Alzheimer risk genes modulate the relationship between plasma apoE and cortical PiB binding

ABSTRACT

Objective: We investigated the association between apoE protein plasma levels and brain amyloidosis and the effect of the top 10 Alzheimer disease (AD) risk genes on this association.

Methods: Our dataset consisted of 18 AD, 52 mild cognitive impairment, and 3 cognitively normal Alzheimer’s Disease Neuroimaging Initiative 1 (ADNI1) participants with available \[^{11}C\]-Pittsburgh compound B (PiB) and peripheral blood protein data. We used cortical pattern matching to study associations between plasma apoE and cortical PiB binding and the effect of carrier status for the top 10 AD risk genes.

Results: Low plasma apoE was significantly associated with high PiB SUVR, except in the sensorimotor and entorhinal cortex. For BIN1 rs744373, the association was observed only in minor allele carriers. For CD2AP rs9349407 and CR1 rs3818361, the association was preserved only in minor allele noncarriers. We did not find evidence for modulation by CLU, PICALM, ABCA7, BIN1, and MS4A6A.

Conclusions: Our data show that BIN1 rs744373, CD2AP rs9349407, and CR1 rs3818361 genotypes modulate the association between apoE protein plasma levels and brain amyloidosis, implying a potential epigenetic/downstream interaction. Neuronal Genet 2015;1:e22; doi: 10.1212/NXG.0000000000000022

GLOSSARY

AD = Alzheimer disease; ADNI = Alzheimer’s Disease Neuroimaging Initiative; A\(\beta\) = \(\beta\)-amyloid; CDR = Clinical Dementia Rating; GWAS = genome-wide association studies; MCI = mild cognitive impairment; MMSE = Mini-Mental State Examination; MVb = multivesicular body; NIA = National Institute on Aging; PiB = Pittsburgh compound B; SNP = single nucleotide polymorphism; SUVR = standardized uptake value ratio.

The strongest genetic risk factor for late-onset Alzheimer disease (AD) is the apolipoprotein E gene (APOE), with the APOE e4 allele carrying the greatest risk via effects on \(\beta\)-amyloid (A\(\beta\)) metabolism. APOE codes for the apolipoprotein E protein (apoE), which plays an essential role in cholesterol metabolism, neuronal trafficking, synaptogenesis, and blood–brain barrier integrity. ApoE-containing lipoproteins bind A\(\beta\) and promote its clearance and degradation. The apoE4 isoform is preferentially degraded by astrocytes, leading to reduced overall brain apoE levels, reduction of A\(\beta\) clearance, and higher A\(\beta\) levels in the brain and plasma. Hence apoE protein plasma levels are significantly lower in patients with AD and in APOE e4 carriers.

Amyloid PET tracers provide reliable in vivo visualization of cortical fibrillar A\(\beta\) plaque deposition. The APOE e4 genotype has been shown to positively associate with brain amyloid levels.

*Co-first authors.

From the University of California Berkeley (A.L.), Berkeley; Oakland University William Beaumont School of Medicine (K.S.H.), Rochester, MI; Department of Neurology (K.S.H., N.G., A.E.B., E.T., G.C., L.G.A.), David Geffen School of Medicine at UCLA, Los Angeles, CA; Drexel University College of Medicine (L.M.R.), Philadelphia, PA; Northwestern University Feinberg School of Medicine (J.E.), Chicago, IL; Veterans Affairs Greater Los Angeles Healthcare System (E.T., G.C.), Los Angeles, CA; School of Nursing (K.G.), UCLA, Los Angeles, CA; Department of Radiology and Imaging Sciences, Center for Neuroimaging (A.J.S., L.G.A.), Department of Neurology (L.G.A.), and Department of Medical and Molecular Genetics (L.G.A.), School of Medicine, Indiana University, Indianapolis; Department of Pathology and Laboratory Medicine (L.M.S., J.Q.T.), University of Pennsylvania School of Medicine, Philadelphia; Department of Public Health and Neuroscience (W.J.J.), UC Berkeley, CA; and Department of Veterans’ Affairs Medical Center (M.W.W.), San Francisco, CA.

Coinvestigators are listed at Neurology.org/ng.

