

Supplementary Table 1

Model	β	SE	L_CI	U_CI	P	r^2p
M1						$r^2=0.653$
AGE	-0.11	0.020	-0.15	-0.07	1.09×10^{-8}	0.025
GENDER	-1.28	0.035	-1.35	-1.21	1.95×10^{-204}	0.506
TIME DRAWN	-0.46	0.033	-0.52	-0.39	3.22×10^{-40}	0.125
TANNER STAGE	-0.25	0.015	-0.28	-0.22	1.55×10^{-53}	0.165
M1 + HEIGHT + WEIGHT						$r^2=0.666$
AGE	-0.10	0.019	-0.14	-0.07	6.78×10^{-8}	0.022
GENDER	-1.25	0.043	-1.33	-1.16	1.41×10^{-145}	0.394
TIME DRAWN	-0.45	0.033	-0.52	-0.39	3.41×10^{-40}	0.125
TANNER STAGE	-0.23	0.015	-0.26	-0.20	6.88×10^{-47}	0.145
HEIGHT	0.10	0.025	0.05	0.15	4.28×10^{-5}	0.013
WEIGHT	-0.16	0.024	-0.21	-0.12	6.95×10^{-12}	0.035
M2						$r^2=0.666$
AGE	-0.10	0.019	-0.14	-0.07	6.61×10^{-8}	0.022
GENDER	-1.31	0.063	-1.43	-1.18	7.26×10^{-82}	0.244
TIME DRAWN	-0.45	0.033	-0.51	-0.39	5.85×10^{-40}	0.125
TANNER STAGE	-0.23	0.016	-0.26	-0.20	2.33×10^{-44}	0.138
HEIGHT	0.12	0.030	0.06	0.18	6.43×10^{-5}	0.012
F-MASS	-0.11	0.020	-0.15	-0.07	6.32×10^{-8}	0.012
L-MASS	-0.16	0.039	-0.23	-0.08	5.77×10^{-5}	0.022

Table shows regression analyses of different covariates versus CTX in 1325 participants aged 15.5 (580 males, 745 females). M1= adjustment for average age, gender, tanner stage and whether the individual attended the clinic in the morning or afternoon. M1 + HEIGHT + WEIGHT = Model 1 in addition to height and weight; M2 = Model 1 in addition to total-body lean mass and fat mass. n= sample size; SE = standard error; β = SD change CTX per SD change in exposure (excluding categorical variables: gender, time drawn and tanner stage); L_CI = lower 95% confidence estimate of β ; U_CI = upper 95% confidence estimate of β ; P = strength of evidence against the null hypothesis of no association between the outcome and exposure variable; r^2p = partial proportion of the variance in the outcome explained by the exposure in the specified model. r^2 = total proportion of variance in the CTX explained by the all exposures in the specified model

