

Genome-wide association study identifies variants at *CLU* and *CR1* associated with Alzheimer's disease

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The gene encoding apolipoprotein E (*APOE*) on chromosome 19 is the only confirmed susceptibility locus for late-onset Alzheimer's disease. To identify other risk loci, we conducted a large genome-wide association study of 2,032 individuals from France with Alzheimer's disease (cases) and 5,328 controls. Markers outside *APOE* with suggestive evidence of association ($P < 10^{-5}$) were examined in collections from Belgium, Finland, Italy and Spain totaling 3,978 Alzheimer's disease cases and 3,297 controls. Two loci gave replicated evidence of association: one within *CLU* (also called *APOJ*), encoding clusterin or apolipoprotein J, on chromosome 8 (rs11136000, OR = 0.86, 95% CI 0.81–0.90, $P = 7.5 \times 10^{-9}$ for combined data) and the other within *CR1*, encoding the complement component (3b/4b) receptor 1, on chromosome 1 (rs6656401, OR = 1.21, 95% CI 1.14–1.29, $P = 3.7 \times 10^{-9}$ for combined data). Previous biological studies support roles of *CLU* and *CR1* in the clearance of β amyloid (A β) peptide, the principal constituent of amyloid plaques, which are one of the major brain lesions of individuals with Alzheimer's disease.

Alzheimer's disease is a neurological disorder primarily affecting the elderly that manifests through memory disorders, cognitive decline and loss of autonomy. Two principal types of neuropathologic lesions are observed: (i) neurofibrillary degeneration resulting from the intraneuronal accumulation of hyperphosphorylated Tau proteins and (ii) amyloid deposits resulting from the extracellular accumulation of amyloid plaques, which are primarily composed of A β peptides. Currently, the processes leading to the formation of these lesions and their combined association with Alzheimer's disease are not adequately understood¹.

Genetic studies have provided significant insights into the molecular basis of Alzheimer's disease. Rare hereditary early-onset forms of the disease have been linked to mutations in three different genes: *APP*, encoding amyloid precursor protein, on chromosome 21; *PS1*, encoding presenilin 1, on chromosome 14; and *PS2*, encoding presenilin 2, on chromosome 1 (ref. 2). These mutations, however, explain less than 1% of all cases of Alzheimer's disease, whereas the vast majority (especially for late-onset forms of the disease) have other, more complex genetic determinants³.

Genetic studies have led to the consistent identification of the $\epsilon 4$ allele of *APOE* as a susceptibility locus for late-onset Alzheimer's disease⁴. Twin studies suggest that genes may have a role in more than 60% of Alzheimer's disease susceptibility⁵ and that *APOE* may account for as much as 50% of this genetic susceptibility⁶. More than 550 other genes have been proposed as candidates for Alzheimer's disease susceptibility, but thus far none have been confirmed to have a role in Alzheimer's disease pathogenesis⁷.

As with other multifactorial diseases, this knowledge gap has motivated more comprehensive investigations using genome-wide association studies (GWAS). The first GWAS of case-control Alzheimer's disease data collections have examined a relatively small number of cases (<1,000)⁸⁻¹². Similar to studies done on other multifactorial disorders, these GWAS have shown that, except in the case of *APOE*, larger samples will be necessary to locate common genetic factors of Alzheimer's disease. Here, we report results from a large two-stage GWAS of late-onset Alzheimer's disease.

In the first stage of this study, we undertook a GWA analysis of 537,029 SNPs in 2,032 French Alzheimer's disease cases and 5,328 French controls. Patients with probable Alzheimer's disease were ascertained by neurologists. Individuals without symptoms of

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Table 1 Association of SNPs at the *CLU* locus with Alzheimer's disease in the Stage 1 and Stage 2 samples

