

Genetic risk for Alzheimer disease predicts hippocampal volume through the human lifespan

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Abstract

Objective

To test the hypothesis that genetic risk for Alzheimer disease (AD) may represent a stable influence on the brain from early in life, rather than being primarily age dependent, we investigated in a lifespan sample of 1,181 persons with a total of 2,690 brain scans, whether higher polygenic risk score (PGS) for AD and presence of *APOE* ϵ 4 was associated with lower hippocampal volumes to begin with, as an offset effect, or possibly faster decline in older age.

Methods

Using general additive mixed models, we assessed the relations of PGS for AD, including variants in *APOE* with hippocampal volume and its change in a cognitively healthy longitudinal lifespan sample (age range: 4–95 years, mean visit age 39.7 years, SD 26.9 years), followed for up to 11 years.

Results

AD-PGS and *APOE* ϵ 4 in isolation showed a significant negative effect on hippocampal volume. The effect of a 1 sample SD increase in AD-PGS on hippocampal volume was estimated to -36.4 mm^3 (confidence interval [CI]: $-71.8, -1.04$) and the effect of carrying ϵ 4 allele(s) -107.0 mm^3 (CI: $-182.0, -31.5$). Offset effects of AD-PGS and *APOE* ϵ 4 were present in hippocampal development, and interactions between age and genetic risk on volume change were not consistently observed.

Conclusions

Endophenotypic manifestation of polygenic risk for AD may be seen across the lifespan in cognitively healthy persons, not being confined to clinical populations or older age. This emphasizes that a broader population and age range may be relevant targets for attempts to prevent AD.

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Glossary

AD = Alzheimer disease; CI = confidence interval; DBP = diastolic blood pressure; GAF = genetic ancestry factor; LCBC = Center for Lifespan Changes in Brain and Cognition; PGS = polygenic risk score; SBP = systolic blood pressure; SNP = single nucleotide polymorphism.

Among the earliest signs of Alzheimer disease (AD) are deficits in memory and orientation, tied to hippocampal dysfunction.¹ Although variants at multiple genetic loci identified in case-control studies are associated with AD risk,² little is known about how these affect individuals devoid of AD diagnosis at an endophenotypic level. Although the major AD genetic risk factor, the *APOE* $\epsilon 4$ allele,^{3,4} has been associated with smaller hippocampi at various ages also in healthy persons,^{5,6} longitudinal data are scarce and restricted to older adults.^{7,8} It is unknown whether polygenic risk scores (PGSs) calculated from established AD risk variants translate to differences in neural characteristics at different life stages in healthy persons. Of interest, higher AD-PGS was recently found related to lower left hippocampal volumes in young adults, also when variants in *APOE* were excluded.⁶ The negative effect of common genetic polymorphisms on late-life disease, including AD, has often been interpreted in terms of age-specific mechanisms.^{9,10} However, there is growing evidence for continuous influence of early life factors on later-life cognitive function and neural foundations.^{11–14} Do AD genetic risk factors exert their effects only at later life stages resulting in brain atrophy and clinical symptoms or affect neural structures throughout the lifespan in the population at large? We hypothesized a relation throughout the lifespan, with higher AD-PGS, including the presence of *APOE* $\epsilon 4$, showing association with lower hippocampal volumes early in life, as an offset effect. We tested whether AD genetic risk factors derived from genome-wide association studies² had an effect on hippocampal volume and volumetric changes in cognitively healthy individuals through the lifespan and whether genetic risk interacted with age.

Methods

Sample

A total of 2,690 valid scans from 1,181 cognitively healthy participants, aged 4.1–95.7 years (mean visit age 39.7 years, SD 26.9 years), were drawn from 5 Norwegian studies. The studies included 4 substudies coordinated by the Center for Lifespan Changes in Brain and Cognition (LCBC); The Norwegian Mother and Child Cohort Neurocognitive Study,¹⁵ Neurocognitive Development,¹⁶ Cognition and Plasticity Through the Lifespan,¹⁷ Neurocognitive Plasticity,¹⁸ and a study run collaboratively by LCBC and Oslo University Hospital, Novel Biomarkers¹⁹ (see e-Methods for details, links.lww.com/NXG/A316). The majority of participants were followed longitudinally, scan intervals ranging 0.2–11.0 years (mean = 2.9 years, SD = 2.6 years). The sample is partly overlapping with a lifespan sample used in previous

reports.^{12,14} All were screened, and dementia, previous stroke with sequela, Parkinson disease, and other neurodegenerative diseases likely to affect cognition were the exclusion criteria, with additional inclusion and exclusion criteria being applied per study (see e-Methods). Sample descriptions for the total sample binned by time points are given in table 1. Additional descriptions including distribution of substudy samples per time point are given in table e-1.

