Common variants at *MS4A4/MS4A6E*, *CD2AP*, *CD33* and *EPHA1* are associated with late-onset Alzheimer's disease

The Alzheimer Disease Genetics Consortium (ADGC) performed a genome-wide association study of late-onset Alzheimer disease using a three-stage design consisting of a discovery stage (stage 1) and two replication stages (stages 2 and 3). Both joint analysis and meta-analysis approaches were used. We obtained genome-wide significant results at *MS4A4A* (rs4938933; stages 1 and 2, meta-analysis P (P_M) = 1.7×10^{-9}, joint analysis P (P_J) = 1.7×10^{-9}; stages 1, 2 and 3, P_M = 8.2×10^{-12}), *CD2AP* (rs9349407; stages 1, 2 and 3, P_M = 8.6×10^{-9}), *EPHA1* (rs11767557; stages 1, 2 and 3, P_M = 6.0×10^{-10}) and *CD33* (rs3865444; stages 1, 2 and 3, P_M = 1.6×10^{-9}). We also replicated previous associations at *CR1* (rs6701713; P_M = 4.6×10^{-10}, P_J = 5.2×10^{-11}), *CLU* (rs1532278; P_M = 8.3×10^{-8}, P_J = 1.9×10^{-8}), *BIN1* (rs7561528; P_M = 4.0×10^{-14}, P_J = 5.2×10^{-14}) and *PICALM* (rs561655; P_M = 7.0×10^{-11}, P_J = 1.0×10^{-10}), but not at *EXOC3L2*, to late-onset Alzheimer's disease susceptibility[1–3].

Alzheimer's disease is a neurodegenerative disorder affecting more than 13% of individuals aged 65 years and older and 30–50% of individuals aged 80 years and older[4,5]. Early work identified mutations in *APP*, *PSEN1* and *PSEN2* that cause early-onset autosomal dominant Alzheimer's disease[6–9] and variants in *APOE* that affect late-onset Alzheimer's disease (LOAD) susceptibility[10]. Recent genome-wide association studies (GWAS) identified variants in *CR1*, *CLU*, *PICALM* and *BIN1* as LOAD susceptibility loci[1–3]. However, because LOAD heritability estimates are high ($h^2 \approx$ 60–80%)[11], much of the genetic contribution to this condition remains unknown.

To identify genetic variants associated with risk for Alzheimer's disease, the ADGC assembled a discovery dataset (stage 1, 8,309 individuals with LOAD (cases) and 7,366 cognitively normal elders (CNEs) as controls) using data from eight cohorts and a ninth newly assembled cohort from the 29 National Institute on Aging (NIA)-funded Alzheimer Disease Centers (ADCs) (**Supplementary Tables 1,2** and **Supplementary Note**), with data coordinated by the National Alzheimer Coordinating Center (NACC) and samples coordinated by the National Cell Repository for Alzheimer Disease (NCRAD). For the stage 2 replication, we used four additional datasets and additional samples from the ADCs (3,531 LOAD cases and 3,565 CNEs). The stage 3 replication used the results of association analyses provided by three other consortia, including 6,992 LOAD cases and 24,666 mixed-age controls, reported in a companion manuscript[12]. For stages 1 and 2, we used both a meta-analysis approach that integrated results

from the association analyses of individual datasets and a joint analysis approach in which genotype data from each study were pooled. The latter method has improved power over the meta-analysis in the absence of between-study heterogeneity[13] and has a more direct correction for confounding sampling bias[14]. We were limited to meta-analysis for stage 3 analyses.

Because the cohorts were genotyped using different platforms, we used imputation to generate a common set of 2,324,889 SNPs. We applied uniform stringent quality control measures to all datasets to remove low-quality and redundant samples and problematic SNPs (**Supplementary Tables 3,4** and Online Methods). We performed an association analysis assuming an additive model on the log odds ratio scale with adjustment for population substructure using logistic regression for case-control data and generalized estimating equations (GEE) with a logistic model for family data. We combined results from individual datasets in the meta-analysis using the inverse variance method, applying a genomic control to each dataset. We performed the joint analysis using GEE and incorporated terms to adjust for population substructure and site-specific effects (Online Methods). For both approaches, we also examined an extended model of covariate adjustment that adjusted for age (age at onset or death in cases and age at exam or death in controls), sex and number of *APOE* ε4 alleles (0, 1 or 2). Genomic inflation factors (λ) for both the discovery meta-analysis and the joint analysis and extended models were less than 1.05, indicating that there was not substantial inflation of the test statistics (**Supplementary Table 3** and **Supplementary Fig. 1**). Association findings from the meta-analysis and joint analysis were comparable.

