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RESEARCH PAPER

Candidate-gene analysis of white matter hyperintensities on neuroimaging

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Received 13 October 2014

Revised 8 March 2015

Accepted 11 March 2015

Published Online First

2 April 2015

ABSTRACT

Background White matter hyperintensities (WMH) are a common radiographic finding and may be a useful endophenotype for small vessel diseases. Given high heritability of WMH, we hypothesised that certain genotypes may predispose individuals to these lesions and consequently, to an increased risk of stroke, dementia and death. We performed a meta-analysis of studies investigating candidate genes and WMH to elucidate the genetic susceptibility to WMH and tested associated variants in a new independent WMH cohort. We assessed a causal relationship of WMH to methylene tetrahydrofolate reductase (MTHFR).

Methods Database searches through March 2014 were undertaken and studies investigating candidate genes in WMH were assessed. Associated variants were tested in a new independent ischaemic cohort of 1202 WMH patients. Mendelian randomization was undertaken to assess a causal relationship between WMH and MTHFR.

Results We identified 43 case-control studies interrogating eight polymorphisms in seven genes covering 6,314 WMH cases and 15,461 controls. Fixed-effects meta-analysis found that the C-allele containing genotypes of the aldosterone synthase CYP11B2 T(−344)C gene polymorphism were associated with a decreased risk of WMH (OR=0.61; 95% CI, 0.44 to 0.84; p=0.003). Using mendelian randomisation the association among MTHFR C677T, homocysteine levels and WMH, approached, but did not reach, significance (expected OR=1.75; 95% CI, 0.90–3.41; observed OR=1.68; 95% CI, 0.97–2.94). Neither CYP11B2 T(−344)C nor MTHFR C677T were significantly associated when tested in a new independent cohort of 1202 patients with WMH.

Conclusions There is a genetic basis to WMH but anonymous genome wide and exome studies are more likely to provide novel loci of interest.

INTRODUCTION

White matter hyperintensities (WMH) are defined as diffuse white matter abnormalities detected on T2-weighted or fluid attenuated inversion recovery (FLAIR) MRI and appearing as regions of low attenuation on brain CT scans.¹ They are a common radiological finding, particularly in older individuals, with a reported prevalence of up to 95%.^{2,3} These lesions, of presumed vascular origin, may represent an endophenotype for small cerebral vessel diseases such as stroke and dementia; thus, WMH could be used in early diagnosis and guided

management of these conditions.⁴ WMH has consistently been associated with increasing age and hypertension,⁵ as well as smoking,⁶ previous stroke⁷ or TIA,⁸ and elevated homocysteine (Hcy) levels.^{9,10} Studies have reported high heritability estimates ranging between 55% and 71%,^{11–13} implying a significant genetic component to WMH development.

To date, a single WMH GWAS has been published by the CHARGE consortium¹⁴ that identified six single nucleotide polymorphisms (SNPs) in a novel locus on chromosome 17q25 associated with WMH burden in stroke-free participants.¹⁴ The most significantly associated SNP on 17q25 was rs3744028 with a reported p value of 1.0×10^{-9} after adjustment for hypertension.¹⁴ This association between the 17q25 locus and WMH has recently been tested in a cohort of ischaemic patients with stroke, where it has replicated in association with WMH volume but not lacunar stroke status.¹⁵ The latter may suggest that these two diseases have distinct pathogeneses of cerebral microangiopathy. Rs3744028 was again found to be significantly associated with increased WMH burden (effect size=0.12; SE=0.04; p=0.003), although this SNP was not the most significantly associated in this study population (rs9894383; effect size=0.13; SE=0.04; p=0.0006).

Several statistically underpowered small candidate gene studies on WMH have been published, but the results remain invalid due to low power. By consolidating data from these smaller studies, a literature-based meta-analysis is considered to be the next best way to increase power and find a true genetic risk association. We conducted a comprehensive meta-analysis of all case-control studies investigating candidate genes in WMH, tested our findings in a new independent WMH cohort and sought to identify a causal relationship with methylene tetrahydrofolate reductase (MTHFR).

METHODS

A comprehensive search strategy in electronic databases (PubMed, Google Scholar, Embase) was undertaken using a range of search terms for WMH (*leukoaraiosis, white matter hyperintensities, white matter lesions, white matter disease, age-related white matter changes, homocysteine, hyperhomocysteinaemia*) in combination with the



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To cite: Tran T, Cotlarciuc I, Yadav S, et al. *J Neurol Neurosurg Psychiatry* 2016;**87**:260–266.

Boolean operator AND/OR (*genetics, genotype, genes or polymorphism*). Further searches were conducted for each gene identified, using specific gene names combined with the WMH search terms. Additional studies were found by hand searching reference lists of relevant papers. For duplicate papers, the largest cohort was selected. A variety of different methods were utilised to quantify WMH levels or volumes, but in general visual rating scales were more commonly used than automated programmes. Most papers reported genotype frequencies within the Hardy-Weinberg equilibrium. Our study complied with PRISMA guidelines.

Study selection

Inclusion criteria were: (1) case-control studies where WMH was reported as a grade on a standardised scale or as a volume, (2) WMH was objectively confirmed by MRI or CT brain scans, (3) genotype frequency was reported for WMH cases and controls. Studies were excluded if they did not explicitly distinguish WMH from other brain lesions such as lacunes and microinfarcts. For the Mendelian randomisation part of this study, additional selection was based on plasma Hcy levels for cases and controls in the studies of participants of European descent reporting SDs associated with mean Hcy levels.