To check for possible recruitment bias/age selectivity, the correlations of age and AD-PGS and *APOE* $\epsilon 4$ allelic variation (coded as both 0, 1, or 2 copies of the $\epsilon 4$ allele and the absence or presence of the $\epsilon 4$ allele, 0 or 1) were calculated. For AD-PGS and age, there was a modest correlation, $r = -0.04$ (confidence interval [CI]: $-0.08, -0.00$), suggesting some age selectivity. No correlations reaching statistical significance or trends toward such were observed for *APOE* $\epsilon 4$ allelic variation, correlations with age being $r = -0.01$ (CI: $-0.05, 0.02$) for *APOE* $\epsilon 4$ coded as no (0)/yes (1) and $r = 0.00$ (CI: $-0.04, 0.04$) for *APOE* coded as no $\epsilon 4$ allele (0)/1 $\epsilon 4$ allele (1)/2 $\epsilon 4$ alleles (2). This suggests that there was no substantial sample selection age bias with regard to genetic risk for AD. The proportion of *APOE* $\epsilon 4$ carriers in the present study appears largely in line with that which would be expected for the population in a Scandinavian country.²⁰

Standard protocol approvals, registrations, and patient consents

The studies were approved by the Regional Ethical Committee of South East Norway. Written informed consent was obtained from participants aged 16 years and older and from parents of participants younger than 16 years, and oral consent was obtained from participants aged 12 years and older.

Genotyping

Buccal swab and saliva samples were collected for DNA extraction, followed by genome-wide genotyping using the Global Screening Array (Illumina, Inc., San Diego, CA). For a full description of genotyping and postgenotyping methods, including quality control and imputation of untyped markers, please see e-Methods (links.lww.com/NXG/A316). AD-PGS of our sample was calculated using the allelic effect sizes from Lambert et al.² Shared single nucleotide polymorphisms (SNPs) between our data and Lambert et al.² were pruned to be nearly independent using the program PLINK²¹ with the following parameters $-\text{clump-p1 } 1.0 -\text{clump-p2 } 1.0 -\text{clump-kb } 500 -\text{clump-r2 } 0.1$. The linkage disequilibrium structure was based on the European subpopulation from the 1000 Genomes Project Phase 3.²² Because of the complexity of the

Table 1 Sample descriptives

	Time point 1 (n = 1,181, 738 F)		Time point 2 (n = 677, 384 F)		Time point 3 (n = 442, 252 F)		Time point 4 (n = 180, 106 F)		Time point 5 (n = 117, 69 F)		Time point 6 (n = 87, 51 F)		Time point 7 (n = 6, 4 F)	
	Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range	Mean	Range
Age	32.5	4.1–89.8	36.5	5.5–91.8	44.4	10.7–93.6	62.6	21.1–95.7	58.7	21.3–82.8	62.6	23.1–84.0	75.6	72.7–79.8
Interval	—	—	2.3	0.2–8.9	4.6	0.4–10.0	2.7	0.6–11.0	1.1	0.8–3.8	2.9	1.8–4.5	4.0	2.8–4.8
Hip. vol	8,127	4,551–11,446	7,940	4,506–10,831	7,922	4,444–10,885	7,549	4,428–10,747	7,733	5,888–10,801	7,482	5,584–10,735	7,024	6,071–7,568
PGS AD	0.214	0.0891–0.368	0.214	0.0943–0.340	0.217	0.0943–0.340	0.213	0.0943–0.323	0.211	0.0943–0.308	0.212	0.0943–0.308	0.206	0.0144–0.231
APOE ε4	0.323	—	0.344	—	0.353	—	0.300	—	0.256	—	0.264	—	0.500	—
ε4 homo	0.029	—	0.030	—	0.029	—	0.022	—	0.026	—	0.023	—	0.167	—