In stage 1, the strongest signal was from the *APOE* region (rs4420638; P_M = 1.1×10^{-266}, P_J = 1.3×10^{-253}; **Supplementary Table 5**). Excluding the *APOE* region, SNPs at nine distinct loci yielded P_M or $P_J \leq 10^{-6}$ (**Table 1**; all SNPs with P < 10^{-4} are shown in **Supplementary Table 5**). SNPs from these nine loci were carried forward to stage 2. Five of these loci had not previously been associated with LOAD at a genome-wide significance level of $P \leq 5.0 \times 10^{-8}$ (loci in *MS4A*, *EPHA1*, *CD33*, *ARID5B* and *CD2AP*). Because the companion study[12] identified SNPs at *ABCA7* to be within a new LOAD locus, we included *ABCA7*-region SNPs in our stage 2 analysis and provided our results to researchers from that study. For all loci listed in **Table 1**, we did not detect evidence for effect heterogeneity (**Supplementary Fig. 2**). One newly associated locus (in *MS4A*) was significant in the stage 1+2 analysis. Four other loci approached but

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Table 1 Genome-wide association results for LOAD in the ADGC stage 1 and stage 2 datasets

SNP	Chr.:Mb	Nearest gene	MA	MAF	# SNPs	ADGC discovery (stage 1)				ADGC replication (stage 2)				Combined analysis (stage 1+2)			
						OR_M (95% CI)	P_M	OR_J (95% CI)	P_J	OR_M (95% CI)	P_M	OR_J (95% CI)	P_J	OR_M (95% CI)	P_M	OR_J (95% CI)	P_J
rs6701713	1:207.8	CR1[a]	A	0.20	7	1.18 1.11–1.25	1.4×10^{-8}	1.19 1.12–1.26	3.5×10^{-9}	1.13 1.04–1.23	0.004	1.13 1.04–1.24	0.004	1.16 1.11–1.22	4.6×10^{-10}	1.17 1.12–1.23	5.2×10^{-11}
rs7561528	2:127.9	BIN1[a]	A	0.35	10	1.18 1.13–1.24	2.9×10^{-11}	1.18 1.12–1.24	7.7×10^{-11}	1.15 1.07–1.24	1.4×10^{-4}	1.15 1.07–1.24	1.0×10^{-4}	1.17 1.13–1.22	4.2×10^{-14}	1.17 1.12–1.22	5.2×10^{-14}
rs9349407	6:47.5	CD2AP	C	0.27	1	1.14 1.08–1.21	1.2×10^{-6}	1.14 1.08–1.20	5.3×10^{-6}	1.07 0.98–1.17	0.118	1.08 0.99–1.18	0.074	1.12 1.07–1.18	1.0×10^{-6}	1.12 1.07–1.17	2.1×10^{-6}
rs11767557	7:143.1	EPHA1[b]	C	0.19	1	0.85 0.80–0.90	7.3×10^{-8}	0.84 0.79–0.89	3.1×10^{-8}	0.94 0.86–1.03	0.169	0.93 0.85–1.02	0.133	0.87 0.83–0.92	2.4×10^{-7}	0.87 0.83–0.91	4.9×10^{-8}
rs1532278	8:27.5	CLU[a]	T	0.36	2	0.90 0.85–0.95	5.6×10^{-5}	0.89 0.85–0.94	2.0×10^{-5}	0.87 0.81–0.94	2.6×10^{-4}	0.87 0.81–0.94	2.7×10^{-4}	0.89 0.85–0.93	8.3×10^{-8}	0.89 0.85–0.92	1.9×10^{-8}
rs2588969	10:63.6	ARID5B	A	0.37	0	0.88 0.84–0.93	1.1×10^{-6}	0.88 0.85–0.94	6.9×10^{-7}	1.05 0.97–1.13	0.234	1.05 0.98–1.13	0.189	0.93 0.89–0.97	0.001	0.93 0.89–0.97	7.7×10^{-4}
rs4938933	11:60.0	MS4A4A	C	0.39	22	0.88 0.84–0.92	5.2×10^{-8}	0.87 0.83–0.92	4.5×10^{-8}	0.90 0.84–0.97	0.005	0.90 0.84–0.97	0.004	0.88 0.85–0.92	1.7×10^{-9}	0.88 0.85–0.92	1.7×10^{-9}
rs561655	11:85.8	PICALM[a]	G	0.34	36	0.88 0.84–0.92	1.2×10^{-7}	0.88 0.84–0.93	4.6×10^{-7}	0.86 0.80–0.92	8.4×10^{-5}	0.86 0.80–0.92	3.7×10^{-5}	0.87 0.84–0.91	7.0×10^{-11}	0.87 0.84–0.91	1.0×10^{-10}
rs3752246	19:1.1	ABCA7[d]	G	0.19	2	1.16 1.08–1.24	1.0×10^{-5}	1.15 1.08–1.23	1.9×10^{-5}	1.13 1.03–1.24	0.012	1.13 1.03–1.25	0.009	1.15 1.09–1.21	5.8×10^{-7}	1.15 1.09–1.21	5.0×10^{-7}
rs3865444	19:51.7	CD33[c]	A	0.30	1	0.88 0.84–0.93	8.2×10^{-7}	0.88 0.84–0.93	1.9×10^{-6}	0.91 0.85–0.99	0.021	0.92 0.85–0.99	0.029	0.89 0.86–0.93	1.1×10^{-7}	0.89 0.86–0.93	2.0×10^{-7}