Data extraction

Some studies quantified WMH grade on a standardised scale and presented this as dichotomous data, so for each genotype the number of participants in the highest and lowest WMH grade groups was extracted. Where studies subdivided WMH into two categories: deep (or subcortical) WMH and periventricular WMH, without providing data for WMH as a whole, data for the deep WMH was analysed. In cases where the scale cut-off could be chosen, the upper grade group included Fazekas scale 2 or 3 or equivalent.¹⁶ For continuous WMH grade data, the mean grade and SD for each genotype were taken, and for studies presenting WMH data as a volume, we extracted the mean volume and SD for each genotype.

Data analysis for meta-analysis

Data were analysed using Review Manager V.5.2. Using a Mantel-Haenszel statistical method, a pooled OR and 95% CI were calculated for each SNP-WMH association. Statistical significance was set at $p < 0.05$. Where significant heterogeneity was detected, a random-effects analysis model was utilised in order to account for this interstudy heterogeneity. In all other cases, a fixed effects model was used. Heterogeneity was assessed with an I^2 test for each meta-analysis (significance set at $p < 0.10$) and an iterative analysis was performed where significant heterogeneity was found. Publication bias was assessed with Funnel plots and by performing Egger's regression analysis (two-tailed tests) using Comprehensive Meta-Analysis V.2.0 (CMA).

Data analysis for Mendelian randomisation

Mendelian randomisation allows the testing of a causal effect of observed data in the presence of confounding factors. Review Manager was used to calculate an OR and 95% CI for MTHFR-WMH grade association using the TT vs CC model so as to be in keeping with the model used by Casas *et al*¹⁷ who report a weighted mean difference in the Hcy level between TT and CC-genotype to be 1.93 $\mu\text{mol/L}$ in their meta-analysis. A pooled mean difference in Hcy levels between cases with WMH and controls was calculated and then converted into a corresponding OR of WMH for that specific increase in Hcy level

using CMA software. The expected OR was then calculated using the following formula:¹⁷

$$\text{Expected OR} = X^{Y/Z}$$

where X=the OR of risk of WMH for a Z $\mu\text{mol/L}$ increase in plasma Hcy levels.

And Y=the mean difference in the Hcy level ($\mu\text{mol/L}$) between TT and CC-genotype participants.

To calculate the 95% CI for the expected OR, we took the natural log of this number to determine the logged OR. The 95% CI for this logged OR is calculated by taking $1.96 \times \text{SE}$ on either side of this logged OR.¹⁸ The SE is taken as the square root of the sum of the reciprocals of the number of cases and controls. The exponential function in Excel was used to convert the upper and lower CI limits into the 95% CI limits for the original OR.

Replication of the associated genetic variants

Any associated genetic variants were tested in a cohort of 1202 ischaemic stroke cases of European ancestry with genome-wide genotyping available ('MGH,' 'ISGS,' 'ASGS,' 'SWISS' cohorts) that was previously used to replicate chromosome 17q25 locus association with WMH.¹⁵ In this cohort, WMH volume (WMHv) was measured using a previously validated, semiautomated volumetric method (facilitated by *MRIcro*, University of Nottingham School of Psychology, Nottingham, UK; <http://www.mricro.com>).¹⁹ For this analysis, MRI scans obtained from a 1.5 T scanner were converted from DICOM into Analyse format, and the contiguous, supratentorial axial T2 FLAIR sequences were cross-referenced with diffusion-weighted images (DWI) and examined to exclude hyperintensities that represent oedema, acute ischaemia or chronic infarcts. WMHs were manually outlined as regions of interest, and their intersections with automatically derived intensity thresholds were manually examined and touched up by a trained reader. The total WMHv was calculated by doubling the measurement from the hemisphere unaffected by stroke, or by adding bilateral WMHv values in participants with infratentorial DWI lesions. To control for variation in head size, the intracranial area (ICA) was calculated from two middle sagittal T1 slices and used to normalise WMHv by multiplying it by the individual-to-the-mean ICA ratio.^{20, 21} Specific study characteristics and genotyping information of this cohort are previously described.¹⁵

RESULTS

Our initial search identified 1248 studies with 20 additional records from references of relevant articles. Removal of duplicates and matches to our predefined inclusion and exclusion criteria resulted in 43 studies which had available data for meta-analysis interrogating nine polymorphisms in 10 different genes (figure 1). The majority used MRI to assess WMH in (mostly Caucasian) participants aged between 60 and 80 years. Most of the genes studied were involved in the renin-angiotensin-aldosterone system (ACE, angiotensinogen (AGT), angiotensin II receptor 1, aldosterone synthase). The other identified genes had roles in Hcy levels (MTHFR), antiatherosclerosis (paraoxonase 1) and cholesterol regulation (apolipoprotein E). Other gene polymorphisms studied were brain-derived neurotrophic factor/Val66Met²² and nitric oxide synthase 3/G894T,^{23, 24} but some studies had yet to be replicated²² and others did not have data available for meta-analysis²⁴ (table 1).