Supplementary Table 2

LOCUS	RSID	POS	GENE	PMID	r^2	EA	ALSPAC (n=3382)			YFS (n=1558)			GOOD (n=938)		
							β^*	SE	P	β^*	SE	P	β^*	SE	P
8q24.12	rs4355801	119993054	TNFRSF11B	19079262 & 18455228	0.90	A	-0.01	0.021	6.41x10 ⁻¹	-0.08	0.036	3.47x10 ⁻²	0.08	0.043	7.17x10 ⁻²
	rs7839059	120045723	TNFRSF11B	23437003	0.42	A	-0.04	0.022	9.58x10 ⁻²	-0.13	0.042	2.14x10 ⁻³	0.03	0.045	5.58x10 ⁻¹
	rs2062375	120046973	TNFRSF11B	20548944	0.97	C	-0.03	0.021	1.47x10 ⁻¹	-0.12	0.037	7.79x10 ⁻⁴	0.06	0.043	1.48x10 ⁻¹
	rs2062377 [□]	120076601	TNFRSF11B	19801982 & 22504420	1.00	A	-0.03	0.021	1.75x10 ⁻¹	-0.10	0.037	9.00x10 ⁻³	0.06	0.043	1.77x10 ⁻¹
	rs6469792	120077552	TNFRSF11B	19079262	0.74	C	-0.04	0.021	3.93x10 ⁻²	-0.10	0.036	3.96x10 ⁻³	0.07	0.043	8.06x10 ⁻²
	rs11995824	120081881	TNFRSF11B	19801982	0.79	G	-0.04	0.021	4.53x10 ⁻²	-0.11	0.036	2.16x10 ⁻³	0.04	0.043	3.07x10 ⁻¹
	rs6469804	120114010	TNFRSF11B	18445777 & 19079262	0.88	A	-0.03	0.021	2.41x10 ⁻¹	-0.09	0.037	1.27x10 ⁻²	0.04	0.043	3.15x10 ⁻¹
	rs6993813	120121419	TNFRSF11B	18445777 & 19079262	0.69	C	-0.04	0.021	8.27x10 ⁻²	-0.11	0.036	3.09x10 ⁻³	0.04	0.043	3.89x10 ⁻¹
13q14.11	rs9533090 [□]	41849449	AKAP11	19801982 & 22504420	1.00	T	0.00	0.021	8.23x10 ⁻¹	0.00	0.036	9.22x10 ⁻¹	0.03	0.045	5.16x10 ⁻¹
	rs9594738	41850145	TNFSF11	18445777 & 19079262	1.00	T	0.00	0.021	8.29x10 ⁻¹	0.00	0.036	9.10x10 ⁻¹	0.03	0.045	5.46x10 ⁻¹
	rs9533093	41859597	TNFSF11	19079262	0.23	T	0.04	0.024	8.60x10 ⁻²	0.06	0.046	1.80x10 ⁻¹	0.06	0.055	2.66x10 ⁻¹
	rs9594759	41930593	TNFSF11	18445777 & 19079262	0.68	T	0.00	0.021	9.71x10 ⁻¹	-0.01	0.037	7.94x10 ⁻¹	0.02	0.045	6.06x10 ⁻¹
	rs1021188 [□]	42014133	TNFSF11	21124946 & 23437003	0.00	C	-0.08	0.027	3.84x10 ⁻³	-0.02	0.056	6.75x10 ⁻¹	-0.06	0.060	2.93x10 ⁻¹
18q21.33	rs884205 [□]	58205837	TNFRSF11A	19801982 22504420	1.00	A	-0.06	0.024	7.22x10 ⁻³	-0.02	0.041	5.62x10 ⁻¹	-0.11	0.051	2.75x10 ⁻²
	rs3018362	58233073	TNFRSF11A	19079262	0.68	A	-0.05	0.022	1.62x10 ⁻²	-0.01	0.037	7.93x10 ⁻¹	-0.08	0.045	7.18x10 ⁻²

Table showing association of previously reported RANK (*TNFRSF11A*), RANKL (*TNFSF11*) and OPG (*TNFRSF11B*) variants with cortical thickness in ALSPAC (n=3382), GOOD (n=938) and Young Finns (n=1558). (POS) = position in the genome based on hg18; (GENE) = closest gene; (PMID) = accession number of the publication in Pubmed which described the association with BMD; (r^2) = the pairwise LD estimate in CEU populations between the SNP in bold and all other SNPs in that locus; (EA) = effect allele; (β^*) = effect size; (SE) = standard error of β^* ; (P) = P-value; (PMID) = accession number of the publication in Pubmed which described the association with BMD. *Effect estimates expressed as adjusted SD per copy of the effect allele (EA). Note: rs9533090[□] is found upstream of *TNFSF11*, but is closest to *AKAP11*. □ - denotes the variants which were used to generate allele scores (i.e. independent signals).