Age and interval are given in years. Interval is interval since first visit. Hippocampus volume (Hip. vol.) denotes number of voxels (mm³) in the hippocampal segmentation bilaterally. Polygenic risk score (PGS) for Alzheimer disease (AD) (threshold $p < 0.5$) for readability of the table given as values multiplied by 1,000, i.e., 0.214 is 0.000214. APOE ε4 denotes the proportion of participants with 1 or more APOE ε4 alleles. ε4 homozygotes (ε4 homo) denote the proportion with 2 ε4 alleles. Note that since the sample derives from a combination of studies, it is not so that those who have the greatest number of time points necessarily have the longest interval since the first visit.

major histocompatibility complex region (build hg19; chr6: 25652429-33,368,333), we removed SNPs in this region except the most significant one before pruning. Previous studies^{23–25} have shown that PGSs constructed using SNPs with association p value < 0.5 from Lambert et al.² have the largest effect on the risk of AD. Hence, we used the same threshold in the pruned set for computing the AD-PGS. To investigate the consistency of results across different p value thresholds, SNPs with $p < 5e-08$ in the pruned set were additionally used for constructing a PGS for our samples. Analyses were recomputed limiting markers to those showing genome-wide significant association (i.e., $p < 5e-08$) with AD risk in Lambert et al.² Because of its known large effect, we computed AD-PGS with and without markers in the APOE region (build hg19; chr19: 44,909,011-45,912,650). To test the effect of APOE itself, we modeled the counts of APOE ε4 alleles directly by counting the number of alternative alleles at rs429358,^{26,27} as 0, 1, or 2. To control the possible effect of vascular factors on the relation between AD-PGS and hippocampus volume, we included PGSs for systolic and diastolic blood pressure (SBP-PGS and DBP-PGS) as additional covariates in our models. Both SBP-PGS and DBP-PGS were computed using the same protocol as that used for AD-PGS, but based on the public data from Liu et al.²⁸ We computed the genetic ancestry factors (GAFs) using principal component methods.²⁹ For the present analysis, only participants of European ancestry were included, excluding 89 persons for whom we had genotype data.

MRI data acquisition

Participants were scanned at a total of 4 Siemens scanners at 2 sites. The pulse sequences used for morphometric analyses were 1 to 2 3D sagittal T1-weighted MPRAGE sequences. For details on the scanners and pulse sequences used, see e-Methods (links.lww.com/NXG/A316). Other MRI volumes were recorded, including sequences intended for and examined by a radiologist, to rule out and medically follow up incidental neuroradiologic findings. Distribution of scans from the different scanners per time point is given in table e-1. See e-Methods for further details.

Image analysis

All scans were reviewed for quality and automatically corrected for spatial distortion.³⁰ Images were processed with the FreeSurfer software package (version 6.0), including automatic hippocampal volumetric segmentation.³¹ To allow assessment of differences between scanners, 24 participants were scanned on all 3 scanners from Oslo University Hospital on the same day. Linear regression analyses were run testing the concordance between hippocampal volumes between scanners, yielding excellent agreement. Thus, including scanner as a covariate in the analyses would account for offset differences between scanners. See e-Methods for further details (links.lww.com/NXG/A316).

Statistical analyses

Analyses were run in R version 3.6.0. General additive mixed models using the package mgcv version 1.8-28 were used to

derive age functions with a random intercept term per participant. Hippocampal volumes were predicted from (1) a smooth function of age and a linear function of AD-PGS, (2) a smooth function of age and linear functions of AD-PGS and presence/absence of 1 or 2 *APOE* $\epsilon 4$ alleles, (3) smooth functions of age and AD-PGS and a tensor interaction term of age and AD-PGS, and (4) a linear function of the presence/absence of 1 or 2 *APOE* $\epsilon 4$ alleles and a smooth interaction between age and presence/absence of 1 or 2 *APOE* $\epsilon 4$ alleles. In all models, scanner, intracranial volume, sex, and the first 5 GAFs, in addition to genotyping batch (1; $n = 1,014$, 2; $n = 166$), were entered as covariates. DBP-PGS and SBP-PGS were added as additional covariates (repeating steps 2 and 4), to further investigate whether any effect was unique to AD-PGS and *APOE*. We did not have any hypothesis regarding differential hemispheric effects, and bilateral hippocampal volumes were entered in analyses. However, we additionally ran analyses for the left and right hippocampi separately (repeating steps 2 and 4), and to allow for comparison with others,⁶ results presented as e-Methods (links.lww.com/NXG/A316). We computed 95% CIs for the effect of PGS and *APOE* allelic variation. The reported effect sizes are mm^3 difference in hippocampal volume per SD of PGS and when carrying 1 or more $\epsilon 4$ alleles vs not carrying any.