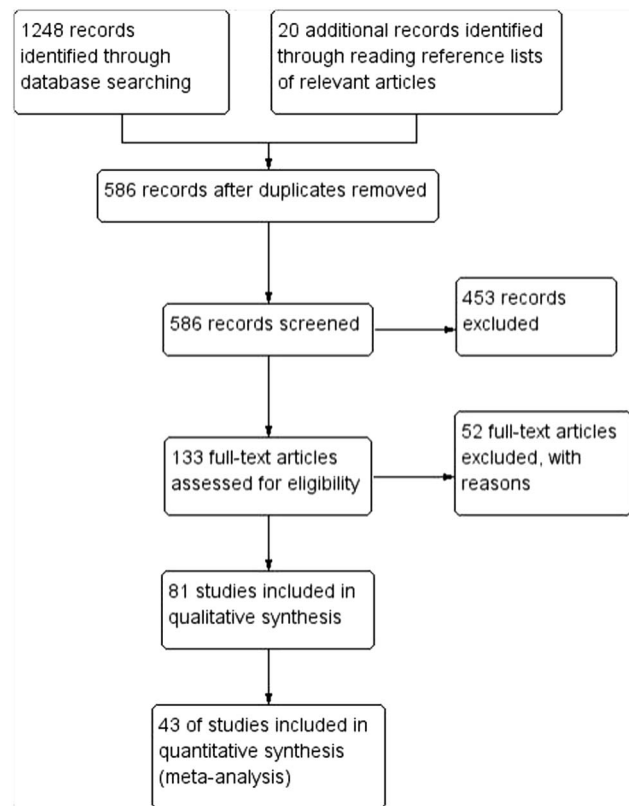


Figure 1 PRISMA flow chart demonstrating the search strategy.

MTHFR677 cytosine/thymine (TT vs CT/CC)

Six studies investigated the association between the MTHFR C677T polymorphism (TT vs CT/CC) and WMH (n=4002).^{25–30} Five studies (n=2988) assessed the genotype difference between the lower and upper WMH grade groups^{25 26 28–30} (figure 2A) and one study (n=1014) compared the mean WMH volume between different genotypes²⁷ (figure 2B). A fixed-effects meta-analysis demonstrated a trend of increased risk burden of WMH with MTHFR TT compared to the CT/CC genotype (OR=1.19; 95% CI 0.95 to 1.51; p=0.14; I²=28%; p=0.24) when the totality of data was plotted (see figure 2). Only one study measured the WMH volume, thus limiting further analysis (standardised mean difference=-0.09; 95% CI -0.29 to 0.11; p=0.37).

Aldosterone synthase CYP11B2(-344)C (CC/CT vs TT)

Two studies (n=1153) evaluated the association between the aldosterone synthase CYP11B2(-344)C polymorphism and

dichotomous graded WMH, and the fixed-effects meta-analysis demonstrated that the C-allele-containing genotypes were at a reduced risk of white matter lesions (OR=0.61; 95% CI 0.44 to 0.84; p=0.003; I²=0%; p=0.70).^{31 32}

Apolipoprotein E (ε4 allele-containing genotypes vs others)

There were 31 studies/substudies that investigated the association between apoE (ε4 allele-containing genotypes vs other genotypes) and WMH (n=11 270).^{29 33–57} Twenty-two of these studies (n=7622) assessed the genotype difference between the lower and upper WMH grade groups,^{29 33–41 50–58} three studies (n=187) compared the mean WMH grade^{42–44} and six studies (n=3461) compared the mean WMH volume between different genotype groups.^{45–49}

The fixed-effects meta-analysis demonstrated no association between apoE ε4-allele carriage status and having severe WMH on neuroimaging (WMH grade, dichotomous data, OR=0.98; 95% CI 0.87 to 1.11; p=0.78; I²=5%; p=0.39). Within participants with WMH, there was no significant predominance of the ε4 allele-containing genotypes (WMH grade, continuous data, pooled standardised mean difference=0.29; 95% CI -0.03 to 0.61; p=0.07; I²=0%; p=0.87; WMH volume, pooled standardised mean difference=0.06; 95% CI -0.02 to 0.14; p=0.14; I²=47%; p=0.09).

ApoE/ε2 allele-containing genotypes vs others

Three studies (n=817) evaluated the risk of WMH in apolipoprotein E ε2-containing genotypes compared to other apolipoprotein E genotypes^{38 41 52} and a random-effects meta-analysis reported no significant association (OR=1.42; 95% CI 0.46 to 4.43; p=0.54) but significant heterogeneity was detected (I²=84; p=0.002). Iterative analysis revealed that the source of inter-study heterogeneity was attributable to Smith 2004,⁵² the exclusion of which resulted in pooled OR=2.59; 95% CI 1.60 to 4.19; p=0.0001; I²=0%; p=0.92.

ACE (DD vs ID/II)

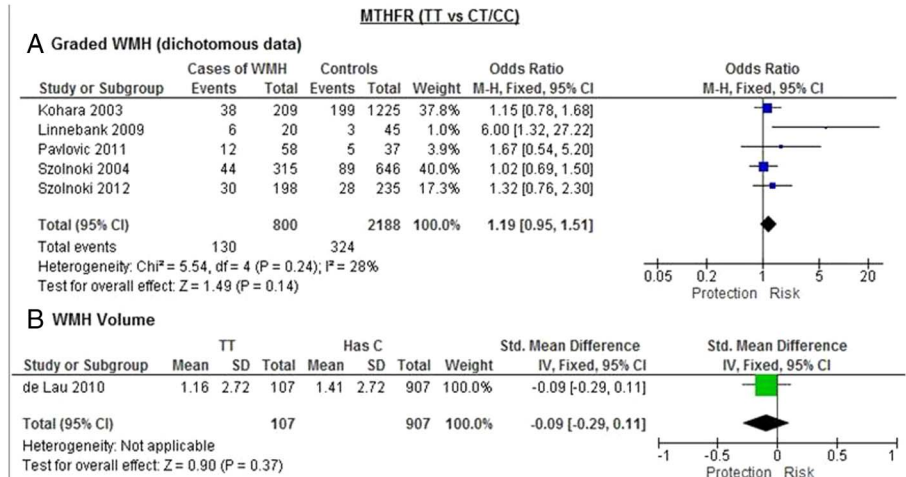
Nine studies/substudies evaluated the association between ACE (DD vs ID/II model) and WMH (n=2615).^{29 31 33 59–63} Eight studies (n=2121) assessed the genotype difference between the lower and upper WMH grade groups^{29 31 33 60–63} and 1 study (n=494) compared the mean WMH volume between different genotypes.⁵⁹ The random-effects meta-analysis suggested no increased risk of WMH with ACE DD-genotype compared to those with ID or II genotype (OR=1.46; 95% CI 0.92 to 2.31; p=0.11), but there was substantial heterogeneity detected between studies (I²=67%; p=0.004). No one study contributed to the heterogeneity as determined by iterative analysis. One study assessed WMH volume and also reported non-significance

