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Cerebrovascular disease

RESEARCH PAPER

Cognitive impairment after lacunar stroke: systematic review and meta-analysis of incidence, prevalence and comparison with other stroke subtypes

Stephen David James Makin,1 Sarah Turpin,2 Martin S Dennis,1 Joanna M Wardlaw3

ABSTRACT

Background  Cognitive impairment and dementia are common after stroke. It is unclear if risk differs between ischaemic stroke subtypes. Lacunar strokes might be less likely to affect cognition than more severe, larger cortical strokes, except that lacunar strokes are associated with cerebral small vessel disease (SVD), which is the commonest vascular cause of dementia.

Methods  We searched MEDLINE and PsychINFO for studies of mild cognitive impairment (MCI) or dementia after lacunar or cortical ischaemic stroke. We calculated the OR for cognitive impairment/dementia in lacunar versus non-lacunar stroke, and their incidence and prevalence in lacunar stroke as a pooled proportion.

Findings  We identified 24 relevant studies of 7575 patients, including 2860 with lacunar stroke; 24% had MCI or dementia post stroke. Similar proportions of patients with lacunar and non-lacunar stroke (16 studies, n=6478) had MCI or dementia up to 4 years after stroke (OR 0.72 (95% CI 0.43 to 1.20)). The prevalence of dementia after lacunar stroke (six studies, n=1421) was 20% (95% CI 9 to 33) and the incidence of MCI or dementia (four studies, n=275) was 37% (95% CI 23 to 53). Data were limited by short follow-up, subtype classification methods and confounding.

Interpretation  Cognitive impairment appears to be common after lacunar strokes despite their small size, suggesting that associated SVD may increase their impact. New prospective studies are required with accurate stroke subtyping to assess long term outcomes while accounting for confounders.

INTRODUCTION

Dementia is common soon after stroke1 but we know little of the mechanisms or whether the risk varies with stroke subtype. Stroke risk factors, amounts and regions of the brain affected, and suggested stroke mechanisms all vary with stroke subtype, and could influence the risk of cognitive impairment after stroke. For example, as cortical ischaemic strokes often affect a large area of brain, they may carry a higher risk of cognitive impairment than the smaller, less neurologically severe, lacunar strokes. Alternatively, lacunar strokes may carry a higher risk of cognitive impairment than would be expected on the basis of the lacunar infarct alone as they are part of the spectrum of cerebral small vessel disease (SVD). SVD, which affects the brain diffusely and is the commonest vascular cause of cognitive impairment,2 could be unmasked by a new lacunar stroke. A recent systematic review3 included studies of patients after lacunar stroke who had undergone detailed neuropsychological assessments but did not provide information on whether lacunar stroke patients are at greater risk than other stroke subtypes, or on potential confounding factors.

We sought to establish the incidence and prevalence of cognitive impairment and dementia after lacunar stroke, in the short and long term, and its magnitude in comparison with cortical stroke, by systematically reviewing the available literature. We also aimed to evaluate study quality and whether they had accounted for confounding factors that may affect performance on cognitive tests such as depression, lower premorbid IQ and pre-existing dementia or SVD.

METHODS

Search methods

Following guidance from experienced librarians from the Cochrane Stroke Group, we searched MEDLINE (1991 to present) and PsychINFO (1991 to present) with the terms (a) ‘cognition’ ‘dementia’ ‘vascular dementia’ or ‘multi-infarct dementia’ and (b) ‘stroke’ or ‘lacunar stroke’ (last searched 7 June 2012, using OVID SP V03.07.00.119, details in the online supplementary information). We checked references in review articles and hand searched the previous 5 years of Stroke and Age and Aging. We complied with the meta-analysis of observational systematic reviews (MOOSE)4 group guidelines.

Study criteria

In our primary analysis we included studies that compared cognition in patients with lacunar stroke with those with cortical stroke. Our secondary analysis to determine the incidence and prevalence of mild cognitive impairment (MCI) and dementia analysed all studies that measured cognition in patients with lacunar stroke.

We included studies that assessed MCI and dementia in adult humans with symptomatic lacunar stroke (defined below). We excluded studies of asymptomatic patients with lesions on imaging and autopsy studies.

In order to establish the proportion of lacunar stroke patients with MCI or dementia, we included studies that tested an unselected group of patients. We excluded studies that tested a very selected group of patients—for example, a study of only patients with a particular radiological finding such as a recent lacunar infarct in the thalamus.

We included English and non-English language studies. We excluded studies that were published only in abstract and those which had tested a group of stroke patients, including some with lacunar stroke, but not presented the results according to stroke subtype.

**Definitions of lacunar stroke, dementia and MCI**

We set the reference standard for stroke subtyping as a non-risk factor dependent clinically based approach (such as the Oxfordshire Community Stroke Project, OSCP) accompanied by MRI, including diffusion weighted imaging (DWI), to identify the recent acute ischaemic lesion. We also accepted studies which made the diagnosis of stroke subtype in other ways, such as clinical syndrome plus CT scanning instead of MR with DWI. We defined a lacunar stroke as one diagnosed as ‘lacunar’ on clinical grounds (a typical lacunar syndrome, or using methods devised by Bamford et al in the OCSP or similar method) with or without imaging verification, or as a clinically evident stroke classed as ‘small vessel’ by the risk factor based classifications such as the Trial of ORG 10172 in Acute Stroke Treatment (TOAST) classification. We defined non-lacunar stroke as a sudden onset of neurological symptoms classed as ‘cortical stroke’ clinically with or without imaging verification, or a stroke classed as ‘large artery atherosclerosis’, ‘cardioembolism’, ‘cryptogenic’ or ‘other’ on TOAST, or other risk factor based classifications. We noted the definition used in each study so as to perform sensitivity analyses.