Data availability

There are data access restrictions in accordance with ethical guidelines and national data protection legislation. The data will be made available pending reasonable request, appropriate institutional data protection security measures, and ethical approval.

Results

In the lifespan sample, hippocampal volumes increased in early development and declined in older age, as shown in figure 1A. We found a significant negative effect of AD-PGS on hippocampal volume, as shown in figure 1B. The estimated effect of AD-PGS on hippocampal volume was -36.4 mm^3 (CI: $-71.8, -1.0$) associated with 1 sample SD higher AD-PGS. The interaction term between age and AD-PGS did not reach statistical significance. The interaction term between age and AD-PGS represents how the effect of AD-PGS varies with age and is nonlinear in both variables. The point estimate of the interaction was less than 10% of the main effect of AD-PGS for AD-PGS within 2 SDs of the sample mean over the full age range considered. At age 20 years, this interaction term estimated an additional offset effect -0.27 mm^3 (CI: $-2.87, 2.33$) at AD-PGS 1 SD below the mean and 0.23 mm^3 (CI: $-2.39, 2.85$) at 1 SD above the mean, and at age 80 years, the corresponding estimates were 0.96 mm^3 (CI: $-3.12, 5.04$) and -0.86 mm^3 (CI: $-4.96, 3.24$). Of note, this implies that the absolute volume difference between those with lower and higher AD-PGS appears to be about the same in young adults (~ 20 s) as in the oldest adults (~ 80 s), and even the extreme ends of the CIs represent quite moderate differences.

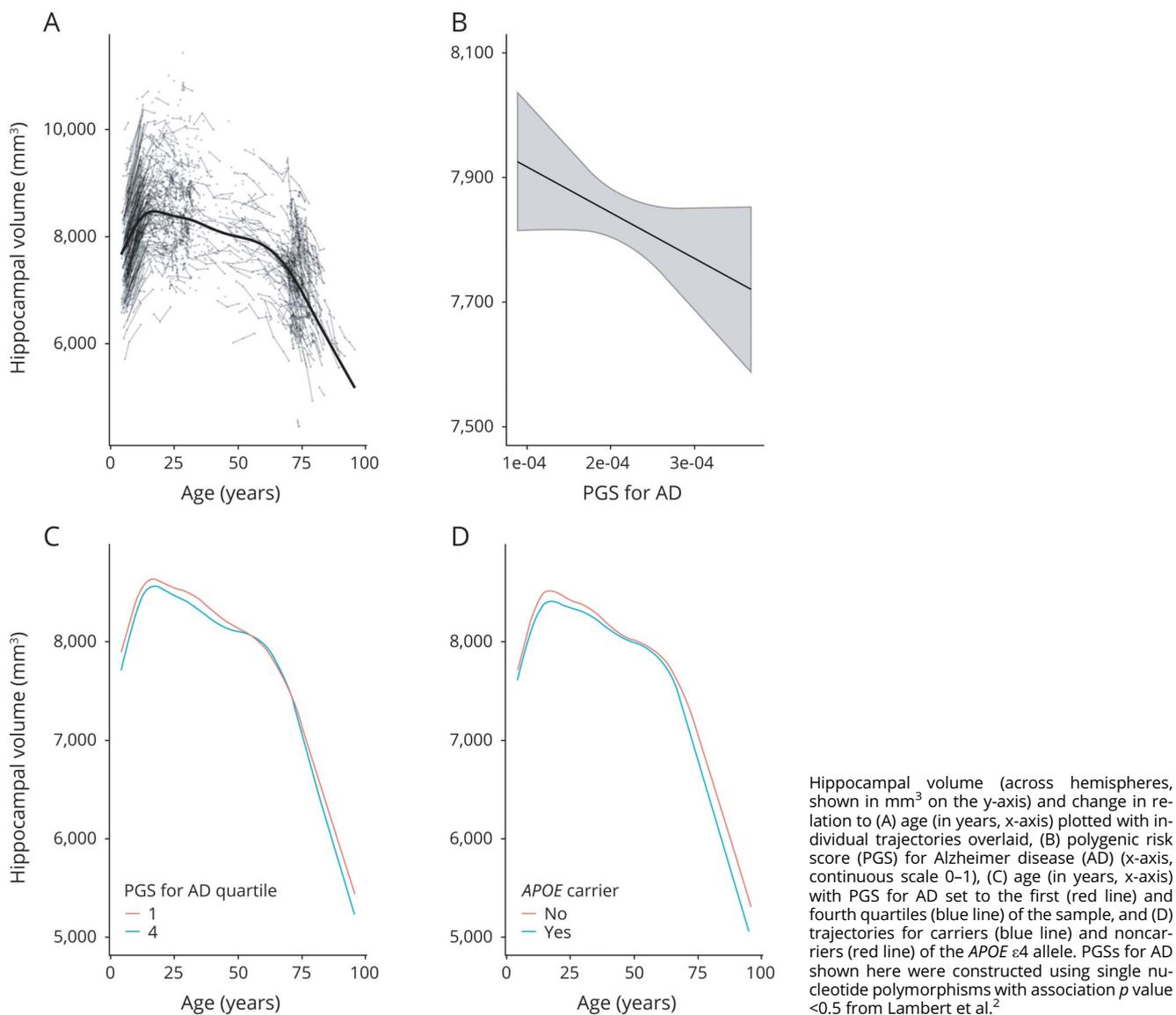
Although the analysis was based on the continuous AD-PGS score, inspection of the age trajectories for values of AD-PGS at 1 SD below and above mean (figure 1C) did not give any consistent indication of greater effects of higher PGS for AD with age. Rather, although there was time after age 60 years when the hippocampal curves merge for the different subgroups, they were otherwise mostly ordered throughout life so that those with PGS for AD 1 SD below mean had the highest volume, whereas those with the PGS for AD 1 SD below mean have the lowest volume. Adding PGSs for blood pressure (DBP and SBP) as additional covariates, the effect of AD-PGS on hippocampal volume was still significant, see e-Methods for further details (links.lww.com/NXG/A316).