Table 1 Summary table demonstrating each gene, polymorphism, model used, number of cases and controls and resulting OR, CI and p values

Gene	Polymorphism	Model used	Cases	Controls	OR	95% CI	p Value
Apolipoprotein	E4	E4 carriers vs non-carriers	2614	5008	0.98	0.87 to 1.11	0.78
Apolipoprotein	E2	E2 carriers vs non-carriers	248	569	1.42	0.46 to 4.43	0.54
ACE	Insertion/deletion	DD vs ID/II	756	1365	1.46	0.92 to 2.31	0.11
MTHFR	C677T	TT vs CT/CC	800	2188	1.19	0.95 to 1.51	0.14
Angiotensinogen	M235T	TT/MT vs MM	328	806	1.12	0.84 to 1.50	0.44
Angiotensinogen II receptor 1	A1166C	CC vs AC/AA	105	354	1.23	0.59 to 2.54	0.58
Aldosterone synthase CYP11B2	T(-344)C	CC/CT vs TT	197	956	0.61	0.44 to 0.84	0.003
Paraoxonase 1	L55M	LL/LM vs MM	77	266	1.42	0.61 to 3.28	0.41

MTHFR, methylene tetrahydrofolate reductase.

Figure 2 Meta-analysis, forest plot and pooled OR of risk from studies investigating the association between WMH and methylene tetrahydrofolate reductase, MTHFR (TT vs CT/CC, recessive model). (A) Graded WMH, dichotomous data (B) WMH volume. MTHFR, methylene tetrahydrofolate reductase; WMH, white matter hyperintensities.



(standardised mean difference = -0.07; 95% CI -0.27 to 0.13; $p = 0.48$).

Angiotensinogen Met235Thr (TT/MT vs MM)

Four studies ($n = 1134$) evaluated the association between AGT M235T (TT/MT vs MM model) and dichotomous graded WMH and fixed-effects meta-analysis found no association between them (OR = 1.12; 95% CI 0.84 to 1.50; $p = 0.77$; $I^2 = 0\%$; $p = 0.44$).^{31 60 61 64}

Angiotensin II receptor 1 A1166C (CC vs AC/AA)

Two studies ($n = 459$) investigated whether the angiotensin II receptor 1 (AGTR1) A1166C polymorphism was associated with dichotomous WMH grade. Using the dominant model (CC vs AC/AA) and a fixed-effects analysis, our meta-analysis found no association (OR = 1.23; 95% CI 0.59 to 2.54; $p = 0.58$; $I^2 = 23\%$; $p = 0.26$).^{31 60}

Paraoxonase 1 L55M (LL/LM vs MM)

Two studies ($n = 343$) evaluated the association between the paraoxonase 1 (PON1) gene and dichotomous graded WMH and fixed-effects meta-analysis found no association between them (OR = 1.42; 95% CI 0.61 to 3.28; $p = 0.41$; $I^2 = 0\%$; $p = 0.33$).^{65 66}

Mendelian randomisation

Ninety-seven studies and five additional records were identified in the search for papers investigating the difference in plasma Hcy levels between WMH cases and controls. After 21 duplicate records were removed, the remaining 81 were screened and 77 were excluded according to the predefined inclusion and exclusion criteria.

Ethnic differences in plasma Hcy levels are well documented with East Asians consistently reported to have significantly lower Hcy levels compared to Caucasians⁶⁷⁻⁷⁰ (table 2). Given these ethnic disparities, we considered it appropriate to exclude studies of East Asian (ie, Japanese, Korean) participants from the Mendelian randomisation, as the majority of studies were conducted in participants of European descent.

The remaining four studies covering 745 Caucasian participants were meta-analysed and a comparison of WMH cases vs controls found a pooled mean difference in plasma Hcy levels of 3.71 $\mu\text{mol/L}$ (95% CI 2.79 to 4.63; $p < 0.00001$; $I^2 = 0\%$) (figure 3). CMA V2.0 software was used to calculate the corresponding pooled OR of risk of WMH for this mean difference in Hcy levels using a fixed effects analysis model, OR = 2.93 (95% CI 2.18 to 3.94).

In a meta-analysis of 42 studies, we had previously examined the effect of MTHFR on plasma Hcy levels in healthy participants ($n = 15\ 635$) and reported the weighted mean difference in the Hcy level between TT and CC-genotype to be 1.93 $\mu\text{mol/L}$ (95% CI 1.38 to 2.47; $p < 0.0001$).¹⁷ Using these three pieces of data, the expected OR was calculated using the following formula:

$$\text{Expected OR} = 2.93^{1.93/3.71} = 1.75$$

where:

- ▶ 2.93 is the OR of risk of WMH for a 3.709 $\mu\text{mol/L}$ increase in plasma Hcy levels,
- ▶ 1.93 is the mean difference in the Hcy level ($\mu\text{mol/L}$) between TT and CC-genotype participants.