	rs2279590						
	N		MAF		HWE	Association test	
	Cases	Controls	Cases	Controls		OR (95% CI)	P value
Stage 1	2,025	5,328	0.36	0.41	3.1×10^{-1}	0.83 (0.77–0.90)	1.0×10^{-6}
Stage 2	3,803	3,097	0.38	0.41	4.9×10^{-1}	0.88 (0.81–0.95)	8.2×10^{-4}
Belgium	1,071	505	0.38	0.41	3.0×10^{-1}	0.82 (0.69–0.98)	3.1×10^{-2}
Finland	587	645	0.40	0.42	4.3×10^{-1}	0.92 (0.78–1.09)	3.5×10^{-1}
Italy	1,410	1,206	0.38	0.41	3.4×10^{-1}	0.87 (0.77–0.97)	1.3×10^{-2}
Spain	738	806	0.38	0.40	5.6×10^{-1}	0.91 (0.79–1.06)	2.4×10^{-1}
Stage 1 & 2	5,828	8,425	0.37	0.41	2.9×10^{-1}	0.86 (0.82–0.91)	8.9×10^{-9}
APOE $\epsilon 4$ carriers	3,060	1,714	0.36	0.40	7.5×10^{-1}	0.81 (0.74–0.89)	1.9×10^{-5}
APOE $\epsilon 4$ non carriers	2,727	6,697	0.39	0.41	1.8×10^{-1}	0.92 (0.85–0.98)	1.3×10^{-2}
	rs11136000						
Stage 1	2,016	5,266	0.35	0.39	6.0×10^{-1}	0.83 (0.77–0.90)	1.5×10^{-6}
Stage 2	3,775	3,154	0.35	0.38	2.6×10^{-1}	0.88 (0.81–0.95)	8.8×10^{-4}
Belgium	987	467	0.35	0.37	5.2×10^{-2}	0.80 (0.66–0.97)	2.2×10^{-2}
Finland	596	640	0.38	0.41	2.5×10^{-1}	0.92 (0.78–1.08)	3.1×10^{-1}
Italy	1,454	1,241	0.35	0.38	6.0×10^{-1}	0.88 (0.80–0.99)	2.8×10^{-2}
Spain	738	806	0.35	0.37	6.1×10^{-1}	0.91 (0.79–1.06)	2.4×10^{-1}
Stage 1 + 2	5,791	8,420	0.35	0.38	2.7×10^{-1}	0.86 (0.81–0.90)	7.5×10^{-9}
APOE $\epsilon 4$ carriers	3,053	1,707	0.36	0.41	9.0×10^{-1}	0.81 (0.74–0.88)	2.7×10^{-5}
APOE $\epsilon 4$ non-carriers	2,693	6,699	0.39	0.41	1.9×10^{-1}	0.91 (0.85–0.97)	7.0×10^{-3}
	rs9331888						
Stage 1	2,025	5,328	0.31	0.28	8.9×10^{-1}	1.19 (1.11–1.30)	1.8×10^{-5}
Stage 2	3,862	3,180	0.31	0.29	7.9×10^{-1}	1.12 (1.04–1.21)	2.9×10^{-3}
Belgium	1,072	501	0.29	0.28	9.3×10^{-1}	1.16 (0.96–1.40)	1.2×10^{-1}
Finland	586	638	0.38	0.37	1.3×10^{-1}	1.09 (0.92–1.29)	3.2×10^{-1}
Italy	1,474	1,241	0.30	0.26	9.9×10^{-1}	1.22 (1.07–1.39)	4.5×10^{-3}
Spain	730	800	0.29	0.28	5.4×10^{-1}	1.06 (0.90–1.24)	4.8×10^{-1}
Stage 1 + 2	5,887	8,508	0.30	0.28	3.3×10^{-1}	1.16 (1.10–1.23)	9.4×10^{-8}
APOE $\epsilon 4$ carriers	3,098	1,723	0.32	0.29	8.6×10^{-1}	1.21 (1.10–1.33)	7.8×10^{-5}
APOE $\epsilon 4$ non-carriers	2,748	6,770	0.30	0.28	3.1×10^{-1}	1.09 (1.02–1.18)	1.8×10^{-2}

P values and ORs with the associated 95% CI have been calculated under an additive model using logistic regression models adjusted for age, gender and centers when necessary. MAF, minor allele frequency; HWE, P value for the test of Hardy-Weinberg equilibrium in controls.

dementia from French Three-City (3C) prospective population-based cohort were obtained as controls (**Supplementary Note**). Samples were genotyped with Illumina Human 610-Quad BeadChip and subjected to standard quality control procedures. The resulting GWA data were then analyzed with logistic regression taking into account sex and age and using principal components to adjust for possible population stratification. The genomic control parameter was 1.20 before this adjustment but 1.04 afterward. Comparison of the observed and expected χ^2 distributions (**Supplementary Fig. 1**) did not indicate substantial inflation of the test statistics after adjustment. We carried out additional tests to establish the robustness of the statistical results as described in the online methods section.