Analysis with *APOE* status as the predictor of hippocampal volume likewise showed a significant negative effect of the presence of the $\epsilon 4$ allele. The estimated effect of carrying 1 or more $\epsilon 4$ alleles on hippocampal volume was -107.0 mm^3 (CI: $-182.0, -31.5$). The interaction term between age and *APOE* allelic variance did not reach statistical significance. $\epsilon 4$ allele carriers had smaller hippocampal volumes in young age, as an offset effect. There was not indication of faster atrophy in older age, as carriers in young adulthood (~ 20 s) appeared to display lower hippocampal volumes, a difference that seemed somewhat lower in middle age, but more pronounced again in older adulthood (~ 80 s; figure 1D). At age 20 years, the difference in hippocampal volume between carriers and noncarriers was -101 mm^3 (CI: $-224, 23$); at age 50 years, this difference was -22.6 mm^3 (CI: $-186, 141$); and at age 80 years, it was -278 mm^3 (CI: $-478, -77.7$). Adding PGSs for blood pressure (DBP and SBP) as additional covariates did not weaken the effect of *APOE* status on hippocampal volume, see e-Methods for further details (links.lww.com/NXG/A316).

AD-PGS excluding the *APOE* region did not show a significant effect on hippocampal volume, although a trend in the same direction as the full model was observed (estimate -31.8 mm^3 , CI: $-67.4, 3.92$). There was no evidence for an age interaction on using the AD-PGS excluding markers in the *APOE* region. The estimated interactions corresponding to values of AD-PGS excluding the *APOE* region at 1 SD below the sample mean varied from -0.1 mm^3 (CI: $-1.1, 2.2$) at age 20 years to 0.2 mm^3 (CI: $-2.2, 2.6$) at age 80 years, and the corresponding values at other ages were very similar. For values of AD-PGS excluding the *APOE* region at 1 SD above the sample mean, the interaction varied from 0.1 mm^3 (CI: $-1.4, 1.6$) at age 20 years to -0.1 mm^3 (CI: $-2.6, 2.3$) at age 80 years, and also here the corresponding values at other ages were very similar.