Chr.:Mb, chromosome:position (build 19); MA, minor allele; MAF, minor allele frequency; # SNPs, the number of SNPs for which $P \leq 1 \times 10^{-6}$ in meta-analysis from the combined analysis in stage 1+2; OR_M, odds ratio in meta-analysis; OR_J, odds ratio in joint analysis; P_J, P value in joint analysis; P_M, P value in meta-analysis. Genes with previous case-control genome-wide statistically significant associations: CR1 (ref. 3), CLU (refs. 1, 3), PICALM (ref. 1), BIN1 (ref. 2). Association signals represent SNPs with the strongest associations within each locus showing $P \leq 10^{-6}$ in the stage 1 dataset or in or near previously reported genes, excluding the APOE region (Supplementary Table 5). [a]Genes with previously published case-control association signals at $P \leq 5.0 \times 10^{-8}$. [b]This locus did not meet this level of statistical significance[2]. [c]The locus previously reported in a family-based association study as genome-wide significant[15]. [d]Locus identified in the companion study[12] with genome-wide significant evidence for association.

did not reach genome-wide significance in the stage 1+2 analyses and were carried forward to stage 3. For three of these (loci in *CD33*, *EPHA1* and *CD2AP*), the stage 3 analysis strengthened the evidence for association. However, stage 2 and 3 results did not support the stage 1 results for *ARID5B* (**Table 2**).

Our stage 1+2 analysis identified the *MS4A* gene cluster as a new LOAD locus ($P_M = 1.7 \times 10^{-9}$, $P_J = 1.7 \times 10^{-9}$) (**Table 1** and **Fig. 1a**). The minor allele (minor allele frequency (MAF) = 0.39) was protective and had identical odds ratios (ORs) in both the meta-analysis and the joint analysis (OR_M and $OR_J = 0.88$, 95% CI 0.85–0.92). In the stage 1+2 analysis, other SNPs gave smaller P values when compared to discovery SNP, rs4938933, and the most significant SNP was rs4939338 ($P_M = 2.6 \times 10^{-11}$, $P_J = 4.6 \times 10^{-11}$, OR_M and $OR_J = 0.87$, 95% CI 0.84–0.91) (**Supplementary Table 5**). In the accompanying manuscript[12], genome-wide significant results were also obtained at the *MS4A* locus (rs670139; $P_M = 5.0 \times 10^{-12}$) using an independent sample. In a combined analysis of ADGC results and those from the companion study[12], the evidence for this locus at rs4938933 increased to $P_M = 8.2 \times 10^{-12}$ (**Table 2**; $OR_M = 0.89$, 95% CI 0.87–0.92; **Fig. 1a**).

SNPs in the *CD2AP* locus also met our stage 1 criteria for additional analysis (**Fig. 1b**). Stage 2 data modestly strengthened this association, but the results did not reach genome-wide significance. The stage 3 analysis yielded a genome-wide significant result for rs9349407 ($P_M = 8.6 \times 10^{-9}$), which identified *CD2AP* as a new LOAD locus. The minor allele (MAF = 0.27) at this SNP increased the risk for LOAD ($OR_M = 1.11$, 95% CI 1.07–1.15) (**Table 2** and **Fig. 1b**).

Another locus studied further in stages 2 and 3 centered on *EPHA1*. Previous work provided suggestive evidence that this is a LOAD risk locus, although the associations did not previously reach genome-wide significance ($P = 1.7 \times 10^{-6}$)[2]. Here, results from stages 1 and 2 for rs11767557, located in the promoter region of *EPHA1*, reached genome-wide significance in the joint analysis. The addition of stage 3 results increased the evidence for association ($P_M = 6.0 \times 10^{-10}$; **Table 2** and **Fig. 1c**). The minor allele (MAF = 0.19) for this SNP is protective ($OR_M = 0.90$, 95% CI 0.86–0.93). We observed no evidence for heterogeneity at this locus (**Supplementary Fig. 2d**; heterogeneity $P = 0.58$).

In stages 1 and 2, we also obtained strong evidence for association for SNPs in *CD33*, a gene located approximately 6 Mb from *APOE*, but our results did not reach genome-wide significance. The addition of stage 3 data confirmed that *CD33* is a LOAD risk locus (rs3865444; stages 1, 2 and 3, $P_M = 1.6 \times 10^{-9}$). The minor allele (MAF = 0.30) for this SNP is protective ($OR_M = 0.91$, 95% CI 0.88–0.93; **Tables 1,2** and **Fig. 1d**). A single SNP (rs3826656) in the 5 region of *CD33* was previously reported as a genome-wide significant Alzheimer's disease–related locus using a family-based approach ($P = 6.6 \times 10^{-6}$) (ref. 15). We were unable to replicate this finding ($P_M = 0.73$, $P_J = 0.39$ in the stage 1 analysis for rs3826656). Though rs3826656 is only 1,348 bp from our top SNP (rs3865444), these two sites have only weak linkage disequilibrium (LD) ($r^2 = 0.13$).