Table 2 Ethnic differences in plasma homocysteine level between East Asian and Caucasian participants

Study	Sample	n	Mean tHcy \pm SD ($\mu\text{mol/L}$)	p Value	Association
Anand <i>et al</i> ⁶⁸	Europeans	326	10.0 \pm 3.8	0.02	Chinese had significantly lower Hcy levels.
	Chinese	317	9.2 \pm 3.8		
Carmel <i>et al</i> ⁶⁹	White	237	14.8*	<0.05	Whites had higher Hcy concentrations than Asian-Americans.
	Asian-Americans	68	12.8*		
Senaratne <i>et al</i> ⁷⁰	Caucasians	106	10.8 \pm 0.6	<0.001	East Asians had significantly lower plasma Hcy compared to Caucasians.
	East Asians (Chinese, Japanese)	17	7.6 \pm 0.5		

*SD not reported by study.
tHcy, total homocysteine.

Model	Study name	Statistics for each study						
		Difference in means	Standard error	Variance	Lower limit	Upper limit	Z-Value	p-Value
	Censori 2007	4.300	1.187	1.409	1.974	6.626	3.623	0.000
	Clarke 2000	2.000	1.146	1.312	-0.245	4.245	1.746	0.081
	Hogenvost 2002	4.000	0.709	0.502	2.611	5.389	5.643	0.000
	Pavlovic 2011	4.000	0.981	0.962	2.078	5.922	4.078	0.000
Fixed		3.709	0.471	0.222	2.785	4.633	7.869	0.000

Figure 3 Meta-analysis of studies investigating the mean difference in homocysteine levels ($\mu\text{mol/L}$) between white matter hyperintensities (WMH) cases and controls.

To calculate the 95% CI for the expected OR of 1.75, we took the natural log of 1.75 to get the logged OR of 0.56. The 95% CI for this logged OR was -0.108 to 1.227 and was calculated by taking $1.96 \times \text{SE}$ on either side of 0.56.¹⁸ The SE was 0.34, which was calculated as the square root of the sum of the reciprocals of the number of cases and controls. Using the exponential function in excel, these limits were converted into the 95% CI limits for the original OR of 1.75 giving EXP (-0.108) = 0.897 to EXP (1.227) = 3.412. From our meta-analysis of two studies investigating the association between MTHFR (TT vs CC) and WMH grade, the observed OR for WMH was 1.68 (95% CI 0.97 to 2.94, $p=0.07$). Despite the results not reaching statistical significance, the similarity between the expected and observed ORs supports a likely causal relationship between MTHFR and WMH.

Replication of associated genetic variants

We examined the association between CYP11B2T(-344)C and MTHFR C677T polymorphisms and WMH quantified on the MRI using a validated, semiautomated volumetric protocol in an independent cohort of 1202 ischaemic stroke cases with WMH. There was no association between either polymorphism and the WMH volume in this cohort (CYP11B2T(-344)C, $P=0.5755$; MTHFR C677T, $P=0.68$).

DISCUSSION

In this largest study to date of candidate genes in WMH burden, we interrogated 6253 WMH cases and 15 239 controls for eight polymorphisms in seven genes (APO E/ $\epsilon 4$ and $\epsilon 2$, ACE insertion/deletion, MTHFR C677T, AGT M235T, AGTR1 A1166C, CYP11B2 T344C, PON1 L55M). Our analysis demonstrated a likely genetic effect for ischaemic white matter disease, with an apparent inverse association between CYP11B2 and WMH. A trend for positive association between MTHFR and WMH severity could not be completely interrogated, given the relatively small sample size of the available studies as compared to other well-powered genetic studies on stroke.

The MTHFR gene is involved in plasma Hcy levels and may contribute to endothelial dysfunction, which is one of the suggested mechanisms behind WMH.⁷¹ While the association between the MTHFR TT-genotype and WMH fell shy of statistical significance, the totality of the data suggested a trend for association. The Mendelian randomisation approach allowed us to investigate any potential relationship between MTHFR and WMH in more depth and evaluate for potential causality. The particular strengths of this method are that confounding factors are equally distributed among genotypes, which facilitates causality to be tested in their presence, whereas measurement error bias, reverse causality and selection biases are largely

overcome.⁷² Using this approach yielded similar values for the expected and observed OR, and there was considerable overlap of their 95% CIs. Given that the two values are derived from meta-analyses of the different study types (genetic association vs observational)—either of which is prone to a different source of bias—might be suggestive of a causal association between Hcy and WMH burden.⁷³ However, this analysis was insufficiently powered, and future studies using an adequate sample size may prove more conclusive.

A number of study limitations need to be documented. Publication bias is always a concern in meta-analysis. However, funnel plots were produced for each gene-WMH association and Egger's regression analysis (two-tailed test) was performed to assess publication bias. Given that the majority of included studies reported non-significant results, substantial publication bias is considered unlikely, although it can never be completely excluded. Further, of the 78 studies investigating the association of candidate genes and WMH, just under half did not have usable data for meta-analysis. It may be that the authors of these papers did not consider the data to be interesting enough to report (selective outcome reporting). The vast majority of these studies found no association and their inclusion would have strengthened our finding of no relationship between WMH and any of the studied gene polymorphisms. Some studies reported data according to a genetic model they had chosen rather than reporting event rates for each genotype separately, which limited our ability to incorporate their studies into other genetic models. There was considerable variation in WMH measurement methods used between studies, which introduces methodological heterogeneity. Assessing WMH using visual rating scales can be subjective and observer dependent,³ although most papers reported good inter-rater agreement. The inclusion of CT and MRI studies adds another source of interstudy heterogeneity. CT has been shown to be less sensitive at detecting WMH and its use may result in an underestimation of the true WMH load within those studies. However, removal of these studies did not lead to a substantial change in the pooled OR; thus, we considered it appropriate to include them. Finally, results could be confounded by failure to adjust for age, intracranial volume and vascular risk factors in all studies. Additionally, consideration ought to be given in analyses to the known WMH risk factors since genes may be exerting their effect through these factors. The variable disease status of the study populations could have introduced heterogeneity into our analysis. For example, six studies in our APO E4 analysis were conducted in participants with probable or pathologically confirmed Alzheimer's disease.^{74 75} Combining these studies with those of asymptomatic participants could have confounded our results. Finally, a number of covariates which we are not able to assess because of a lack of complete data sets may influence our final results.