Recomputing the analyses with an AD-PGS limited to SNPs only showing genome-wide significant association (i.e., $p < 5e-08$) with AD risk in Lambert et al.² confirmed a significant negative effect of AD-PGS on hippocampal volume ($-44.8 \text{ mm}^3/\text{AD-PGS SD}$, CI: $-80.7, -8.9$). This result was reduced to a trend when excluding markers in the *APOE* region ($-35.6 \text{ mm}^3/\text{PGS SD}$, CI: $-76.7, 5.5$). However, significant age interactions appeared in the PGS analyses limited

Figure 1 Hippocampal volume in relation to age and genetic risk



to genome-wide significant markers, both with and without markers in the *APOE* region (figure 2). Especially when excluding the *APOE* region, there seemed to be a somewhat more negative effect of higher AD-PGS on hippocampal volume in older age (above 80 years). For a more detailed description of these interaction effects, see e-Methods (links. lww.com/NXG/A316).

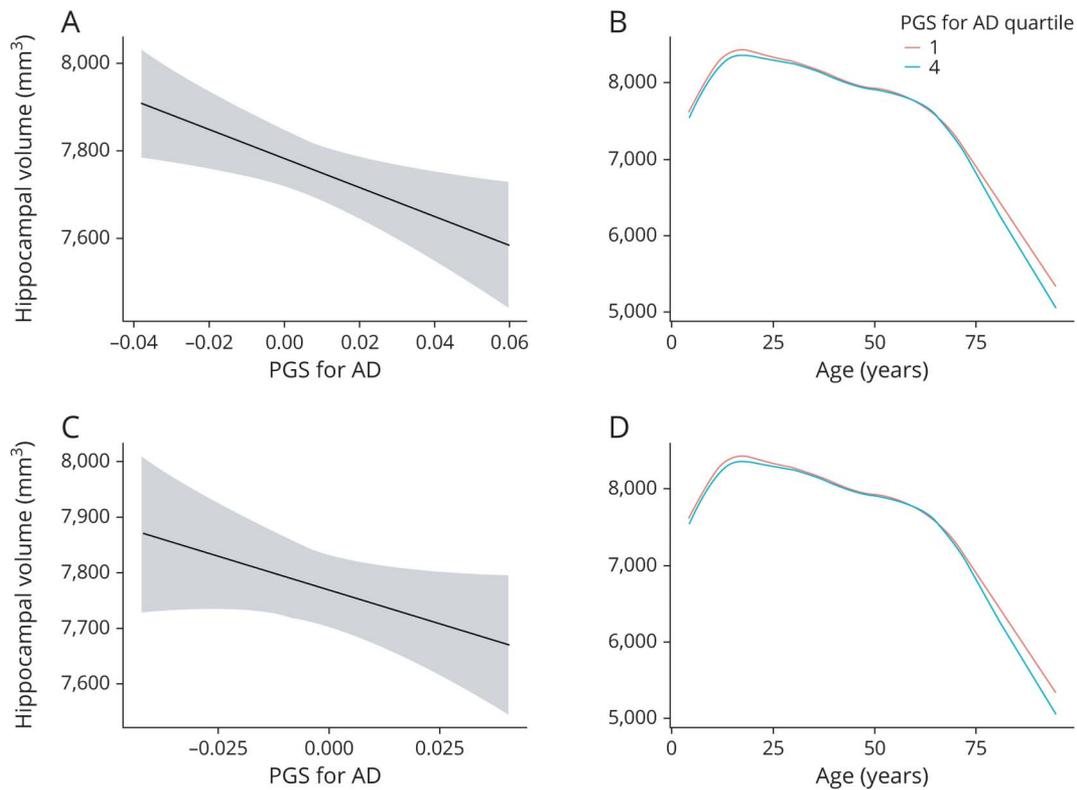
Discussion

This study shows that genetic risk for AD is associated with effects on hippocampal volume throughout life, having neurodevelopmental offset effects observable from childhood in cognitively healthy well-functioning participants. Higher AD-PGS and carrying the *APOE* ϵ 4 allele specifically were associated with lower developmental hippocampal volume offsets,

and persons with higher genetic risk for AD remained having lower hippocampal volumes with age. Notably, this main effect was consistently observed for PGSs constructed using SNPs with 2 different association *p* values.² The association of hippocampal volume and AD-PGS through the lifespan found here can contribute to build knowledge on the lifespan trajectories of polygenic AD risk at an endophenotypic level. However, and as expected, the effect was not large, and when using the AD-PGSs computed without the *APOE* region as predictor, the effect was reduced to a trend.