Researchers in the accompanying study[12] report highly significant evidence for the association of an *ABCA7* SNP, rs3764650, with LOAD ($P_M = 4.5 \times 10^{-17}$), from a meta-analysis that included data from our study. In our stage 1+2 analysis, we obtained suggestive evidence for association with the *ABCA7* SNP rs3752246 ($P_M = 5.8 \times 10^{-7}$, $P_J = 5.0 \times 10^{-7}$), which is a missense variant (p.Gly1527Ala) that may alter the function of the ABCA7 protein (see **Supplementary Table 6** for functional SNPs in LD with SNPs yielding P_M or $P_J < 10^{-4}$).

Our stage 1+2 analyses also confirmed the association of previously reported loci (in *BIN1*, *CR1*, *CLU* and *PICALM*) with LOAD (**Table 1**). For each locus, supporting data were P values that were less than $P = 5.0 \times 10^{-8}$ in one or both types of analysis.

Table 2 Meta-analysis of stage 1+2 with stage 3 (CHARGE/GERAD/EADI1 Consortia[2]) GWAS results

Gene:SNP	Cases	Controls	Total	OR_M (95% CI)	P_M	OR_J (95% CI)	P_J
CD2AP: rs9349407							
ADGC	11,840	10,931	22,771	1.12 (1.07–1.18)	1.0×10^{-6}	1.12 (1.07–1.17)	2.1×10^{-6}
External	6,922	18,896	25,818	1.09 (1.03–1.15)	0.002	—	—
ADGC + External	18,762	29,827	48,589	1.11 (1.07–1.15)	8.6×10^{-9}	—	—
EPHA1: rs11767557							
ADGC	11,840	10,931	22,771	0.87 (0.83–0.92)	2.4×10^{-7}	0.87 (0.83–0.91)	4.9×10^{-8}
External	6,922	24,666	31,588	0.91 (0.87–0.96)	2.9×10^{-4}	—	—
ADGC + External	18,762	35,597	54,359	0.90 (0.86–0.93)	6.0×10^{-10}	—	—
ARID5B: rs2588969							
ADGC	11,840	10,931	22,771	0.93 (0.89–0.97)	0.001	0.93 (0.89–0.97)	7.8×10^{-4}
External	6,922	18,896	25,818	1.06 (1.01–1.11)	0.018	—	—
ADGC + External	18,762	29,827	48,589	0.99 (0.95–1.02)	0.362	—	—
MS4A4A: rs4938933							
ADGC	11,840	10,931	22,771	0.88 (0.85–0.92)	1.7×10^{-9}	0.88 (0.85–0.92)	1.7×10^{-9}
External	6,922	18,896	25,818	0.92 (0.88–0.97)	5.4×10^{-4}	—	—
ADGC + External	18,762	29,827	48,589	0.89 (0.87–0.92)	8.2×10^{-12}	—	—
CD33: rs3865444							
ADGC	11,840	10,931	22,771	0.89 (0.86–0.93)	1.1×10^{-7}	0.89 (0.86–0.93)	2.0×10^{-7}
External	6,922	18,896	25,818	0.92 (0.88–0.97)	0.002	—	—
ADGC + External	18,762	29,827	48,589	0.91 (0.88–0.93)	1.6×10^{-9}	—	—

Meta-analysis using an external replication case-control sample (stage 3) for SNPs from previously unidentified loci at which associations did not exceed the genome-wide statistical significance threshold ($P = 5.0 \times 10^{-8}$) in the ADGC meta-analysis (stage 1+2). Results for *MS4A* are also included to show association results from the ADGC and accompanying manuscript[12]. The external replication dataset is described in the accompanying paper[12] and includes the stage 1 discovery sample and the CHARGE sample[2] but does not include results from the TGEN, ADNI and MAYO cohorts (**Supplementary Tables 1** and **2**).

We also examined SNPs with statistically significant GWAS results reported by others (*GAB2* (ref. 16), *PCDH11X*[17], *GOLM1* (ref. 18) and *MTHFD1L*[19]; **Supplementary Table 7**). Stage 1 data were used, except for *PCDH11X*, for which stage 1+2 data were used because Affymetrix platforms do not contain the appropriate SNP. Only SNPs in the *APOE*, *CR1*, *PICALM* and *BIN1* loci had $P < 10^{-6}$. For *MTHFD1L*[19], we obtained modest independent association evidence at rs11754661 (previously reported $P = 4.7 \times 10^{-8}$; this study $OR_M = 1.16$, 95% CI 1.04–1.29, $P_M = 0.006$, $OR_J = 1.19$, 95% CI 1.08–1.32, $P_J = 7.5 \times 10^{-4}$). For the remaining sites, we obtained only nominal evidence ($P < 0.05$) or no evidence of association. For the *GAB2* locus[16] at rs10793294 (previously reported $P = 1.60 \times 10^{-7}$), we obtained nominal statistically significant results ($P_M = 0.017$, $P_J = 0.029$). The association for rs5984894 in the