Despite undertaking, to the best of our knowledge, the largest meta-analysis to date along with studying a new independent WMH cohort, the genetics of this condition remains unclear. Future genetic studies not using an a priori hypothesis may shed further light on this field.

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Acknowledgements PS is a former Department of Health (UK) Senior Fellow. PB holds a Department of Health (UK) Senior Fellowship.

Contributors PS conceived the work. TT undertook the primary search. NR undertook the replication. All authors provided data and contributed to the final draft of the manuscript.

Funding NH is funded by a MRC (UK) PhD studentship and SY through a British Council/UK-India Education Research Initiative (UKIERI) awarded to PS.

Competing interests None.

Provenance and peer review Not commissioned; externally peer reviewed.

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REFERENCES

- O'Sullivan M. Leukoaraiosis. *Pract Neurol* 2008;8:26–38.
- De Leeuw F, De Groot J, Achten E, et al. Prevalence of cerebral white matter lesions in elderly people: a population based magnetic resonance imaging study. the rotterdam scan study. *J Neurol Neurosurg Psychiatry* 2001;70:9–14.
- Grueter BE, Schulz UG. Age-related cerebral white matter disease (leukoaraiosis): a review. *Postgrad Med J* 2012;88:79–87.
- Assareh A, Mather KA, Schofield PR, et al. The genetics of white matter lesions. *CNS Neurosci Ther* 2011;17:525–40.
- Dufouil C, de Kersaint-Gilly A, Besancon V, et al. Longitudinal study of blood pressure and white matter hyperintensities the EVA MRI cohort. *Neurology* 2001;56:921–6.
- Jeerakathil T, Wolf PA, Beiser A, et al. Stroke risk profile predicts white matter hyperintensity volume the framingham study. *Stroke* 2004;35:1857–61.
- DeBette S, Markus H. The clinical importance of white matter hyperintensities on brain magnetic resonance imaging: Systematic review and meta-analysis. *BMJ* 2010;341:c3666.
- Hendon H, Godfrey O, Lucas C, et al. Risk factors and leukoaraiosis in stroke patients. *Acta Neurol Scand* 1996;94:137–44.
- Wright CB, Paik MC, Brown TR, et al. Total homocysteine is associated with white matter hyperintensity volume the northern manhattan study. *Stroke* 2005;36:1207–11.
- Rost NS, Rahman R, Sonni S, et al. Determinants of white matter hyperintensity volume in patients with acute ischemic stroke. *J Stroke Cerebrovasc Dis* 2010;19:230–5.
- Atwood LD, Wolf PA, Heard-Costa NL, et al. Genetic variation in white matter hyperintensity volume in the framingham study. *Stroke* 2004;35:1609–13.
- Carmelli D, DeCarli C, Swan GE, et al. Evidence for genetic variance in white matter hyperintensity volume in normal elderly male twins. *Stroke* 1998;29:1177–81.
- Turner ST, Jack CR, Fornage M, et al. Heritability of leukoaraiosis in hypertensive sibships. *Hypertension* 2004;43:483–7.
- Fornage M, DeBette S, Bis JC, et al. Genome-wide association studies of cerebral white matter lesion burden. *Ann Neurol* 2011;69:928–39.
- Adib-Samii P, Rost N, Traylor M, et al. 17q25 locus is associated with white matter hyperintensity volume in ischemic stroke, but not with lacunar stroke status. *Stroke* 2013;44:1609–15.
- Paternoster L, Chen W, Sudlow CL. Genetic determinants of white matter hyperintensities on brain scans: a systematic assessment of 19 candidate gene polymorphisms in 46 studies in 19,000 subjects. *Stroke* 2009;40:2020–6.
- Casas JP, Bautista LE, Smeeth L, et al. Homocysteine and stroke: evidence on a causal link from Mendelian randomisation. *Lancet* 2005;365:224–32.
- Bland JM, Altman DG. Statistics notes: the odds ratio. *BMJ* 2000;320:1468.
- Rost NS, Rahman RM, Biffi A, et al. White matter hyperintensity volume is increased in small vessel stroke subtypes. *Neurology* 2010;75:1670–7.
- Chen YW, Gurol ME, Rosand J, et al. Progression of white matter lesions and hemorrhages in cerebral amyloid angiopathy. *Neurology* 2006;67:83–7.
- Gurol ME, Irizarry MC, Smith EE, et al. Plasma beta-amyloid and white matter lesions in AD, MCI, and cerebral amyloid angiopathy. *Neurology* 2006;66:23–9.
- Taylor WD, Zuchner S, McQuoid DR, et al. The brain-derived neurotrophic factor VAL66MET polymorphism and cerebral white matter hyperintensities in late-life depression. *Am J Geriatr Psychiatry* 2008;16:263–71.
- Hassan A, Gormley K, O'Sullivan M, et al. Endothelial nitric oxide gene haplotypes and risk of cerebral small-vessel disease. *Stroke* 2004;35:654–9.
- Henskens LH, Kroon AA, van Bostel MP, et al. Associations of the angiotensin II type 1 receptor A1166C and the endothelial NO synthase G894T gene polymorphisms with silent subcortical white matter lesions in essential hypertension. *Stroke* 2005;36:1869–73.
- Kohara K, Fujisawa M, Ando F, et al. MTHFR gene polymorphism as a risk factor for silent brain infarcts and white matter lesions in the japanese general population: the NILS-LSA study. *Stroke* 2003;34:1130–5.