Funding information and disclosures are provided at the end of the article. Go to Neurology.org/ng for full disclosure forms. The Article Processing Charge was paid by the authors.

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivatives License 4.0 (CC BY-NC-ND), which permits downloading and sharing the work provided it is properly cited. The work cannot be changed in any way or used commercially.
APOE e4 carriers show significantly higher prevalence of Pittsburgh compound B (PiB) uptake than noncarriers across all disease stages.\textsuperscript{8,9}

ApoE protein plasma levels were reported to be lower in research subjects with high PiB–PET binding in one study\textsuperscript{7} but not in another.\textsuperscript{10} Such discrepant results are reported to reconcile and raise the question whether other genetic and perhaps even environmental factors influence this association.

Genome-wide association studies (GWAS) have identified novel AD risk variants. Of those, PICALM rs3851179, BIN1 rs7561528, and CR1 rs1408077 have been associated with cortical and hippocampal atrophy,\textsuperscript{11} and ABCA7 rs3764650,\textsuperscript{12} ABCA7 rs3752246,\textsuperscript{13} BIN1 rs744373,\textsuperscript{14} CR1 rs6701713,\textsuperscript{11} CR1 rs3818361,\textsuperscript{10} CR1 rs6656401, and CLU rs3818361\textsuperscript{14} have been associated with PET amyloid deposition. An association between postmortem amyloid burden and CD2AP rs9349407 was reported by one group,\textsuperscript{12} but no significant relationship with PiB binding in nondemented elderly was observed by others.\textsuperscript{13}

The exact mechanism through which these genes exert an association with amyloid burden is not clear. In this study, we assess the associations between the plasma levels of the APOE gene product—the apoE protein—and cortical PiB binding, and examine whether the presence of other AD risk variants modulates this association.

**METHODS**

**Standard protocol approvals, registrations, and patient consents.** Data used in the preparation of this article were obtained from the Alzheimer’s Disease Neuroimaging Initiative (ADNI) database (http://adni.loni.usc.edu). ADNI was launched in 2003 by the National Institute on Aging (NIA), the National Institute of Neurological and Communicative Disorders and Stroke and the AD and Related Disorders Association (NINCDS-ADRDA) criteria. Participants with AD were required to have Mini-Mental State Examination (MMSE) scores between 20 and 26 and a Clinical Dementia Rating (CDR) score of 0.5 to 1 at baseline. Qualifying MCI participants had memory complaints but no significant functional impairment, scored between 24 and 30 on the MMSE, had a global CDR score of 0.5, a CDR memory score of 0.5 or greater, and objective memory impairment on Wechsler Memory Scale—Logical Memory II test. Healthy control participants had MMSE scores between 24 and 30, a global CDR score of 0, and did not meet criteria for MCI and AD. Participants were excluded if they refused or were unable to undergo MRI; had other neurologic disorders, active depression, or a history of psychiatric diagnosis, alcohol, or substance dependence within the past 2 years; had less than 6 years of education; or were not fluent in English or Spanish. The full list of inclusion/exclusion criteria may be accessed on pages 23–29 of the online ADNI protocol (see http://www.adni-info.org/Scientists/ADNIScientists/Home.aspx). Written informed consent was obtained from all participants. The subset of 18 AD, 52 MCI, and 3 cognitively normal ADNI participants who received [\textsuperscript{11}C]-PiB have been included in this study.

**PiB analyses.** We downloaded all available PiB ADNI1 scans from the ADNI Web site (http://adni.loni.usc.edu) in October 2008. A detailed description of PiB-PET acquisition may be found at www.adni-info.org. Briefly, ADNI PiB images were collected at 12 ADNI sites. [\textsuperscript{11}C]-PiB with minimum 90% radiochemical purity and minimum specific activity of 300 Ci/ mmol was synthesized. Participants were injected with 15 ± 1.5 mCi PiB. Dynamic acquisition frames were obtained on a PET scanner 50–70 minutes after injection. A PiB standardized uptake value ratio (SUVR) image was obtained by averaging the individual 50–70 minutes after injection frames.