A negative cross-sectional association between hippocampal volumes and polygenic risk for AD—also excluding the *APOE* region—has been previously reported in a smaller sample of young adults,⁶ suggesting that the relationship between hippocampal volume and AD risk results from multiple genetic factors and not exclusively variability in the *APOE* region. We

Figure 2 Hippocampal volume and PGS for AD limited to SNPs showing genome-wide significant association



Upper panel: including *APOE*. Hippocampal volume (across hemispheres, shown in mm³ on the y-axis) and change in relation to (A) AD-PGS including the *APOE* region as calculated based on effect sizes from Lambert et al.² at $p < 5e-08$ (x-axis, continuous scale 0–1). (B) Age (in years, x-axis) with AD-PGS at $p < 5e-08$ including the *APOE* region set to the first (red line) and fourth (blue line) quartiles of the sample. Lower panel: excluding *APOE*. Hippocampal volume (across hemispheres, shown in mm³ on the y-axis) and change in relation to (C) AD-PGS excluding the *APOE* region as calculated based on effect sizes from Lambert et al.² at $p < 5e-08$ (x-axis, continuous scale 0–1). (D) Age (in years, x-axis) with AD-PGS at $p < 5e-08$ excluding the *APOE* region set to the first (red line) and fourth (blue line) quartiles of the sample. AD = Alzheimer disease; PGS = polygenic risk score; SNP = single nucleotide polymorphism.

do not consider those findings in conflict with the present, where a trend toward the same was observed. However, as can be seen from the associations reported, the effect of *APOE* on hippocampal volumes in cognitively healthy persons of varying ages appears stronger than that of other genetic variants.

Having low vs high genetic risk for AD was associated with roughly equivalent difference in hippocampal volume at age 25 and 80 years. In the current study, the effect of the PGS constructed using SNPs with association p value < 0.5 from Lambert et al.,² shown to have the largest effect on the risk of AD,^{23–25} did not significantly interact with age. However, the PGS effect was not equally apparent at all ages among older adults, and notably, age interactions were observed for the same PGS constructed using SNPs with the lowest association p value, also when excluding variants in the *APOE* region. Possibly, different extent of effects at different ages may illuminate why one small recent study found associations of an AD genetic risk score and hippocampal atrophy over a 2-year interval in healthy older adults ($n = 45$), although no association with individual variation in volume was seen at baseline ($n = 66$).⁸ AD-PGS did correlate modestly with age in our sample, and there may be some sample selection bias in healthy older age samples, so it would be interesting to see

whether age interactions could be consistently observed if persons were followed for even longer and at higher ages. It should be noted that a limitation of the current study is that sample density was lower in the middle of the lifespan than in young and older age groups. We cannot, based on the current sample, exclude a mixture of neurodevelopmental offset and aging effects, although the neurodevelopmental offset effects were the most consistently supported by the present results.

There was only a trend for effects of *APOE* allelic variants to interact with age, and this trend was not clearly indicative of faster atrophy for *APOE* $\epsilon 4$ carriers in older age. Cross-sectional studies have shown that *APOE* $\epsilon 4$ ^{3,4} carriers tend to have lower hippocampal/medial temporal lobe volumes at various ages also in healthy persons,⁵ and notably, this has been identified even in neonates.³² However, it has been difficult to interpret whether these developmental structural brain differences actually do represent long-term risk factors, as longitudinal imaging data have been lacking. The only such study we know of was restricted to older adults aged 55–75 years,⁷ showing greater hippocampal atrophy in $\epsilon 4$ carriers across a 5-year interval. The present study hence partly confirms, yet nuances and extends previous reports, in showing a main effect through the lifespan, i.e., also an offset effect.