- Linnebank M, Moskau S, Jurgens A, et al. Association of genetic variants of methionine metabolism with methotrexate-induced CNS white matter changes in patients with primary CNS lymphoma. *Neuro Oncol* 2009;11:2–8.
- de Lau LM, van Meurs JB, Uitterlinden AG, et al. Genetic variation in homocysteine metabolism, cognition, and white matter lesions. *Neurobiol Aging* 2010;31:2020–2.
- Pavlovic AM, Pekmezovic T, Obrenovic R, et al. Increased total homocysteine level is associated with clinical status and severity of white matter changes in symptomatic patients with subcortical small vessel disease. *Clin Neurol Neurosurg* 2011;113:711–15.
- Szolnoki Z, Somogyvari F, Kondacs A, et al. Specific APO E genotypes in combination with the ACE D/D or MTHFR 677TT mutation yield an independent genetic risk of leukoaraiosis. *Acta Neurol Scand* 2004;109:222–7.
- Szolnoki Z, Szaniszló I, Szekeres M, et al. Evaluation of the MTHFR A1298C variant in leukoaraiosis. *J Mol Neurosci* 2012;46:492–6.
- Brenner D, Labreuche J, Pico F, et al. The renin-angiotensin-aldosterone system in cerebral small vessel disease. *J Neurol* 2008;255:993–1000.
- Verpillat P, Alperovitch A, Cambien F, et al. Aldosterone synthase (CYP11B2) gene polymorphism and cerebral white matter hyperintensities. *Neurology* 2001;56:673–5.
- Amar K, MacGowan S, Wilcock G, et al. Are genetic factors important in the aetiology of leukoaraiosis? results from a memory clinic population. *Int J Geriatr Psychiatry* 1998;13:585–90.
- Bracco L, Piccini C, Moretti M, et al. Alzheimer's disease: role of size and location of white matter changes in determining cognitive deficits. *Dement Geriatr Cogn Disord* 2005;20:358–66.
- Dufouil C, Alperovitch A, Tzourio C. Influence of education on the relationship between white matter lesions and cognition. *Neurology* 2003;60:831–6.
- Dufouil C, Godin O, Chalmers J, et al. Severe cerebral white matter hyperintensities predict severe cognitive decline in patients with cerebrovascular disease history. *Stroke* 2009;40:2219–21.
- Hogervorst E, Ribeiro HM, Molyneux A, et al. Plasma homocysteine levels, cerebrovascular risk factors, and cerebral white matter changes (leukoaraiosis) in patients with alzheimer disease. *Arch Neurol* 2002;59:787–93.
- Hong ED, Taylor WD, McQuoid DR, et al. Influence of the MTHFR C677T polymorphism on magnetic resonance imaging hyperintensity volume and cognition in geriatric depression. *Am J Geriatr Psychiatry* 2009;17:847–55.
- Kalman J, Juhasz A, Csaszar A, et al. Increased apolipoprotein E4 allele frequency is associated with vascular dementia in the hungarian population. *Acta Neurol Scand* 1998;98:166–8.
- Kuller LH. Risk factors for dementia in the cardiovascular health study cognition study. *Rev Neurol* 2003;37:122–6.
- Lemmens R, Gorner A, Schrooten M, et al. Association of apolipoprotein E epsilon2 with white matter disease but not with microbleeds. *Stroke* 2007;38:1185–8.
- Bartess-Faz D, Junque C, Clemente IC, et al. MRI and genetic correlates of cognitive function in elders with memory impairment. *Neurobiol Aging* 2001;22:449–59.
- Bornebroek M, Haan J, Van Duinen SG, et al. Dutch hereditary cerebral amyloid angiopathy: structural lesions and apolipoprotein E genotype. *Ann Neurol* 1997;41:695–8.
- Doody RS, Azher SN, Haykal HA, et al. Does APO epsilon4 correlate with MRI changes in Alzheimer's disease? *J Neurol Neurosurg Psychiatry* 2000;69:668–71.
- Carmelli D, DeCarli C, Swan GE, et al. The joint effect of apolipoprotein E epsilon4 and MRI findings on lower-extremity function and decline in cognitive function. *J Gerontol A Biol Sci Med Sci* 2000;55:M103–9.
- DeCarli C, Reed T, Miller BL, et al. Impact of apolipoprotein E epsilon4 and vascular disease on brain morphology in men from the NHLBI twin study. *Stroke* 1999;30:1548–53.
- de Leeuw FE, Richard F, de Groot JC, et al. Interaction between hypertension, apoE, and cerebral white matter lesions. *Stroke* 2004;35:1057–60.
- Godin O, Tzourio C, Maillard P, et al. Apolipoprotein E genotype is related to progression of white matter lesion load. *Stroke* 2009;40:3186–90.
- Wen HM, Baum L, Cheung WS, et al. Apolipoprotein E epsilon4 allele is associated with the volume of white matter changes in patients with lacunar infarcts. *Eur J Neurol* 2006;13:1216–20.
- Vuorinen M, Solomon A, Rovio S, et al. Changes in vascular risk factors from midlife to late life and white matter lesions: a 20-year follow-up study. *Dement Geriatr Cogn Disord* 2011;31:119–25.
- Stenest V, Hofoss D, Johnsen L, et al. White matter lesion load increases the risk of low CSF Aβeta42 in apolipoprotein E-varepsilon4 carriers attending a memory clinic. *J Neuroimaging* 2011;21:e78–82.
- Smith EE, Gurol ME, Eng JA, et al. White matter lesions, cognition, and recurrent hemorrhage in lobar intracerebral hemorrhage. *Neurology* 2004;63:1606–12.