Several *APOE*-linked SNPs gave strong evidence of disease association (**Supplementary Table 1**). Outside of *APOE*, one marker (rs11136000) within *CLU* on chromosome 8p21-p12 showed significance ($P = 9.0 \times 10^{-8}$) in the association test. This slightly surpassed the criteria for genome-wide significance as evaluated with a conservative Bonferroni correction ($P < 9.3 \times 10^{-8}$). We observed markers in several chromosome regions with suggestive evidence of association ($P < 10^{-5}$) as shown in a Manhattan plot (**Supplementary Fig. 2**). The results of the entire GWAS are available online. Finally, we imputed genotypes using the HapMap CEU samples to increase the number of SNPs examined in these regions. The genotyped and imputed markers from the regions that gave $P < 10^{-5}$ are shown in **Supplementary Table 2**.

In the second stage, we sought replication of association by genotyping a selection of markers at the loci identified in **Supplementary Table 2** in additional collections totaling 3,978 probable Alzheimer's disease cases and 3,297 controls obtained from Belgium, Finland, Italy and Spain (**Supplementary Table 3** and **Supplementary Note**). Genotyping of the second-stage samples was performed using either Taqman or Sequenom assays. Data were analyzed using logistic regression under an additive genetic model taking into account sex and age at diagnosis (cases) or at confirmation of the absence of dementia (controls). Five SNPs at two loci showed association, with P values ranging from 1.6×10^{-2} to 8.2×10^{-4} . The first of these loci encompasses *CLU* on 8p21-p12, and the second spans the gene encoding complement component (3b/4b) receptor 1 (*CR1*) on 1q32.

At *CLU*, three markers (rs2279590, rs11136000, rs9331888) showed statistically significant association with Alzheimer's disease in the replication collections, even with a conservative Bonferroni correction for all 11 SNPs tested in stage 2 ($P < 4.5 \times 10^{-3}$, **Table 1**). The odds ratios were statistically homogeneous across the stage 1 and 2 study collections (**Supplementary Table 4**). We found strong evidence for association in the combined GWA and replication datasets taking into account the sample origin in the logistic regression, with two markers exceeding the criterion for genome-wide significance ($P < 9.3 \times 10^{-8}$). For the marker showing the strongest evidence of association (rs11136000), the odds ratio for the minor

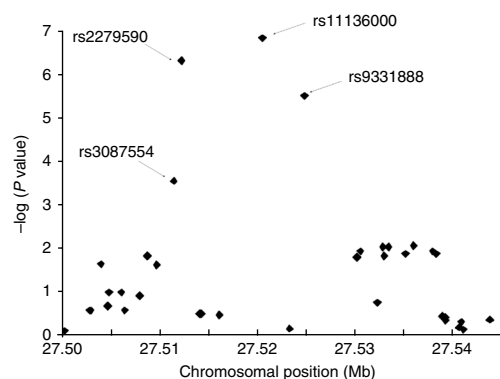


Figure 1 Schematic overview of *CLU* and LD patterns at the *CLU* locus. *P* values for association of SNPs (resulted from imputation and with minor allele frequency (MAF) >5%) encompassing the *CLU* locus with Alzheimer's disease risk under an additive model with adjustment for age and gender are plotted against physical distance. The LD plot is shown for SNPs at the *CLU* locus in controls (stage 1 data in Haploview 4.0, solid spine haplotype block definition, r^2 color scheme).