PiB scans were normalized to the mean PiB retention value of the cerebellar cortex (SUVR\textsubscript{C3,C6}). After normalization, the PiB SUVR\textsubscript{C3,C6} data were spatially coregistered to the participants’ baseline MRI scan using the Minitracc algorithm and 9-parameter (9P) transformation (3 translations, 3 rotations, 3 scales). The PiB data were smoothed with a 15-mm kernel and convolved onto the 3D hemispheric models derived with the cortical pattern matching technique as previously described.\textsuperscript{16,17}

**Genetic analyses.** Genotypic data were extracted from the publicly accessible GWAS data from the ADNI Web site (http://adni.loni.usc.edu). The detailed GWAS genotyping protocol has been previously described.\textsuperscript{18} Briefly, single nucleotide polymorphism (SNP) genotyping was completed on all ADNI participants for more than 620,000 target SNPs using a total of 7 ml of blood from all participants. Genomic DNA was extracted using the QiAamp DNA Blood Maxi Kit (Qiagen, Inc, Valencia, CA) and analyzed using Human 610-Quad BeadChip (Illumina, Inc, San Diego, CA) according to the manufacturer’s protocols (Infinium HD Assay; Super Protocol Guide; rev. A, May 2008).

APOE genotyping was carried out by PCR amplification, HflI restriction enzyme digestion, and subsequent standard gel resolution and visualization processes.\textsuperscript{19,20} We classified participants as carriers and noncarriers for the following AD risk SNPs: BINI rs744373 and rs7561528; CLU rs11136000, rs2279590, and rs6313888; PICALM rs3851179 and rs541458; CR1 rs6818361; ABCA7 rs3764650; MS4A6A rs610932; CD33 rs3826656 and rs3865444; MS4A4E rs670139; CD2AP rs9349407 and rs3865444. The following SNPs had balanced carrier/noncarrier status and were further analyzed:
Plasma biomarkers. Plasma collection was conducted according to ADNI standard operating protocols. Briefly, morning fasting plasma samples were obtained at each visit. The plasma samples were then sent to Rules-Based Medicine (RBM, Austin, TX) for measurement of 190 protein analytes with an immunoassay panel developed on the Luminex xMAP (Austin) platform. Each plate was run with 3 levels of quality control, and each analyte had a validation report; samples from the total ADNI plasma cohort were run on 15 plates. ApoE protein plasma levels were freely downloaded from http://adni.loni.usc.edu.

RESULTS Demographic comparisons. All demographic variables and comparisons are shown in table 1.

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Demographic statistics of participant pool, indicated by the pooled sample and genotype</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age, y</strong></td>
<td><strong>Sex</strong></td>
</tr>
<tr>
<td><strong>Pooled sample</strong></td>
<td>75.3 ± 8</td>
</tr>
<tr>
<td><strong>BNI rs744373 carriers</strong></td>
<td>74.7 ± 7</td>
</tr>
<tr>
<td><strong>BNI rs744373 noncarriers</strong></td>
<td>76.2 ± 9</td>
</tr>
<tr>
<td><strong>x² (p value)</strong></td>
<td>0.479</td>
</tr>
<tr>
<td><strong>CD2AP rs9349407 carriers</strong></td>
<td>75.7 ± 8</td>
</tr>
<tr>
<td><strong>CD2AP rs9349407 noncarriers</strong></td>
<td>74.9 ± 8</td>
</tr>
<tr>
<td><strong>x² (p value)</strong></td>
<td>0.478</td>
</tr>
<tr>
<td><strong>CD33 rs3826656 carriers</strong></td>
<td>75.5 ± 7</td>
</tr>
<tr>
<td><strong>CD33 rs3826656 noncarriers</strong></td>
<td>75.2 ± 9</td>
</tr>
<tr>
<td><strong>x² (p value)</strong></td>
<td>0.412</td>
</tr>
<tr>
<td><strong>CLU rs2279590 carriers</strong></td>
<td>26.1 ± 3</td>
</tr>
<tr>
<td><strong>CLU rs2279590 noncarriers</strong></td>
<td>76.0 ± 7</td>
</tr>
<tr>
<td><strong>x² (p value)</strong></td>
<td>0.484</td>
</tr>
<tr>
<td><strong>CLU rs9331888 carriers</strong></td>
<td>77.2 ± 8</td>
</tr>
<tr>
<td><strong>CLU rs9331888 noncarriers</strong></td>
<td>72.4 ± 7</td>
</tr>
<tr>
<td><strong>x² (p value)</strong></td>
<td>0.412</td>
</tr>
<tr>
<td><strong>CR1 rs3818361 carriers</strong></td>
<td>76.4 ± 8</td>
</tr>
<tr>
<td><strong>CR1 rs3818361 noncarriers</strong></td>
<td>74.6 ± 8</td>
</tr>
<tr>
<td><strong>x² (p value)</strong></td>
<td>0.48</td>
</tr>
</tbody>
</table>