- 53 Skoog I, Hesse C, Aevansson O, *et al.* A population study of apoE genotype at the age of 85: relation to dementia, cerebrovascular disease, and mortality. *J Neurol Neurosurg Psychiatry* 1998;64:37–43.
- 54 Schmidt H, Schmidt R, Fazekas F, *et al.* Apolipoprotein E e4 allele in the normal elderly: neuropsychologic and brain MRI correlates. *Clin Genet* 1996;50:293–9.
- 55 Sawada H, Udaka F, Izumi Y, *et al.* Cerebral white matter lesions are not associated with apoE genotype but with age and female sex in alzheimer's disease. *J Neurol Neurosurg Psychiatry* 2000;68:653–6.
- 56 Nebes RD, Vora JJ, Meltzer CC, *et al.* Relationship of deep white matter hyperintensities and apolipoprotein E genotype to depressive symptoms in older adults without clinical depression. *Am J Psychiatry* 2001;158:878–84.
- 57 Benedictus MR, Goos JD, Binnewijzend MA, *et al.* Specific risk factors for microbleeds and white matter hyperintensities in Alzheimer's disease. *Neurobiol Aging* 2013;34:2488–94.
- 58 Hirono N, Yasuda M, Tanimukai S, *et al.* Effect of the apolipoprotein E epsilon4 allele on white matter hyperintensities in dementia. *Stroke* 2000;31:1263–8.
- 59 Slegers K, den Heijer T, van Dijk EJ, *et al.* ACE gene is associated with Alzheimer's disease and atrophy of hippocampus and amygdala. *Neurobiol Aging* 2005;26:1153–9.
- 60 Sierra C, Coca A, Gomez-Angelats E, *et al.* Renin-angiotensin system genetic polymorphisms and cerebral white matter lesions in essential hypertension. *Hypertension* 2002;39(2 Pt 2):343–7.
- 61 Gormley K, Bevan S, Markus HS. Polymorphisms in genes of the renin-angiotensin system and cerebral small vessel disease. *Cerebrovasc Dis* 2007;23:148–55.
- 62 Hassan A, Lansbury A, Catto AJ, *et al.* Angiotensin converting enzyme insertion/deletion genotype is associated with leukoaraiosis in lacunar syndromes. *J Neurol Neurosurg Psychiatry* 2002;72:343–6.
- 63 Mizuno T, Makino M, Fujiwara Y, *et al.* Renin-angiotensin system gene polymorphism in Japanese stroke patients. *International Congress Series* 2003;1252:83–90.
- 64 Schmidt R, Schmidt H, Fazekas F, *et al.* Angiotensinogen polymorphism M235T, carotid atherosclerosis, and small-vessel disease-related cerebral abnormalities. *Hypertension* 2001;38:110–15.
- 65 Hadjigeorgiou GM, Malizos K, Dardiotis E, *et al.* Paraoxonase 1 gene polymorphisms in patients with osteonecrosis of the femoral head with and without cerebral white matter lesions. *J Orthop Res* 2007;25:1087–93.
- 66 Schmidt R, Schmidt H, Fazekas F, *et al.* MRI cerebral white matter lesions and paraoxonase PON1 polymorphisms: three-year follow-up of the austrian stroke prevention study. *Arterioscler Thromb Vasc Biol* 2000;20:1811–16.
- 67 Albert MA, Glynn RJ, Buring JE, *et al.* Relation between soluble intercellular adhesion molecule-1, homocysteine, and fibrinogen levels and race/ethnicity in women without cardiovascular disease. *Am J Cardiol* 2007;99:1246–51.
- 68 Anand SS, Yusuf S, Vuksan V, *et al.* Differences in risk factors, atherosclerosis, and cardiovascular disease between ethnic groups in Canada: the study of health assessment and risk in ethnic groups (SHARE). *Lancet* 2000;356:279–84.
- 69 Carmel R, Green R, Jacobsen DW, *et al.* Serum cobalamin, homocysteine, and methylmalonic acid concentrations in a multiethnic elderly population: ethnic and sex differences in cobalamin and metabolite abnormalities. *Am J Clin Nutr* 1999;70:904–10.
- 70 Senaratne MP, Macdonald K, De Silva D. Possible ethnic differences in plasma homocysteine levels associated with coronary artery disease between south asian and east asian immigrants. *Clin Cardiol* 2001;24:730–4.
- 71 Szolnoki Z. Pathomechanism of leukoaraiosis. *Neuromolecular Med* 2007;9:21–33.
- 72 Smith GD, Ebrahim S. What can mendelian randomisation tell us about modifiable behavioural and environmental exposures? *BMJ* 2005;330:1076.
- 73 Wald DS, Law M, Morris JK. Homocysteine and cardiovascular disease: Evidence on causality from a meta-analysis. *BMJ* 2002;325:1202.
- 74 Poirier J, Bertrand P, Kogan S, *et al.* Apolipoprotein E polymorphism and Alzheimer's disease. *The Lancet* 1993;342:697–9.
- 75 Rubinsztein DC, Easton DF. Apolipoprotein E genetic variation and Alzheimer's disease. *Dement Geriatr Cogn Disord* 1999;10:199–209.