association with Alzheimer's disease in the replication collections ($P = 8.2 \times 10^{-4}$, **Table 3**). There was no significant heterogeneity by origin (**Supplementary Table 4**), and the odds ratio and *P* values in the combined data were 1.21 (95% CI 1.14–1.29) and 3.7×10^{-9} . We found similar results when using the Mantel-Haenszel method (**Supplementary Table 5**). At this locus, we also detected a statistical interaction with *APOE* $\epsilon 4$ status and risk of disease ($P = 9.6 \times 10^{-3}$) with significant association in both carriers and non-carriers, but with a more significant association in the former. Although the association of the second marker tested at this locus (rs3818361) only showed a suggestive significance overall in the stage 2 collections (OR = 1.11, 95% CI 1.02–1.22, $P = 1.6 \times 10^{-2}$), there was evidence of association of this SNP in the *APOE* $\epsilon 4$ carriers in stage 1, stage 2 and combined samples (respectively, OR = 1.38, 95% CI 1.19–1.60, $P = 2.3 \times 10^{-5}$; OR = 1.29, 95% CI 1.08–1.56, $P = 5.4 \times 10^{-3}$; OR = 1.34, 95% CI 1.20–1.50, $P = 2.9 \times 10^{-7}$). The genotyped markers define two principal haplotypes that account for 97.8% of the observations at the *CRI* locus, and a third haplotype has an estimated frequency of 1.2% in the combined control population (**Table 4**). The odds ratio was highest for the AA haplotype compared to the GG haplotype (OR = 1.22, $P = 3.1 \times 10^{-10}$ for the combined samples).

The association results for *CLU* are supported by a recent meta-analysis of linkage studies of Alzheimer's disease, in which a region of 15.1 Mb on chromosome 8p that encompasses *CLU* had the strongest evidence genome wide for linkage¹³. The biological functions of the *CLU* markers associated with Alzheimer's disease, and of other markers from the region that may be in LD with them, is presently unknown. However, *CLU* is the only described gene located within the association region, and several lines of evidence suggest that it is a strong candidate for involvement in the disease.

APOE and *CLU* are the most abundantly expressed apolipoproteins in the central nervous system^{14,15}, with strong analogies in terms of possible impact on the Alzheimer's disease pathophysiological process. Like *APOE*, *CLU* is present in amyloid plaques^{16,17} and can bind $A\beta$ ^{18,19}. Consistent with this, several experiments in transgenic PDAPP mice lacking *CLU* have shown that this protein may strongly influence $A\beta$ fibrillogenesis and increase $A\beta$ neurotoxicity *in vivo*²⁰. It has also been proposed that *CLU* may participate in $A\beta$ clearance from the brain across the blood-brain barrier, mainly of the highly pathogenic $A\beta 42$ peptide^{21,22}, which is a function similar to that of *APOE*^{23–25}.

In addition, we observed an association between Alzheimer's disease risk and markers spanning *CRI*, the gene that encodes for the main receptor of the complement C3b protein²⁶. The biological function of these *CRI* markers is unknown. The LD block containing these markers

allele was 0.86 (95% CI 0.81–0.90, $P = 7.5 \times 10^{-9}$). Similar results were obtained using a Mantel-Haenszel statistic for the combined analysis (**Supplementary Table 5**).

We detected a statistical interaction between the *APOE* $\epsilon 4$ status and the *CLU* SNPs (ranging from 3.0×10^{-2} to 5.2×10^{-2} from the SNP tested). For rs11136000, although the association was significant in both $\epsilon 4$ carriers and non-carriers, it was more significant in carriers (**Table 1**).

The three *CLU* locus markers replicated in stage 2 are within a linkage disequilibrium (LD) block that encompasses only the *CLU* gene (**Fig. 1**). They define three common haplotypes (frequency >2%) that together account for 98.2% of the observations in stage 1 controls. Compared to the most frequent haplotype, TTC, the other two most frequent haplotypes were all associated with a statistically significant increased disease risk, with similar odd ratios in the stage 1 and stage 2 collections (**Table 2**). The odds ratio was highest for the CCG haplotype compared to the TTC haplotype (OR = 1.22, $P = 5.6 \times 10^{-10}$ for the combined samples).

A second locus of potential interest lies within an LD block that encompasses *CRI* on 1q32 (**Fig. 2**). We tested two SNPs at this locus in the second stage, and one (rs6656401) showed evidence of

Table 2 Association results for haplotypes at the *CLU* locus

Haplotypes	Stage 1				Stage 2				Stage 1 + 2			
	Cases	Controls	OR (95% CI)	<i>P</i> value	Cases	Controls	OR (95% CI)	<i>P</i> value	Cases	Controls	OR (95% CI)	<i>P</i> value
TTC	0.344	0.388	Ref.	–	0.343	0.372	Ref.	–	0.344	0.382	Ref.	–
CCC	0.334	0.329	1.14 (1.04–1.24)	3.0×10^{-3}	0.338	0.327	1.10 (1.01–1.21)	3.3×10^{-2}	0.336	0.328	1.12 (1.06–1.20)	8.0×10^{-5}
CCG	0.302	0.265	1.28 (1.17–1.41)	1.5×10^{-7}	0.277	0.258	1.19 (1.09–1.31)	1.5×10^{-4}	0.286	0.263	1.22 (1.14–1.29)	5.6×10^{-10}