The present study adds to the evidence for early life factors exerting a continuous influence on later-life function,¹¹ showing that genetic factors established to influence late-life neural and cognitive disease work in part in a temporally stable and dimensional manner. That is, genetic risk factors for AD seem not to only manifest at late life in hippocampal atrophy and clinical symptoms but appear to start influencing the neural structures on which cognition rely early on, through the entire lifespan and in the population at large. In this regard, currently observed effects of the *APOE* ε4 allele cannot well be interpreted only within the framework of antagonistic pleiotropy,^{9,10} as this allelic variant appears to have in part similar effects on neural structures of memory function at different ages. As for AD-PGS, currently observed effects are of similar magnitude in young adulthood as in much of older adulthood, meaning that explanations evoking different effects on the hippocampus at different ages may be incomplete.

This does not necessarily imply that effects on behavior may be readily observed through the lifespan. A number of studies have indeed not observed effects of *APOE* status on standard neuropsychological memory tests.^{33,34} However, as recently reported, this does not mean that effects of genetic risk for AD do not manifest early, in more fine-grained behaviors dependent on hippocampal circuitry, such as spatial navigation.³⁴ A related yet different account is the magnification or resource modulation hypothesis,^{35–37} stating that genetic effects are magnified in persons with constrained neural resources, such as older—and putatively also developing—individuals. Here, it is assumed that the function relating brain resources to cognition is nonlinear so that genetic differences exert increasingly larger effects on performance with lesser neural substrates. Smaller hippocampi can be seen as a risk factor. The idea that persons starting with larger brain structures may preserve mental status despite developing AD plaques, and may be said to have a greater reserve, is supported by older postmortem and clinical data.³⁸ Within a brain reserve account, such differences in brain structure may relate to differences in tolerance to pathology before one falls under a functional threshold.³⁹ Hence, these differences may become readily functionally apparent only in older age, but they likely are there to begin with and throughout life in a stable manner. This supports a lifespan model of dementing disorder,⁴⁰ where effects of common genetic variants in part work in a stable manner to be one of several factors affecting risk for cognitive decline and neurodegenerative disease.

In conclusion, endophenotypic expression of genetic risk for AD may be seen in a dimensional and lifespan perspective, not being confined to clinical populations or older age. This emphasizes that a broader population and age range may be relevant targets for attempts to prevent AD. To help define targets for prevention, future studies of the genetic factors established to influence AD are needed. These should investigate which of these factors have an age-independent influence, across manifestations of health and disease, and which may have more pronounced effects in aging.

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Disclosure

The authors report no disclosures. Go to Neurology.org/NG for full disclosures.

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Appendix Authors

Name	Location	Contribution
Kristine B. Walhovd, PhD	University of Oslo, Norway	Designed and conceptualized the study, analyzed the data, and drafted the manuscript for intellectual content
Anders M. Fjell, PhD	University of Oslo, Norway	Designed and conceptualized the study and revised the manuscript for intellectual content
Øystein Sørensen, PhD	University of Oslo, Norway	Role in conceptualization of the study and building of statistical models, interpreted the data, and revised the manuscript for intellectual content
Athanasia Monika Mowinckel, PhD	University of Oslo, Norway	Processing and interpretation of the data and revised the manuscript for intellectual content
Céline Sonja Reinbold, PhD	University of Oslo, Norway	Genotyping analyses and revised the manuscript for intellectual content
Ane-Victoria Idland, MD	University of Oslo, Norway	Role in the acquisition of data and revised the manuscript for intellectual content

Continued

Appendix (continued)

Name	Location	Contribution
Leiv Otto Watne, MD, PhD	University of Oslo, Norway	Role in the acquisition of data and revised the manuscript for intellectual content
Andre Franke, PhD	Christian-Albrechts-University of Kiel, Germany	Genotyping analyses and revised the manuscript for intellectual content
Valerija Dobricic, PhD	University of Lübeck, Germany	Genotyping analyses and revised the manuscript for intellectual content
Fabian Kilpert, PhD	University of Lübeck, Germany	Genotyping analyses and revised the manuscript for intellectual content
Lars Bertram, MD	University of Oslo, Norway; University of Lübeck, Germany	Genotyping analyses, interpretation of data, suggestions for analyses, and revised the manuscript for intellectual content
Yunpeng Wang, PhD	University of Oslo, Norway	Design and conceptualization of the study, analyzed and interpreted the genetic data, and revised the manuscript for intellectual content

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