Abbreviations: AD = Alzheimer disease; HC = healthy control; MCI = mild cognitive impairment; MMSE = Mini-Mental State Examination. Age, sex, and MMSE displayed in mean ± SD.

*A Two-tailed x² test p value with significant differences. Only genes that had balanced distribution between carriers and noncarriers were considered.*
We found no significant differences in age, sex distribution, or level of education between carriers and non-carriers for each gene. There were significantly more MCI than AD and healthy control subjects among CD2AP noncarriers ($p = 0.03$). BIN1 and CD2AP carriers had lower MMSE scores than noncarriers (BIN1 mean MMSE 25.7 vs 26.5, $p = 0.005$; CD2AP mean MMSE 25.3 vs 27.0 $p = 0.047$). Of note, there
were no significant differences in APOE e4 status between carriers and noncarriers for any risk gene.

None of the AD risk genotypes showed significant association with PiB SUVR in the pooled sample. Plasma apoE levels showed an association with PiB SUVR throughout the brain with the exception of the sensorimotor and entorhinal cortex (figure 1) across the pooled sample (left \( p_{corr} = 0.004 \), right \( p_{corr} = 0.008 \)).

Our analyses showed no evidence of modulation of the apoE protein–PiB association by CLU rs9331888 and CLU rs2279590. Plasma apoE levels showed an association with PiB SUVR in CDA2AP rs9349407 and CR1 rs3818361 noncarriers \( (\text{table 2 and figure e-1}) \).

Plasma apoE levels showed an association with PiB SUVR throughout the brain. Yet, when stratified by AD risk genotypes, plasma apoE showed a significant association with PiB SUVR only in noncarriers of the minor allele of CDA2AP rs9349407 and CR1 rs3818361 and only in carriers of the BIN1 rs744373 minor allele. These findings imply a downstream interaction between these genes and the apoE disease-associated pathways and suggest a direct or modulatory role on Aβ accumulation and clearance. Brain and plasma apoE levels are believed to be lower than apoE3 levels because of a difference in their fate after they are endocytosed and routed to multivesicular bodies that function as sorting endosomes. ApoE3 is packaged into recycling endosomes and retroendocytosed back to the surface and secreted, whereas apoE4 is typically trafficked on through to the lysosome and degraded. Plasma apoE levels reflect this difference in processing between apoE3 and apoE4.

Bridging integrator 1, or BIN1, encodes a nucleocytoplasmic adaptor protein abundantly expressed in the CNS that is involved in synaptic vesicle endocytosis, immune response, calcium homeostasis, and apoptosis. The protein encoded by BIN1 has been shown to link the microtubule cytoskeleton and cellular membrane and has been implicated in amyloid precursor protein turnover and hence Aβ production through clathrin-mediated amyloid precursor protein endocytosis. BIN1 rs744373 minor allele carriers have been reported to have an increased risk of AD. Increased BIN1 rs744373 expression is associated with later disease onset and shorter disease duration. Postmortem, no association with amyloid plaque burden

<table>
<thead>
<tr>
<th>Table 2</th>
<th>p values of the association between plasma apoE levels and cortical PiB binding stratified by BIN1, CDA2AP, and CR1 carrier status</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Covariates</strong></td>
<td><strong>Age/sex</strong></td>
</tr>
<tr>
<td></td>
<td>L hemisphere</td>
</tr>
<tr>
<td><strong>BIN1</strong></td>
<td></td>
</tr>
<tr>
<td>Carriers</td>
<td>0.006*</td>
</tr>
<tr>
<td>Noncarriers</td>
<td>NS</td>
</tr>
<tr>
<td><strong>CD2AP</strong></td>
<td></td>
</tr>
<tr>
<td>Carriers</td>
<td>NS</td>
</tr>
<tr>
<td>Noncarriers</td>
<td>0.003*</td>
</tr>
<tr>
<td><strong>CR1</strong></td>
<td></td>
</tr>
<tr>
<td>Carriers</td>
<td>NS</td>
</tr>
<tr>
<td>Noncarriers</td>
<td>0.008*</td>
</tr>
</tbody>
</table>

Abbreviations: NS = not significant; PiB = Pittsburgh compound B.