The results have been calculated using the THEIAS program with adjustment for age, gender and center (see Online Methods). The *P* values for the global association were 1.8×10^{-6} , 2.9×10^{-3} and 2.7×10^{-9} , respectively, for the data from stage 1, stage 2 and stage 1 and 2 combined. The markers as ordered from left to right (5' to 3') are rs2279590, rs11136000 and rs9331888. Minor alleles are underlined.

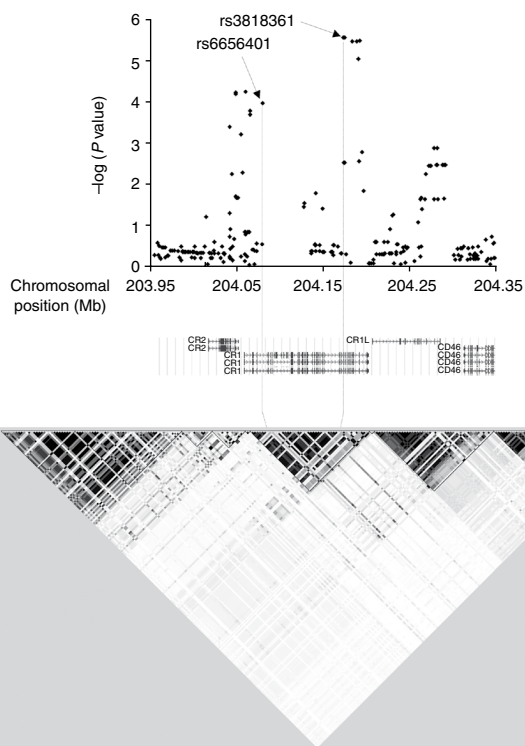


Figure 2 Schematic overview of *CR1* and LD patterns at the *CR1* locus. *P* values for association of SNPs (resulted from imputation and with *MAF* >5%) encompassing the *CR1* locus under an additive model with adjustment for age and gender are plotted against physical distance. The LD plot shown is for SNPs at the *CR1* locus in controls (stage 1 data in Haploview 4.0, solid spine haplotype block definition, r^2 color scheme).

Such a mechanism has been proposed to participate in A β clearance for several reasons: (i) aggregated A β are able to activate and become bound by C3b^{29–31}; (ii) in a transgenic mouse model, an increase in C3 expression coincides with a smaller degree of A β deposition and neuropathology^{32,33}; (iii) conversely, expression of a C3 convertase inhibitor such as the rodent complement receptor 1-related gene/protein γ (*Crry*) results in increased A β deposition and neurodegeneration³³; (iv) using human erythrocytes (the cell type that most abundantly expresses CR1), it has been observed that this cell type is able to sequester A β_{42} (refs. 31,32) and to favor its clearance via the C3b-mediated adherence to erythrocyte CR1 (ref. 32); (v) finally, this mechanism may be deficient in individuals with Alzheimer's disease³¹. Altogether, these data support a protective role for CR1 via the generation and binding of C3b, which may contribute to A β clearance³³.

We calculated the attributable fractions of risk to be 25.5% for *APOE*, 8.9% for *CLU* and 3.8% for *CR1*. As these calculations are based on the combined stage 1 and 2 samples, the estimates are biased upward. Nevertheless, if the estimate that 60–80% of the Alzheimer's disease risk is due to genetic factors⁵ is correct, additional genetic susceptibility loci remain to be identified, which is also true for many other diseases in which loci have been successfully mapped by GWA^{34,35}. In a recent GWAS, the *PCDH11X* gene was found to be a genetic determinant of Alzheimer's disease¹². We examined the effect of 17 SNPs in the *PCDH11X* locus in the stage 1 samples. Although none reached our criteria for suggestive level of significance, we did observe $0.01 < P < 0.05$ for several SNPs (**Supplementary Table 6**). Thus, this locus may also contribute to the Alzheimer's disease risk in our dataset, albeit not as significantly as the other loci that we have identified.

only includes *CR1*. Several observations suggest that pathways involving C3b and CR1 are involved in the Alzheimer's disease process, particularly in A β clearance. Briefly, complement activation leads to the formation of C3-cleaving enzymes, known as C3 convertases, on the surface of the pathogen or protein undergoing complement attack. Cleavage of C3 results in the formation of C3a and C3b fragments. Whereas C3a is involved in the chemotaxis of phagocytes, C3b binds covalently to acceptor molecules and can mediate phagocytosis through CR1^{27,28}.