Supplementary Table 3

LOCUS	RSID	POS	GENE	PMID	r ²	EA	ALSPAC (n=3382)			YFS (n=1558)			GOOD (n=938)		
							β^*	SE	P	β^*	SE	P	β^*	SE	P
8q24.12	rs4355801	119993054	TNFRSF11B	19079262 & 18455228	0.90	A	-0.05	0.018	2.74x10 ⁻³	-0.13	0.036	2.89x10 ⁻⁴	-0.12	0.044	5.11x10 ⁻³
	rs7839059	120045723	TNFRSF11B	23437003	0.42	A	-0.07	0.019	1.04x10 ⁻⁴	-0.14	0.041	4.88x10 ⁻⁴	-0.17	0.046	2.07x10 ⁻⁴
	rs2062375	120046973	TNFRSF11B	20548944	0.97	C	-0.07	0.018	2.97x10 ⁻⁴	-0.16	0.036	1.16x10 ⁻⁵	-0.14	0.044	2.21x10 ⁻³
	rs2062377 [□]	120076601	TNFRSF11B	19801982 & 22504420	1.00	A	-0.06	0.018	1.27x10 ⁻³	-0.16	0.037	2.17x10 ⁻⁵	-0.11	0.044	1.14x10 ⁻²
	rs6469792	120077552	TNFRSF11B	19079262	0.74	C	-0.05	0.018	5.92x10 ⁻³	-0.14	0.036	1.45x10 ⁻⁴	-0.10	0.044	1.81x10 ⁻²
	rs11995824	120081881	TNFRSF11B	19801982	0.79	G	-0.05	0.018	2.66x10 ⁻³	-0.14	0.036	7.79x10 ⁻⁵	-0.14	0.045	1.39x10 ⁻³
	rs6469804	120114010	TNFRSF11B	18445777 & 19079262	0.88	A	-0.05	0.018	3.25x10 ⁻³	-0.15	0.037	2.68x10 ⁻⁵	-0.12	0.044	8.67x10 ⁻³
	rs6993813	120121419	TNFRSF11B	18445777 & 19079262	0.69	C	-0.05	0.018	5.38x10 ⁻³	-0.15	0.036	5.40x10 ⁻⁵	-0.14	0.044	1.30x10 ⁻³
13q14.11	rs9533090 [□]	41849449	AKAP11	19801982 & 22504420	1.00	T	-0.02	0.018	2.09x10 ⁻¹	-0.11	0.035	1.48x10 ⁻³	0.01	0.046	8.98x10 ⁻¹
	rs9594738	41850145	TNFSF11	18445777 & 19079262	1.00	T	-0.02	0.018	2.09x10 ⁻¹	-0.11	0.035	1.47x10 ⁻³	-0.01	0.046	9.01x10 ⁻¹
	rs9533093	41859597	TNFSF11	19079262	0.23	T	-0.01	0.021	7.59x10 ⁻¹	-0.04	0.046	3.56x10 ⁻¹	-0.01	0.055	8.66x10 ⁻¹
	rs9594759	41930593	TNFSF11	18445777 & 19079262	0.68	T	-0.01	0.018	6.20x10 ⁻¹	0.01	0.037	8.67x10 ⁻¹	0.04	0.046	3.79x10 ⁻¹
	rs1021188 [□]	42014133	TNFSF11	21124946 & 23437003	0.00	C	-0.12	0.024	1.97x10 ⁻⁷	-0.16	0.056	3.10x10 ⁻³	-0.30	0.063	1.65x10 ⁻⁶
18q21.33	rs884205 [□]	58205837	TNFRSF11A	19801982 & 22504420	1.00	A	-0.09	0.021	3.05x10 ⁻⁵	0.02	0.040	5.95x10 ⁻¹	-0.04	0.051	4.90x10 ⁻¹
	rs3018362	58233073	TNFRSF11A	19079262	0.68	A	-0.07	0.019	1.93x10 ⁻⁴	-0.02	0.036	5.35x10 ⁻¹	-0.06	0.046	2.07x10 ⁻¹

Table showing association of previously reported RANK (*TNFRSF11A*), RANKL (*TNFSF11*) and OPG (*TNFRSF11B*) variants with cortical BMD in ALSPAC (n=3382), GOOD (n=938) and Young Finns (n=1558). (POS) = position in the genome based on hg18; (GENE) = closest gene; (PMID) = accession number of the publication in Pubmed which described the association with BMD; (r²) = the pairwise LD estimate in CEU populations between the SNP in bold and all other SNPs in that locus; (EA) = effect allele; (β^*) = effect size; (SE) = standard error of β^* ; (P) = P-value; (PMID) = accession number of the publication in Pubmed which described the association with BMD. *Effect estimates expressed as adjusted SD per copy of the effect allele (EA). Note: rs9533090[□] is found upstream of *TNFSF11*, but is closest to *AKAP11*. □ - denotes the variants which were used to generate allele scores (i.e. independent signals).