*Significant.
has been observed for BIN1 rs7561528,12 and BIN1 rs744373,24 consistent with our results in the pooled sample. An association of BIN1 rs744373 minor allele with in vivo amyloid deposition measured with 11C-florbetapir has been reported,14 but this effect disappeared in the presence of the protective PICALM minor allele at rs3851179. BIN1 rs744373 also failed to show an association with CSF Aβ levels.28

In our study, the presence of the BIN1 rs744373 risk variant was associated with preserved and potentially enhanced association between apoE protein plasma levels and PiB binding, lending evidence for a potential downstream functional interaction between the BIN1 and APOE genes. BIN1 rs744373 increases mRNA expression and protein expression.29 As mentioned above, apoE levels are strongly related to the classic clathrin-dependent receptor-mediated endocytosis with impaired recycling of apoE4 isoforms.22 BIN1 is an integral part of the sorting nexin-4/BAR complexes. These complexes have been shown to regulate the Rab5/7 switch that influences the transition from early to late endosomes and controls traffic to the lysosomes. They also seem to regulate Rab4/Rab11 transition to recycling endosomes.30 Aβ and apoE are known for trafficking through Rab5/Rab7 to lysosomes and to Rab11 recycling endosomes for degradation. This occurs in an apoE isoform–dependent manner in which apoE3 promotes greater Aβ and apoE trafficking to the lysosome for clearance.31 Therefore, BIN1 is ideally positioned to regulate both Aβ and apoE influx into lysosomes and trafficking toward degradation or recycling through retroendocytosis. Thus higher BIN1 expression can be hypothesized to increase apoE recycling and elevate protein levels notably, and this would be linked to faster amyloid clearance and less deposition associated with higher apoE levels.

CD2AP, the CD2-associated protein gene, codes for a scaffolding protein that plays a role in the formation oftight junctions, endocytosis, cellular waste management, and immune response.24 CD2AP has an established role in endocytosis, vesicle trafficking within the cell, and formation of the cytoskeleton.32 CD2AP is involved in endosome trafficking,33 binds directly to Rab4, and regulates recycling endosomes,34 which play an essential role in the retroendocytosis of apoE that controls apoE levels.22 Consistent with this, CD2AP-knockout mice have deficits in multivesicular body (MVB) formation and endosomal–lysosomal trafficking.29 Intraneuronal Aβ accumulation occurs in MVBs, thereby impairing their function.35 This supports a possible interaction between CD2AP and Aβ metabolism. CD2AP rs9349407 has been associated with postmortem plaque burden and AD status.36,37 However, studies have failed to find a direct association between this SNP and PiB binding in ADNI.13 Our results further elaborate on this complex relationship.

We found that the association between apoE protein plasma levels and PiB binding is no longer present in minor allele carriers, indicating that CD2AP influences amyloid pathology. Since CD2AP rs9349407 trends toward reducing CD2AP expression in human brain,29 one might hypothesize that decreased CD2AP (or altered interaction with binding partners) could reduce apoE’s isoform-dependent recycling and endosomal–lysosomal traffic that regulates both apoE levels and Aβ clearance in lysosomes.31 The latter has been implicated as a cause for both low plasma apoE4 and defects in Aβ clearance.31 Furthermore, endocytosed apoE isoforms differentially recycle and degrade in lysosomes.22 They also regulate endosomal–lysosomal Aβ clearance.31 Since CD2AP rs9349407 trends toward reducing CD2AP expression in human brain,29 one might hypothesize that decreased CD2AP (or altered interaction with binding partners) could reduce apoE’s isoform-dependent recycling and endosomal–lysosomal traffic that regulates both apoE levels and Aβ clearance in lysosomes.31 Thus, lowering CD2AP could diminish the correlations between PiB and plasma apoE.