Table 3 Association of SNPs at the *CR1* locus with Alzheimer's disease in the Stage 1 and Stage 2 samples

	rs6656401							
	<i>N</i>		<i>MAF</i>		<i>HWE</i>	Association test		<i>P</i> value
	Cases	Controls	Cases	Controls		OR (95% CI)		
Stage 1	2,025	5,324	0.22	0.18	9.9×10^{-1}	1.27 (1.16–1.39)	1.8×10^{-7}	
Stage 2	3,880	3,198	0.20	0.18	5.3×10^{-2}	1.16 (1.06–1.27)	8.2×10^{-4}	
Belgium	1,066	500	0.22	0.18	2.2×10^{-1}	1.24 (0.99–1.24)	5.6×10^{-2}	
Finland	608	654	0.21	0.17	2.9×10^{-1}	1.38 (1.12–1.70)	2.5×10^{-3}	
Italy	1,472	1,243	0.20	0.20	2.2×10^{-1}	1.03 (0.88–1.17)	8.4×10^{-1}	
Spain	734	801	0.20	0.16	4.2×10^{-2}	1.23 (1.02–1.47)	2.6×10^{-2}	
Stage 1 + 2	5,905	8,526	0.21	0.19	1.1×10^{-1}	1.21 (1.14–1.29)	3.5×10^{-9}	
APOE $\epsilon 4$ carriers	2,497	1,632	0.22	0.18	7.5×10^{-1}	1.38 (1.22–1.55)	1.8×10^{-7}	
APOE $\epsilon 4$ non-carriers	2,761	6,780	0.21	0.19	4.3×10^{-2}	1.13 (1.04–1.23)	3.7×10^{-3}	
rs3818361								
Stage 1	2,018	5,324	0.22	0.18	8.5×10^{-1}	1.28 (1.17–1.40)	8.5×10^{-8}	
Stage 2	3,717	3,094	0.22	0.20	1.1×10^{-1}	1.11 (1.02–1.22)	1.6×10^{-2}	
Belgium	972	436	0.24	0.24	9.9×10^{-1}	1.05 (0.84–1.48)	6.8×10^{-1}	
Finland	590	634	0.22	0.19	1.5×10^{-1}	1.26 (1.03–1.57)	2.6×10^{-2}	
Italy	1,423	1,232	0.20	0.20	9.2×10^{-1}	1.03 (0.89–1.18)	7.2×10^{-1}	
Spain	732	792	0.21	0.18	1.0×10^{-2}	1.21 (1.01–1.44)	3.8×10^{-2}	
Stage 1 + 2	5,735	8,418	0.22	0.19	2.2×10^{-1}	1.19 (1.11–1.26)	8.9×10^{-8}	
APOE $\epsilon 4$ carriers	3,032	1,696	0.22	0.18	8.9×10^{-1}	1.34 (1.20–1.50)	2.9×10^{-7}	
APOE $\epsilon 4$ non-carriers	2,661	6,707	0.21	0.19	1.8×10^{-1}	1.12 (1.03–1.22)	6.4×10^{-3}	

P values and ORs with their associated 95% CIs have been calculated under an additive model using logistic regression models adjusted for age, gender and centers when necessary. *MAF*, minor allele frequency; *HWE*, *P* value for the test of Hardy-Weinberg equilibrium in controls.