PCDH11X locus[17] (previously reported $P = 3.9 \times 10^{-12}$) did not replicate ($P_M = 0.89$, $P_J = 0.26$). Likewise, findings at *GOLM1* (ref. 18) for rs10868366 (previously reported $P = 2.40 \times 10^{-4}$) did not replicate ($P_M = 0.71$, $P_J = 0.62$). Another gene consistently implicated in LOAD is *SORL1* (ref. 20), where at rs3781835 (previously reported $P = 0.006$), we obtained modest evidence for association ($OR_M = 0.72$, 95% CI 0.60–0.86, $P_M = 2.9 \times 10^{-4}$, $OR_J = 0.78$, 95% CI 0.59–0.86, $P_J = 3.8 \times 10^{-4}$).

We examined the influence of the *APOE* ε4 allele on the loci in **Table 1** stratified by and in interactions with *APOE* ε4 allele carrier status. After adjustment, all loci had similar effect sizes as the unadjusted analyses, with some loci showing a modest reduction in statistical significance. We previously reported evidence for a *PICALM-APOE* (ref. 21) interaction using a dataset that largely overlaps with the stage 1
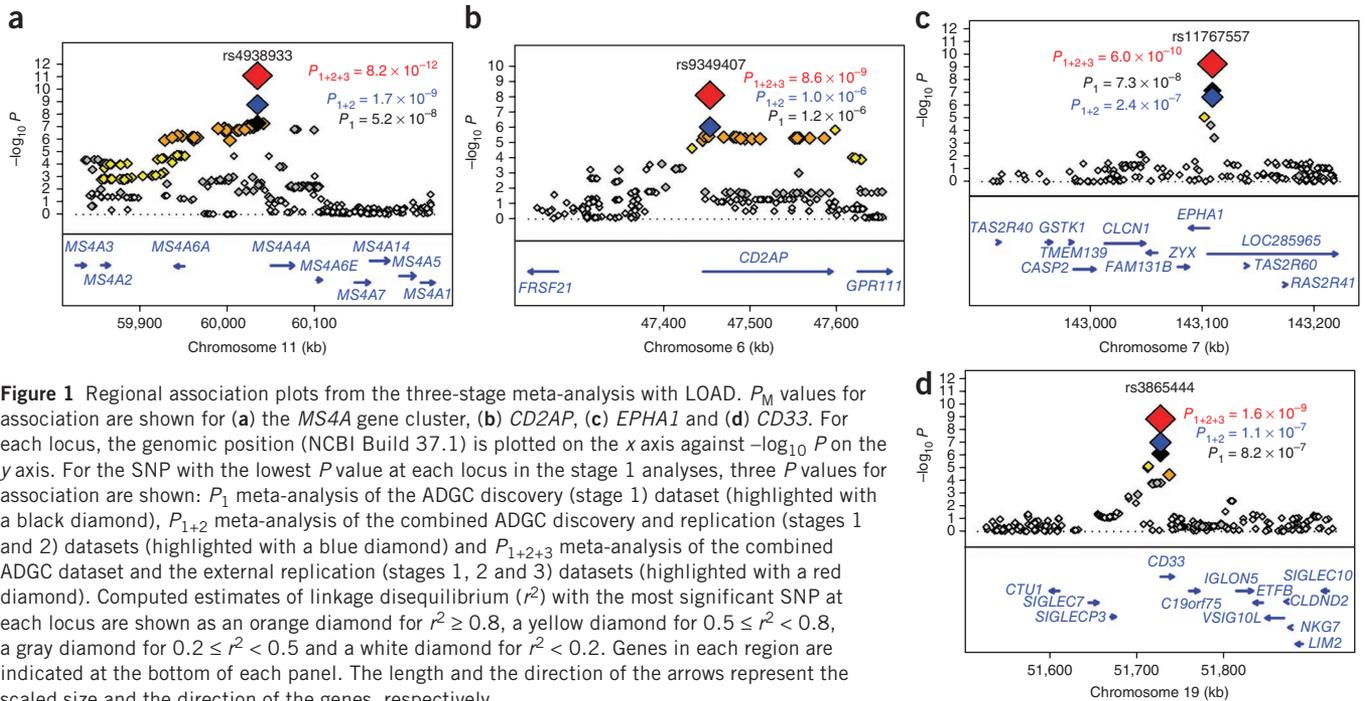
Figure 1 Regional association plots from the three-stage meta-analysis with LOAD. P_M values for association are shown for (**a**) the *MS4A* gene cluster, (**b**) *CD2AP*, (**c**) *EPHA1* and (**d**) *CD33*. For each locus, the genomic position (NCBI Build 37.1) is plotted on the x axis against $-\log_{10} P$ on the y axis. For the SNP with the lowest P value at each locus in the stage 1 analyses, three P values for association are shown: P_1 meta-analysis of the ADGC discovery (stage 1) dataset (highlighted with a black diamond), P_{1+2} meta-analysis of the combined ADGC discovery and replication (stages 1 and 2) datasets (highlighted with a blue diamond) and P_{1+2+3} meta-analysis of the combined ADGC dataset and the external replication (stages 1, 2 and 3) datasets (highlighted with a red diamond). Computed estimates of linkage disequilibrium (r^2) with the most significant SNP at each locus are shown as an orange diamond for $r^2 \geq 0.8$, a yellow diamond for $0.5 \leq r^2 < 0.8$, a gray diamond for $0.2 \leq r^2 < 0.5$ and a white diamond for $r^2 < 0.2$. Genes in each region are indicated at the bottom of each panel. The length and the direction of the arrows represent the scaled size and the direction of the genes, respectively.