The CRI gene encodes complement component 3b/4b receptor 1—a membrane glycoprotein found on erythrocytes, leukocytes, glomerular podocytes, and splenic follicular dendritic cells. This protein mediates cell binding and removal of complement-activated immune complexes.24 CRI serves to mediate clearance of immune complexes and phagocytosis by neutrophils and monocytes48 and plays a role in antigen presentation to B lymphocytes.39 Increased CRI expression has been associated with AD diagnosis and worse cognition.24 The presence of the CRI rs3818361 risk variant is associated with lower PiB binding in 2 separate cognitively normal cohorts including an ADNI1 subsample10 but not in the full ADNI1 sample.14 At the same time, 2 other SNPs, CRI rs6656401 and rs6701713, showed an association with increased amyloid plaque burden postmortem.12,40 In this study we observed that CRI rs3818361 minor allele carrier status leads to a loss of the association between apoE protein plasma levels and PiB binding, which disappeared in the presence of APOE e4 as covariate. An epistatic interaction between the APOE genotype and CRI rs3818361 has been previously reported.41,42

Several strengths and limitations of this study should be acknowledged. ADNI is the premier longitudinal biomarker study in AD. ADNI uses unified subject assessment, MRI, PiB-PET, CSF, and peripheral blood collection protocols and meticulous data quality control across all study sites. One of the major limitations of our study is its small sample size limited by PiB availability in ADNI1. Significant disproportion between carriers and noncarriers for the remaining ADNI risk SNPs led to the exclusion of these variants,
allowing us to test our hypothesis for only 6 SNPs across 5 AD risk genes. Furthermore, other unknown genetic interactions may also affect the plasma apoE–cortical PiB associations. Last but not least, ADNI uses rigorous exclusion criteria typical of clinical trials and the study population is not representative of the general population, which may negatively affect the generalizability of our results.

The ADNI PET core chose cerebellar gray matter as the reference region for intensity normalization of ADNI1 PiB data. This is one of the commonly used approaches. Other regions that have been used in the amyloid PET literature include whole cerebellum, periventricular white matter, whole brainstem, and composite regions derived from more than one of these measures. Recently, a group of PET experts compared the performance of 4 of these normalization regions: whole cerebellum, cerebellar gray matter, pons, and whole cerebellum plus brainstem. The authors concluded that normalization to the pons performed worse than normalization strategies including the cerebellum. The 2 regions that performed best (i.e., had lowest SUVR variance) were whole cerebellum and whole cerebellum plus brainstem.15 Our study uses the cerebellar gray matter, which performed better than pons but resulted in more noise (higher SUVR variance) in the normalized data. Yet despite this potentially suboptimal signal-to-noise ratio, we were able to find significant associations.

Overall, our findings imply an interaction between several AD risk gene minor alleles and APOE genotype–driven brain amyloidosis. In the absence of a direct association with brain amyloidosis, we found that several AD risk genes nonetheless exert a modulatory effect on one of the most fundamental disease-associated pathophysiological events—APOE ε4-facilitated brain amyloid deposition. These findings lend the basis for further exploration of the exact AD-related pathophysiologic mechanisms of these genes and their products, which might ultimately lead to new therapeutic strategies.