Supplementary Table 4

LOCUS	RSID	POS	GENE	PMID	r^2	EA	ALSPAC (n=3382)			YFS (n=1558)			GOOD (n=938)		
							β^*	SE	P	β^*	SE	P	β^*	SE	P
8q24.12	rs4355801	119993054	TNFRSF11B	19079262 & 18455228	0.90	A	0.03	0.016	6.18x10 ⁻²	0.12	0.036	1.08x10⁻³	0.02	0.035	6.70x10 ⁻¹
	rs7839059	120045723	TNFRSF11B	23437003	0.42	A	0.02	0.017	1.94x10 ⁻¹	0.05	0.041	2.44x10 ⁻¹	0.03	0.037	3.58x10 ⁻¹
	rs2062375	120046973	TNFRSF11B	20548944	0.97	C	0.03	0.016	6.34x10 ⁻²	0.08	0.036	1.96x10⁻²	-0.01	0.035	8.37x10 ⁻¹
	rs2062377 [□]	120076601	TNFRSF11B	19801982 & 22504420	1.00	A	0.02	0.016	2.74x10 ⁻¹	0.09	0.037	1.46x10⁻²	-0.01	0.035	8.34x10 ⁻¹
	rs6469792	120077552	TNFRSF11B	19079262	0.74	C	0.01	0.016	5.88x10 ⁻¹	0.08	0.036	1.90x10⁻²	0.01	0.035	8.47x10 ⁻¹
	rs11995824	120081881	TNFRSF11B	19801982	0.79	G	0.01	0.016	5.34x10 ⁻¹	0.08	0.036	2.22x10⁻²	0.01	0.036	6.77x10 ⁻¹
	rs6469804	120114010	TNFRSF11B	18445777 & 19079262	0.88	A	0.02	0.016	3.03x10 ⁻¹	0.09	0.037	9.92x10⁻³	0.00	0.035	9.38x10 ⁻¹
13q14.11	rs9533090 [□]	41849449	AKAP11	19801982 & 22504420	1.00	T	-0.02	0.016	3.36x10 ⁻¹	0.08	0.036	1.71x10⁻²	0.00	0.037	9.42x10 ⁻¹
	rs9594738	41850145	TNFSF11	18445777 & 19079262	1.00	T	-0.02	0.016	3.37x10 ⁻¹	0.08	0.035	1.95x10⁻²	0.01	0.037	8.46x10 ⁻¹
	rs9533093	41859597	TNFSF11	19079262	0.23	T	-0.02	0.018	2.60x10 ⁻¹	0.07	0.046	1.31x10 ⁻¹	-0.01	0.044	8.67x10 ⁻¹
	rs9594759	41930593	TNFSF11	18445777 & 19079262	0.68	T	-0.01	0.016	5.20x10 ⁻¹	0.03	0.036	3.75x10 ⁻¹	-0.04	0.037	3.06x10 ⁻¹
	rs1021188 [□]	42014133	TNFSF11	21124946 & 23437003	0.00	C	0.05	0.020	9.21x10⁻³	0.04	0.056	5.27x10 ⁻¹	0.06	0.050	2.08x10 ⁻¹
18q21.33	rs884205 [□]	58205837	TNFRSF11A	19801982 & 22504420	1.00	A	0.05	0.018	7.03x10⁻³	0.04	0.040	3.26x10 ⁻¹	0.02	0.041	6.53x10 ⁻¹
	rs3018362	58233073	TNFRSF11A	19079262	0.68	A	0.04	0.017	2.91x10⁻²	0.02	0.036	6.15x10 ⁻¹	0.05	0.037	1.83x10 ⁻¹

Table showing association of previously reported RANK (*TNFRSF11A*), RANKL (*TNFSF11*) and OPG (*TNFRSF11B*) variants with Periosteal Circumference in ALSPAC (n=3382), GOOD (n=938) and Young Finns (n=1558). (POS) = position in the genome based on hg18; (GENE) = closest gene; (PMID) = accession number of the publication in Pubmed which described the association with BMD; (r^2) = the pairwise LD estimate in CEU populations between the SNP in bold and all other SNPs in that locus; (EA) = effect allele; (β^*) = effect size; (SE) = standard error of β^* ; (P) = P-value; (PMID) = accession number of the publication in Pubmed which described the association with BMD. *Effect estimates expressed as adjusted SD per copy of the effect allele (EA). Note: rs9533090[□] is found upstream of *TNFSF11*, but is closest to *AKAP11*. □ - denotes the variants which were used to generate allele scores (i.e. independent signals).