Table 4 Association results for haplotypes at the CR1 locus

Haplotypes	Stage 1				Stage 2				Stage 1 + 2			
	Cases	Controls	OR (95% CI)	P value	Cases	Controls	OR (95% CI)	P value	Cases	Controls	OR (95% CI)	P value
GG	0.772	0.813	Ref.	–	0.778	0.786	Ref.		0.776	0.803	Ref.	
<u>GA</u>	0.011	0.009	1.25 (0.87–1.80)	2.2×10^{-1}	0.016	0.017	1.01 (0.77–1.33)	6.4×10^{-1}	0.014	0.012	1.13 (0.89–1.43)	3.9×10^{-1}
<u>AA</u>	0.207	0.170	1.28 (1.17–1.40)	1.4×10^{-7}	0.198	0.182	1.17 (1.07–1.27)	6.8×10^{-4}	0.202	0.175	1.22 (1.15–1.30)	3.1×10^{-10}

The results have been calculated using the THESIAS program with adjustment for age, gender and center (see Online Methods). The *P* values for the global association were 7.5×10^{-7} , 3.0×10^{-3} and 1.5×10^{-10} , respectively, for the data from stage 1, stage 2 and stage 1 and 2 combined. The markers as ordered from left to right (5' to 3') are rs6656401 and rs3818361. Minor alleles are underlined.

In an independent study published in this issue of *Nature Genetics*³⁶, Williams and colleagues report an independent GWAS of late-onset Alzheimer's disease and also report association with markers at *CLU*. In comparing these two studies, the Belgian samples used in the replication phase of each were found to overlap. To test for the independence of these results, we tested our association results with the entire Belgian collection removed from the replication phase. We find that the association to *CLU* remains significant, with OR = 0.89, 95% CI 0.83–0.97 and $P = 5.3 \times 10^{-3}$ for association at rs11136000 in the stage 2 samples when the Belgian collection is excluded. Although this is marginally less significant than the conservative *P* value that we applied at replication phase, the combined stage 1 and 2 samples gave OR = 0.86, 95% CI 0.81–0.91 and $P = 4.8 \times 10^{-8}$ with the Belgian collection excluded, meeting our criterion for genome-wide significance. Thus these two studies provide strong independent evidence of an association of Alzheimer's disease risk with *CLU* markers. In addition, evidence of association with the *CR1* markers at rs3818361 in the GWA data from Williams and her colleagues³⁶ exceeded our criteria for replication in their study ($P = 9.2 \times 10^{-6}$), providing additional confirmation of the relationship between *CR1* markers and late-onset Alzheimer's disease (OR = 1.17, 95% CI 1.09–1.25). We also examined *PILCAM* markers on chromosome 11 for which association has been identified and replicated in the second study. In our stage 1 data, we also detected evidence of association for *PILCAM* markers with *P* values between 10^{-2} and 10^{-3} (rs541458 OR = 0.88, 95% CI 0.81–0.96, $P = 2.8 \times 10^{-3}$ and results available online in GWAS results).

In summary, in addition to the previously known *APOE* locus, we have identified loci at *CLU* and *CR1* that are potentially associated with the risk of late-onset Alzheimer's disease. Biological evidence suggests that the genes at these loci, along with *APOE*, are involved in A β clearance. These data may indicate that whereas familial early-onset forms of Alzheimer's disease are mainly linked to genes implicated in A β overproduction, genetic variants at *APOE* and these newly defined loci may influence susceptibility to late-onset forms of the disease as a result of roles in A β clearance.

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.

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AUTHOR CONTRIBUTIONS

Project conception, design, management. J.-C.L., M.L., P.A. **Phenotype collection, data management.** France: D.C., B.T., L.L., C.B., F.Pasquier, N.F., P.B.-G., O.H., C.L., C.D., C.J., T.L., D.H., K.R., H.B., J.-F.D., C.T., A.A. Belgium: K.S., K.B., S.E., P.D.D., C.V.B. Finland: M.H., S.Helisalmi, H.S. Italy: E.P., P.Bosco, M.M., F.Panza, B.N., P.Bossù, P.P., G.A., D.S., D.G., F.L. Spain: O.C., M.J.B., I.M., A.F., M.M.d.P., V.A. **Genome-wide, validation genotyping.** S.Heath., D.Z., I.G. **Data analysis:** J.-C.L., S.Heath., G.E., M.L., P.A. **Writing group:** J.-C.L., S.Heath., M.L., P.A.

COMPETING INTERESTS STATEMENT

The authors declare competing financial interests: details accompany the full-text HTML version of the paper at <http://www.nature.com/naturegenetics/>.

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ONLINE METHODS

Sample populations. The case-control cohorts are described in the **Supplementary Note**.