AUTHOR CONTRIBUTIONS
Andreas Lazaris is a co-first author of the manuscript, responsible for study design, data processing, statistical analyses, and drafting of the manuscript. Kristy S. Hwang is a co-first author of the manuscript, assisting with study design, data processing, and the completion of integral statistical analyses. Ms. Hwang also authored several segments of the manuscript and took part in all revisions. Naira Goukasian completed some of the final analyses of our data and took part in revising of the manuscript. Leslie M. Ramirez completed some of the initial analyses of our data and took part in revising of the manuscript. Jennifer Eastman completed some of the initial analyses of our data and took part in revising of the manuscript. Anna E. Blanken completed some of the final analyses of our data and took part in revising of the manuscript. Edmond Teng took part in some analysis and interpretations of the data and participated in revising of the manuscript. Karen Gylys provided critical insights for interpretation of our results and participated in revising of the manuscript. Greg Cole provided critical insights for interpretation of our results and participated in revising of the manuscript. Andrew Saykin contributed to the overall ADNI study design, data collection, and genetic analyses. He also provided critical insights for interpretation of our results and participated in revising of the manuscript. Leslie Shaw contributed to the overall ADNI study design, data collection, and proteomic analyses. He also provided critical insights for interpretation of our results and participated in revising of the manuscript. Michael W. Weiner is the principal investigator of ADNI. He has contributed to the overall ADNI study design, data collection, and data analyses. He provided critical insights for interpretation of our results and participated in revising of the manuscript. Liana G. Apostolova is the senior author of this manuscript and is responsible for the study concept and design. She provided major oversight over all analyses, interpretation of results, and participated in writing of the manuscript.
board for Acta Neuropathologica Communications; and has received research support from NIH R01 AG027465-01A2, P50 AG16570, UCLA Alzheimer’s Disease Research Center, and the Lincy Foundation. Greg Cole holds patents for an FDDNP PET probe for protein aggregates, Curcumin formulation for enhanced bioavailability with lipidation and antioxidant stabilization, and Medical Food for brain health, provides unpaid advice to Neurovision Imaging, which is using his and coholders’ patent pending curcumin formulation for retinal imaging; has received research support from NIH NCCIH AT006816, Greater LA VA, and GREECC; receives royalties for the following: FDDNP UCLA patent royalties, Curcumin formulation/UCLAVA patent royalties, and Medical Food—UCLAVA patent royalties; and was an expert witness for legal proceedings regarding WhiteWave (makers of Horizon Milk). Andrew Saykin serves as ADNI Genetics Core Leader; receives research support from Siemens Medical Solutions and Welch Allyn, and from the following Alzheimer’s disease related NIH grants: R01 AG19771, R01 CA101318, R01 LM011360, U01 AG032984, R2C AG036535, and P30 AG101833; has served as a consultant to Siemens Healthcate, Eli Lilly, and Aldrey BioTek; has served on the advisory board for Siemens Healthcare and Eli Lilly; has received honoraria from Siemens Healthcare; and has served on the editorial board of Brain Imaging and Behavior (a Springer Journal). Leslie Shaw has been a consultant to Innotogenics and collaborates on quality assessment activities as part of the Alzheimer’s Disease Neuroimaging Initiative; has been a consultant for Janssen and Novartis as a member of advisory board and has received speaker fees and travel expenses from Eli Lilly and Company; has served on the editorial board for Therapeutic Drug Monitoring; and has received research support support from Eli Lilly, NIH, and the Alzheimer’s Disease Neuroimaging Initiative. John Q. Trojanowski has served on the editorial board of Alzheimer’s & Dementia; may accrue revenue on patents submitted by the University of Pennsylvania wherein he is inventor including: modified avidin-biotin technique; method of stabilizing microtubules to treat Alzheimer’s disease; method of detecting abnormally phosphorylated tau; method of screening for Alzheimer’s disease or disease associated with the accumulation of paired helical filaments; compositions and methods for producing and using homogeneous neuronal cell transplants; rat comprising straight filaments in its brain; compositions and methods for producing and using homogeneous neuronal cell transplants to treat neurodegenerative disorders and brain and spinal