dataset used here. However, using the stage 1+2 data, we did not replicate this finding or see evidence of *APOE* genotype interactions with the loci listed in **Table 1** (data not shown).

Previous work reported an association between LOAD and the chromosome 19 SNP rs597668, which is located 7.2 kb proximal to *EXOC3L2* and 296 kb distal of *APOE*[2]. Although we did observe a signal for this SNP (stage 1, $P_M = 1.5 \times 10^{-9}$, $P_J = 7.7 \times 10^{-10}$) and other SNPs in the *EXOC2L3-MARK4* region, the evidence was completely extinguished for all SNPs after adjustment for *APOE* (Online Methods and **Supplementary Table 8**), suggesting that signal in this region is from *APOE*.

Our observation of genome-wide significant associations at *MS4A4A*, *CD2AP*, *EPHA1* and *CD33* extends our understanding of the genetic architecture of LOAD and confirms the emerging consensus that common genetic variation plays an important role in the etiology of LOAD. With our findings and those in the companion study[12], there are now ten LOAD susceptibility loci (in *APOE*, *CR1*, *CLU*, *PICALM*, *BIN1*, *EPHA1*, *MS4A*, *CD33*, *CD2AP* and *ABCA7*). Examining the amount of genetic effect attributable to these candidate genes, the most strongly associated SNPs at each locus other than that in *APOE* had population attributable fractions between 2.72% and 5.97% (**Supplementary Table 9**), with a cumulative population-attributable fraction for non-*APOE* loci estimated to be as much as 35%; however, these estimates may vary widely between studies[22], and the actual effect sizes are likely to be much smaller than those estimated here because of the 'winner's curse'. Also, the results do not account for interaction among loci and are not derived from appropriate population-based samples.

A recent review of GWAS[23] noted that risk alleles with small effect sizes (0.80 < OR < 1.2) likely exist for complex diseases such as LOAD but remain undetected, even with thousands of samples, because of insufficient power[24]. Our discovery dataset (stage 1, 8,309 cases and 7,366 controls) was well powered to detect associations exceeding the statistical significance threshold of $P < 10^{-6}$ (**Supplementary Table 9**). If there are many loci of more modest effects, some, but not all, will likely be detected in any one study. This likely explains the genome-wide statistical significance for the *ABCA7* locus in the accompanying manuscript[12], which reached only modest statistical significance in our dataset (rs3752246; $P_M = 1.0 \times 10^{-5}$, $P_J = 1.9 \times 10^{-5}$). Finding additional LOAD loci will require larger studies with increased depth of genotyping to test for the effects of both common and rare variants.

URLs. The Alzheimer Disease Genetics Consortium (ADGC), http://alois.med.upenn.edu/adgc/about/overview.html; ADNI database, http://www.loni.ucla.edu/ADNI; ADNI investigators, http://www.loni.ucla.edu/ADNI/Collaboration/ADNI_Manuscript_Citations.pdf; *APOE* Genotyping kit from TIB MOLBIOL, http://www.roche-as.es/logs/LightMix%C2%AE_40-0445-16_ApoE-112-158_V080904.pdf; PLINK, http://pngu.mgh.harvard.edu/~purcell/plink/; PREST, http://utstat.toronto.edu/sun/Software/Prest/; MACH, http://www.sph.umich.edu/csg/abecasis/mach/; EIGENSTRAT, http://genepath.med.harvard.edu/~reich/EIGENSTRAT.htm; The R Project for Statistical Computing, http://www.r-project.org/; Package *GWAF* in R, http://cran.r-project.org/web/packages/GWAF/index.html; Package *gee* in R, http://cran.r-project.org/web/packages/gee/index.html; UCSC Genome Browser, http://genome.ucsc.edu/; METAL, http://www.sph.umich.edu/csg/abecasis/Metal/; FUGUE, http://www.sph.umich.edu/csg/abecasis/fugue/.

METHODS
Methods and any associated references are available in the online version of the paper at http://www.nature.com/naturegenetics/.

Note: Supplementary information is available on the Nature Genetics website.

AUTHOR CONTRIBUTIONS
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ONLINE METHODS

Subjects. A full description of study cohorts is provided in the **Supplementary Note** and **Supplementary Tables 1** and **2**.