Genotyping. DNA samples were transferred to the French Centre National de Génotypage for genotyping. First stage samples that passed DNA quality control were genotyped with Illumina Human610-Quad BeadChips. Genotype data were retained in the study for samples that had been successfully genotyped for >98% of the SNP markers. SNPs with call rates of <98%, with MAF <1% or showing departure from Hardy-Weinberg equilibrium in the control population ($P < 10^{-6}$) were excluded. We removed 134 Alzheimer's disease case and 980 control samples because the individuals who had provided them were found to be first- or second-degree relatives of other study participants or were assessed as being of non-European descent based on genetic analysis using methods described³⁷. This led us to retain 537,029 autosomal SNPs genotyped in 2,032 cases and 5,328 controls. Stage 2 genotyping was performed using Taqman (Applied Biosystems) or Sequenom assays. The primer and probe sequences for the genotyping assays are available upon request. To avoid any genotyping bias, cases and controls were randomly mixed when genotyping, and laboratory personnel were blinded to case or control status. The genotyping success rate was at least 95%, and no departure from Hardy-Weinberg equilibrium was observed for the markers included in the second stage.

Statistical analysis. We evaluated the case and control differences using logistic regression, which optionally incorporated principal components that were significantly associated with disease status to account for possible population stratification as described^{37,38}. We hypothesized that the relatively high genomic control parameter found in the absence of the principal-components adjustment was due to differences in the representation of various French regions in the case and control series. Therefore, we further explored the robustness of our conclusions by incorporating 6,734 anonymized samples from France and other European countries as additional controls (unpublished data and ref. 37). With the inclusion of the additional samples, the genomic control parameter was 1.04 without principle-components adjustment and 1.03 after the adjustment. Inclusion of the additional controls did not substantially modify the association statistics for markers in regions showing suggestive evidence of association ($P < 10^{-5}$) after correction for population structure in the primary analysis (**Supplementary Tables 1 and 2**). Markers in these regions were analyzed at stage 2. We included imputed markers in these regions and selected markers for genotyping at stage 2 if the marker showed at least moderate association with disease status ($P < 10^{-4}$) (either with or without correction for population structure), the marker was not in strong LD with another marker entering stage 2, and a genotyping assay for the marker could be successfully designed.

We did statistical analyses under an additive genetic model using logistic regression taking age, sex and disease status into account and using SAS software version 9.1 (SAS Institute). Population controls that were not genotyped specifically for this study were excluded, as were any samples with missing age or gender data. This gave a maximum of 2,025 Alzheimer's disease cases and 5,328 controls in stage 1 and 3,359 cases and 2,633 controls in stage 2. Information on age and gender of the cases and controls included in these analyses are shown in **Supplementary Table 3**. Homogeneity of the odds ratios in different collections was tested using the Breslow-Day test³⁹. We also used the Mantel-Haenszel method as implemented in Review Manager 5.0 to evaluate odds ratios across collections. Interactions between *CLU* or *CR1* SNPs and *APOE* $\epsilon 4$ polymorphism were tested in logistic regression models adjusted for age, gender and center where the samples were taken. The solid spine haplotype block definition in Haploview 4.0 was used to generate LD blocks of the genomic regions encompassing the *CLU* or *CR1* genes from imputed SNPs (MAF >5%)⁴⁰. Associations of the *CLU* and *CR1* haplotypes were estimated using either logistic regression or a proportional hazards models using THESIAS 3.0, which implements a maximum likelihood model and uses an s.e.m. algorithm⁴¹. The population attributable risk (PAR) fraction was estimated using the following formula: $PAR = F(OR - 1)/(F(OR - 1) + 1)$, where F is the frequency of the deleterious allele in the sample and OR is the odds ratio of Alzheimer's disease risk associated with the deleterious allele.

The criterion for genome-wide significance was defined in stage 1 with a conservative Bonferroni correction ($P < 9.3 \times 10^{-8}$; 0.05/537,029 SNPs tested). In stage 2, the level for significance was defined at $P < 4.5 \times 10^{-3}$ (0.05/11 SNPs tested).

URLs. Haploview: <http://www.broad.mit.edu/mpg/haploview/index.php>; Revman: <http://www.cc-ims.net/revman/>; THESIAS: <http://ecgene.net/genecanvas/uploads/THESIAS3.1/>; HapMap: <http://www.hapmap.org>; website for GWAS results: <http://www.cng.fr/alzheimer/>.

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