cord injuries; diagnostic methods for Alzheimer’s disease by detection of multiple MRNAs; and methods and compositions for determining lipid peroxidation levels in oxidant stress syndromes and diseases; compositions and methods for producing and using homogeneous neuronal cell transplants; method of identifying, diagnosing, and treating alpha-synuclein positive neurodegenerative disorders; mutation-specific functional impairments in distinct tau isoforms of hereditary frontotemporal dementia and parkinsonism linked to chromosome 17: genotyp predicts phenotype; microtubule stabilizing therapies for neurodegenerative disorders; and treatment of Alzheimer’s and related diseases with an antibody; is coinventor on patents submitted the University of Pennsylvania wherein he is inventor that has generated income he has received from the sale of Avid to Eli Lilly including: amyloid plaque aggregation inhibitors and diagnostic imaging agents; and has received research support from the Marian S. Ware Alzheimer Program and Benaroya. William J. Jagust has served as a consultant to Banner Alzheimer Institute, Genentech, Inc, Synarc, Janssen Alzheimer Immunotherapy, F. Hoffman LA Roche, Novartis, and Siemens; has served on the scientific advisory boards of Genentech, Inc and Novartis; has served on the editorial boards of Frontiers in Human Neuroscience, Annals of Neurology, Brain Imaging and Behavior, Alzheimer’s Disease and Associated Disorders, and Neuroimage: Clinical; and has received research support from Avid Radiopharmaceuticals F-AV-45-A14, NIH grants AG034570, AG025303, AG044292, AG012435, AG021028, AG031563, AG019724, AG030084, AG032006, and AG024904, and the Tau Consortium (Rainwater Foundation). Michael W. Weiner has served on the Scientific Advisory Boards for Pfizer, BOLT International, Neurotrophe Bioscience, Alzheon, University of Pennsylvania’s Neurosciences of Behavior Initiative, National Brain Research Center (NBRC), India, LEARN Program at University of North Carolina, Dolby Family Ventures, LP, ADNL, and Eli Lilly; has provided consulting to Synarc, Pfizer, Janssen, KJL Associates, Easton Associates, Harvard University, University of California, Los Angeles (UCLA), Alzheimer’s Drug Discovery Foundation (ADDF), Neutrotrope Bioscience, Avid Radiopharmaceuticals, ClearView Healthcare Partners, Perceptive Informatics, Smartfish AS, Decision Resources, Inc, Atari, Merck, Defined Health, Howard University, Bogen Idea, BioClinica, and Genentech; has received travel funding from Pfizer, Paul Sabatier University, MGI Group France, Travel Dreams, Inc, Neuroscience School of Advanced Studies (NSAS), Danone Trading, BV, CTAD Adv Cones, Kennes, Int, ADRC, UCLA, UCSD, ADCS, Sanofi-Aventis Groupe, University Center Hospital, Toulouse, Atari, AC Immune, Eli Lilly, New York Academy of Sciences (NYAS), National Brain Research Center, Northwestern University, Fidelity Biosciences Research Initiative, University of Pennsylvania, The Alzheimer’s Association, Merck, ADPD, Alzheimer’s Drug Discovery Foundation (ADDF), Tokyo University, Kyoto University, Cornell-Weill University, Rockefeller University, Memorial Sloan-Kettering Cancer Center, Biogen, and India for Johns Hopkins Medicine; has received honoraria from Pfizer, Tohoku University, Consortium for Multiple Sclerosis Centers (CMSC), and Danone Trading, BV; has received research support from research support from Merck and Avid; has been an employee of the University of California, San Francisco and the San Francisco VA Medical Center; has received research support from Merck, Avid, Lilly, Alzheimer’s Diseases Discovery Foundation (ADDF), the Veterans Administration (VA), Department of Defense, from the following grants: 2U2UAG024904, W81XWH-13-1-0259, W81XWH-12-2-0012, R01 AG10897, P41 EB015904, P01 AG19724, R01 AG032306, R01A GO3879, ADNI 2-12-23036, 20110506, R01 MH098062-01, and from NIH/NIA/National Institute of Mental Health, DOD, Alzheimer’s Association, Alzheimer’s Drug Discovery Foundation, Merck, Avid, and the Veterans Administration (VA). Luna G. Apostolova has served as a consultant to Lilly and GE Healthcare; has received speaker honoraria from Eli Lilly; has served on the editorial boards of Alzheimer and Dementia: Diagnosis, Assessment and Disease Monitoring; has served on the speakers bureaus of Eli Lilly and GE Healthcare; and has received research support from GE Healthcare and NIA. Go to Neurology.org/ng for full disclosure forms.

Received June 21, 2015. Accepted in final form August 13, 2015.

REFERENCES


Alzheimer risk genes modulate the relationship between plasma apoE and cortical PiB binding
Andreas Lazaris, Kristy S. Hwang, Naira Goukasian, et al.

*Neurol Genet* 2015:1;
DOI 10.1212/NXG.0000000000000022

This information is current as of October 15, 2015