Covariate data. Age of onset data was available from some cohorts (Alzheimer's Disease Center (ADC), Translational Genomics Research Institute series 2 (TGEN2), National Institute on Aging Late-onset AD (NIA-LOAD), Multi-Institutional Research on Alzheimer's Genetic Epidemiology (MIRAGE), Adult Changes in Thought (ACT), Multi-Site Collaborative Study for Genotype-Phenotype Associations in Alzheimer's Disease (GenADA), University of Pittsburgh (UP) and the Rush University Religious Orders Study/Memory and Aging Project (ROS/MAP)), whereas for others, only age at ascertainment (Washington University (WU) and ADNI), age at diagnosis (Mayo Clinic (MAYO)), or a combination of both age at ascertainment and age at death was available (a subset of autopsy-confirmed samples in the University of Miami/Vanderbilt University/Mt. Sinai School of Medicine (UM/VU/MSSM) cohort). For subjects with autopsy-confirmed diagnosis and no clinical diagnosis, the age at diagnosis was equated to the age at death. For all studies, the age used for CNEs was the age of last exam or age at death. Case and CNE subjects with age at symptom onset or age at death less than 60 were excluded from the analysis. We restricted our association analyses to individuals of European ancestry because there were insufficient subjects from non–European-ancestry groups to obtain meaningful results.

Genotyping, data cleaning and imputation. Genotypes were from either Illumina or Affymetrix high-density SNP microarrays (**Supplementary Table 3**). Genotype data were cleaned by applying minimum call rates (95% and 98%) and minimum minor allele frequencies (0.02 and 0.01) for cohorts genotyped on Affymetrix and Illumina chips, respectively. SNPs not in Hardy-Weinberg equilibrium ($P < 10^{-6}$) were excluded. Subjects where the gender was mis-specified were identified by analysis of X-chromosome SNPs using PLINK[25]. For cohorts genotyped on multiple chips (MIRAGE and UM/UV/MSSM), genotype and sample quality thresholds were applied within subsets of individuals genotyped on each chip. For all other cohorts, quality thresholds were applied per cohort. Relationships among individuals in the family-based cohorts (MIRAGE and NIALOAD) were confirmed by pairwise genome-wide estimates of proportion identity-by-descent (IBD) using PREST software[26]. Any discrepancies identified were reviewed in light of available clinical and pedigree data to determine the most likely relationship consistent with a proportion of IBD, and any remaining scenarios were excluded from analysis. Latent relatedness in the case-control cohorts was identified by proportion IBD using PLINK software[25,27]. Both of each pair of identical samples by IBD ($\hat{\pi} > 0.99$) were dropped, and one subject was selected from each related pair ($0.4 \geq \hat{\pi} > 0.90$), prioritizing non-missing case or non-missing control status and then higher call rate in selection. Duplicate enrollments among studies (**Supplementary Table 4**) were identified using proportion of IBD in a genotyped dataset including all cohorts where pairs with $\hat{\pi} > 0.95$ were considered duplicate enrollments. Duplicates with discordant case or control status by study were dropped from both studies, and those with concordant status were included in only one cohort and selected according to a predetermined priority list of cohorts which considered genotype data, phenotype data and the type of cohort. Genome-wide imputation was performed per cohort using MACH software[28] with HapMap phase 2 (release 22) CEPH Utah pedigree (CEU) reference haplotypes and genotype data passing quality control as inference. Imputation quality was determined as R^2 and only SNPs imputed with $R^2 \geq 0.50$ were included in the analysis.

APOE genotyping. APOE genotypes were determined for the ADC, ACT, NIA-LOAD, UM/VU/MSSM, MAYO and GenADA cohorts using SNPs rs7412 and rs429358; for the MIRAGE cohort using the Roche Diagnostics LightCycler 480 instrument (Roche Diagnostics)[29] LightMix Kit ApoE C112R R158 (TIB MOLBIOL); for TGEN2, ADNI, UP and WU cohorts by pyrosequencing[30] or restriction fragment length polymorphism analysis[31,32]; and for ROSMAP by high-throughput sequencing of codons 112 and 158 in APOE by Agencourt Bioscience Corporation.

Meta-analysis. Presence of intra-study population substructure was evaluated separately by cohort in a two-step process that first removed outliers before estimating population substructure within the remaining population. For the first step, either the STRUCTURE software package[33,34] (UM/VU/MSSM and MIRAGE) or the 'smartpca' script in EIGENSTRAT[35] (remaining cohorts) was used to remove outliers and/or confirm self-reported ethnicity after filtering to remove SNPs in pairwise LD. In the second step, we used EIGENSTRAT[35], often a second time, to estimate principal component loadings for inclusion in association analysis. For each study, the first two, three or four estimated principal components were identified for inclusion as covariates in association analysis (**Supplementary Table 3**). Outlier detection for the ADC, TGEN2, GenADA, ACT, ADNI, ROS/MAP, OHSU, UP, WU and MAYO cohorts was evaluated by comparison to the HapMap 3 CEPH (CEU) population. EIGENSTRAT analyses of family cohort data (NIA-LOAD and MIRAGE) used a sample of unrelated individuals to fit principal components after outliers with respect to European-American ancestry were removed.

