## Supplementary Table 1. Analysis based on <u>controls only</u> (ATBC, CPS-II, CLUE II, and PLCO combined; n=1,312)

SNPª	Risk allele (freq)	Chr	Location <sup>b</sup>	Gene neighborhood	P-value <sup>c,d</sup>	Beta <sup>e</sup>	S.E.
rs2282679	C (0.26)	4	72827247	GC	$1.4 \times 10^{-11}$	-0.39	0.06
rs3829251	A (0.19)	11	70872207	NADSYN1/DHRC7	0.11	-0.10	0.06
rs6599638	G (0.49)	10	124694139	C10orf88	$4.0 \times 10^{-3}$	-0.15	0.05
rs2060793	A (0.41)	11	14866810	CYP2R1	0.25	0.06	0.05

## FOOTNOTES TO SUPPLEMENTARY TABLES 1 AND 2

- a. SNP identifier based on NCBI dbSNP
- b. Chromosomal location based on NCBI Human Genome Build 35 coordinates
- c. The result of 1 df test based on linear regression, using an additive model, after square root transformation of 25-hydroxy vitamin D levels adjusted for age, vitamin D assay batch, study, case-control status, sex, body mass index, season of blood collection, vitamin D supplement intake, dietary vitamin D intake, region/latitude, and chosen eigenvectors to control population stratification.
- d. The result of 1 df test based on logistic regression, using an additive model,— adjusted for age, vitamin D assay batch, study, case-control status, sex, body mass index, season of blood collection, vitamin D supplement intake, dietary vitamin D intake, region/latitude, and chosen eigenvectors to control population stratification.
- e. The regression beta and standard error were based on the square root scale.

## Supplementary Table 2. Binary outcome for vitamin D clinical deficiency (<25 nmol/L) (ATBC, CPS-II, CLUE II, PLCO combined; n=2,907)

SNPª	Risk allele (freq)	Chr	Location <sup>b</sup>	Gene Neighborhood	P-value <sup>c</sup>	OR
rs2282679	C (0.26)	4	72827247	GC	$6.5 \times 10^{-9}$	1.83
rs3829251	A (0.19)	11	70872207	NADSYN1/DHRC7	0.11	1.18
rs6599638	G (0.49)	10	124694139	C10orf88	$8.1 \times 10^{-2}$	1.17
rs2060793	A (0.41)	11	14866810	CYP2R1	0.21	0.89

Joint model: *GC* rs2282679 + *NADSYN1/DHCR7* rs3829251 + *C10orf88* rs6599638 + *CYPR1* rs2060793. Odds ratio of having one copy of high risk allele at each of the four loci is 2.97 (95% CI 2.00 - 4.39). The joint effect from the four SNPs is significant  $(P = 3.2 \times 10^{-8})$ .