency in US and Central Europe, and a number of studies demonstrated that both serum concentrations of 25(OH)D and other biomarkers of bone health status increase during summer only to reduce during winter, while PTH behavior is the exact opposite [55–63].

Our data highlighted how drawing a blood sample during summer was a protective factor against hypovitaminosis D, when compared to spring (Fig. 1). OR for summer shows a reduction of the risk of approximately the 80%.

In our study, almost half the children had normal Ca, P, ALP and PTH levels. This data differs from other pediatric studies, which show a percentage way lower [23, 64]. Normal serum calcium works as an inhibitory signal for parathyroids, stopping PTH secretion, as described by Steingrimsdottir and Aggarwal [65, 66].

Our study differs from other studies on the same matter because of the interest taken in showing the relationship existing between vitamin D and parathyroid hormone. Only a very small percentage of our population was affected by hyperparathyroidism, but a significant data was that we could observe elevated PTH levels in children having 25(OH)D levels over 30 ng/mL and normal calcium, a finding deserving further studies: was it a primary hypersecretion or a signal of an altered bone metabolism?

We also tried to identify the PTH inflection point for this population. With this term, other studies indicated the mean 25(OH)D value at which PTH begins to increase. We did not succeed, probably because of the small number of children with an increased PTH included in our population and the extreme variability of values at which PTH was found to increase.

Biochemical rickets affected a significant percentage of our study population. Surprisingly, even children with only mild VDD were affected by it (Table 8). It is though that rickets is confined to children exposed to extreme undernourishment but, as a matter of fact, it can be also observed in western countries, because of the growing epidemia of childhood obesity and the previously discussed relationship of high body mass index and fat mass with VDD [67].

Limits - Strenghts

Our study was limited by the lack of data on dietary intake of calcium and vitamin D. Type of nutrition in the first few months of life can deeply influence values of VD. Some of these infants and toddlers lived with their biological families prior to adoption and might have been breast-fed (low VD intake, limited Ca absorption); however, many international adoptees spent most of their life in orphanages or with foster families and were primarily formula fed (high VD intake, maximized Ca absorption).

Additionally, we could not retrieve any information about sunshine exposure, use of sun protections, medical conditions (i.e renal and/or hepatic failure), use of hypovitaminizing drugs, use of concealing clothing.

However, our study also has some strengths: it shows the irrelevance of PTH as a predictor of VDD, and identifies, through a multivariate analysis, some of the factors a pediatrician cannot miss when evaluating an internationally adopted child for the first time.

Conclusions

The study indicates a widespread global VD insufficiency compared with proposed threshold levels. This high prevalence implies that higher doses than currently recommended would be needed to reach country specific "desirable" levels.

Our results reinforce the importance of an early evaluation of serum 25(OH)D levels in internationally adopted children to start the adequate strategy of VD supplementation/treatment, avoiding further deterioration of VD status, particularly in older, dark skin children arrived in Italy during winter and spring period.

Additional file

Additional file 1: Post hoc analyses. (DOCX 17 kb)

Abbreviations

25(OH)D: 25-hydroxy-vitamin D; ALP: Alkaline phosphatase; Ca: Calcium; P: Phosphorus; PTH: Parathormone; VD: Vitamin D; VDD: Vitamin D deficiency

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Availability of data and materials

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

PV concepted and designed the study; GS, DB and MC wrote the manuscript; DP and VF collected clinical and laboratory data; CDW and MDA analyzed and interpreted the data. All authors read and approved the final manuscript.

Ethics approval and consent to participate

This study was conducted with the approval of the Institutional Boards of Institute of Pediatrics and Department of Woman and Child Health Sciences on 06/10/15 (record n.0415).

Oral informed consent was obtained by parents of the partecipants.

Consent for publication

All authors have read and approved the content, and they agree to submit it for consideration for publication in the journal.

Competing interests

The authors declare that they have no competing interests.

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