Genotyped and imputed SNP data passing quality control were tested for association with Alzheimer's disease in each dataset using logistic generalized linear model (GLM) for case-control analysis and logistic generalized estimating equations (GEE) for family-based cohorts in R[36–38]. All analyses assumed an additive genetic model, coding genotyped SNPs by the number of minor alleles (0, 1 or 2) and imputed SNPs by the posterior probability of the minor allele (range 0–2). Primary association analyses were adjusted for population substructure (baseline model).

SNP association results for each dataset were meta-analyzed using the inverse variance method implemented in the software package METAL[39]. The meta-analysis P value was estimated by the summarized test statistic after applying a genomic control within each individual study. Heterogeneity among odds ratios in the meta-analysis was assessed using the Cochran's Q and I^2 statistics[40,41].

Regional association plots were prepared for the most strongly associated SNPs in *CR1*, *BIN1*, *CD2AP*, *EPHA1*, *CLU*, *MS4A4A/MS4A6A*, *PICALM*, *ABCA7* and *CD33* using the gene locations from UCSC Genome browser (hg19, GRCh37, Feb 2009 release) and SNP locations from the corresponding dbSNP build 131. Estimates of LD were calculated with the FUGUE software[42] using HapMap phase 2 (release 24, CEU) genotype data and build 131 SNP positions. Forest plots of study-specific effects and analysis results are presented for the same set of SNPs using the 'rmeta' package in R.

Joint analysis. Testing for population substructure across studies was performed in a combined dataset using the set of SNPs genotyped in all study cohorts. After filtering SNPs with pairwise LD (r^2) <0.20, 31,310 SNPs were evaluated using EIGENSTRAT. The top three principal components from EIGENSTRAT were used as covariates in the joint analysis for association in addition to an adjustment for site-specific effects using dummy variables for each cohort. SNP associations with Alzheimer's disease affection status were examined in a pooled analysis of subjects from all cohorts, excluding SNPs missing from one or more individual dataset or with genotypes available on fewer than 98% of individuals overall. In total, 2,312,972 directly genotyped or imputed SNPs common to all datasets were tested for association in 8,309 cases and 7,366 CNEs, including 3,489 individuals in family datasets using GEE analyses in R. Joint analyses of the baseline model, full model and models evaluating robustness to *APOE* included as covariates the principal components from inter-study and intra-study population substructure and a dummy covariate for cohort-specific effects. Genomic inflation factors for the discovery joint analysis in the basic and extended models of covariate adjustment were 1.05 and 1.04, respectively (**Supplementary Table 3**), which were similar to those from meta-analysis.

Secondary analysis. Association results in regions yielding at least one SNP with $P < 10^{-6}$ (follow-up SNPs) were further evaluated for robustness to *APOE* ε4 carrier status in analyses stratified according to presence or absence of *APOE* ε4 and an interaction analysis including effects for SNP, *APOE* ε4 and their interaction. In addition, we examined the *EXOC3L2* region in chromosome 19 previously reported as independent of *APOE* genotype[2] in a full model including covariates for age at onset or age at last exam, gender and the dosage of *APOE* ε4 alleles.

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Internal and external replication analyses. SNPs attaining a $P \leq 1 \times 10^{-6}$ for association with LOAD in the discovery cohort were evaluated in five independent datasets (ADC3, OHSU, MAYO, ROS/MAP and UP) consisting of 3,531 cases and 3,565 CNEs using the same analytical approaches as described above. Replication was performed using both meta-analysis and joint analysis. The datasets included in discovery and replication analyses are summarized in **Supplementary Tables 1** and **2**. Following internal replication, an external replication cohort was sought to evaluate the most strongly associated SNP in each of four newly identified genes (*CD2AP* (rs9349407), *EPHA1* (rs11767557), *ARID5B* (rs2588969) and *CD33* (rs3865444)) for which results did not met genome-wide significance ($P_M > 5 \times 10^{-8}$, $P_J > 5 \times 10^{-8}$) in the combined discovery and replication datasets (stage 1+2). We obtained summarized results from five independent external datasets generously provided by the Genetic and Environmental Risk in Alzheimer's Disease (GERAD) Consortium[1], the European Alzheimer's Disease Initiative (EADI) Consortium[3] and the Cohorts for Heart and Aging Research in Genomic Epidemiology (CHARGE) Consortium[2], and these were used for combined stage 1, 2 and 3 meta analysis. After removing subjects recognized as part of the ADGC cohorts[12], the sample included 6,922 Alzheimer's disease cases and 24,666 controls. These datasets were analyzed using meta-analysis as described above for the stage 1 and 2 datasets. Results from stages 1, 2 and 3 were likewise assessed by meta-analysis as described above.

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