We set the reference standard for cognitive impairment as a full neuropsychological assessment covering all cognitive domains, blinded to stroke subtype. We defined dementia as impairment of cognitive function which interfered with everyday activities, and we defined MCI as impairment of cognitive function not severe enough to interfere with everyday life. We used the term ‘cognitive impairment’ to refer to any impairment of cognitive function whether MCI or dementia.

We included studies which had made the diagnosis of cognitive impairment or dementia in other ways, such as the Mini-Mental State Examination (MMSE). We used the criteria for cognitive impairment, MCI or dementia as applied in individual studies.

**Data extraction**

We used a pretested form to extract available data. We extracted the following data about patient recruitment and assessment methods: inclusion and exclusion criteria (specifically whether the study included patients with prior stroke, prior cognitive impairment or aphasia, and if so whether adapted tests were used); how stroke subtypes were defined and if there were any differences between the subtypes; whether information was sought from relatives about signs of cognitive decline prior to stroke; and the details of assessments, including whether depression or premorbid IQ were accounted for; whether patients had MRI or CT imaging to aid diagnosis of subtype; and the proportion of patients who were cognitively assessed. Secondly, we aimed to calculate the incidence and prevalence of cognitive impairment and dementia after lacunar stroke. We calculated the OR and 95% CI using a DerSimonian Laird random effects model in StatsDirect V.2.7.8. We assessed for heterogeneity by calculating the I² statistic for each meta-analysis and publication bias using a funnel plot. We used a random effects model due to potential heterogeneity in the underlying methodology. No ethics approval was necessary in order to conduct this literature review as it was an literature based.

**RESULTS**

Of 320 potentially suitable papers, 164 assessed cognition in stroke patients but only 57 assessed both cognitive function and stroke subtype; of these, 33 presented the results by subtype but nine did not meet other prespecified criteria (see details in the online supplementary table), leaving 24 that met our inclusion criteria (figure 1). The 24 studies included 7575 subjects with ischaemic stroke (table 1).

Nineteen studies compared lacunar with non-lacunar stroke and/or dementia and/or MCI; if impairments occurred in any particular cognitive domain; and any calculated odds or HRs and whether these were adjusted for risk factors, including SVD on imaging, depression and premorbid IQ.

If a study had published more than one paper, we ensured that each patient only contributed once towards the present analysis. The studies were reviewed and the data extracted by two of the authors (SDJM and ST) with advice from another author (JMW). We did not contact the authors to obtain additional information.

**Statistical analysis**

Firstly we investigated whether patients with lacunar stroke were more likely to have cognitive impairment than patients with non-lacunar stroke by calculating the pooled OR and 95% CI of cognitive impairment in patients with lacunar compared with cortical stroke (the OR would be >1.0 if cognitive impairment were more common in lacunar stroke).

We examined the impact of potential confounders by recording available adjusted outcome measures and by analysis of the following prespecified subgroups: community versus hospital based studies; timing of cognitive assessment after stroke (early=under 1 month, mid=1 month to 1 year and later than 1 year); whether or not patients with previous stroke and/or dementia had been excluded; stroke subtyping method used; cognitive test used; whether depression or premorbid IQ were accounted for; whether patients had MRI or CT imaging to aid diagnosis of subtype; and the proportion of patients who were cognitively assessed. Secondly, we aimed to calculate the incidence and prevalence of cognitive impairment and dementia in patients with lacunar stroke as a pooled proportion and 95% CI, using a DerSimonian Laird random effects model in StatsDirect V.2.7.8. We assessed for heterogeneity by calculating the I² statistic for each meta-analysis and publication bias using a funnel plot. We used a random effects model due to potential heterogeneity in the underlying methodology. No ethics approval was necessary in order to conduct this literature review as it was an literature based.
the index stroke (38% of studies, 19% of patients). Three studies\(^9\)\(^{24}\)\(^{25}\) stated that the assessors were blinded to the stroke subtype (13% of studies, 7% of patients) but the rest did not mention blinding. Some information on background SVD was obtained in 10 studies but the varied methods and reporting precluded its further use: one study (n=81) measured white matter lesion (WML) volume on MRI in all patients;\(^31\) another (n=170) measured WML volume in 133/170 patients who had MRI\(^10\); and eight studies\(^11\)\(^{12}\)\(^{18}\)\(^{21}\)\(^{24}\)\(^{26}\)\(^{30}\)\(^{33}\) scored leukoaraiosis on CT scans of all patients (33% of studies, 16% of patients).

Stroke subtyping methods and use of imaging in subtype diagnosis varied. All studies required a clinical diagnosis of a stroke alongside imaging to exclude haemorrhage. Two studies\(^23\)\(^{31}\) performed MRI on all patients but did not state whether or not this included DWI (8% of studies, 5% of patients), a further 11\(^11\)\(^{13}\)\(^{18}\)\(^{24}\)\(^{26}\)\(^{30}\)\(^{33}\)\(^{34}\)\(^{36}\) performed CT scanning on all patients (46% of studies, 28% of subjects). The remaining studies either performed a mixture of CT and MRI scans or did not give imaging details. Nine studies\(^12\)\(^{16}\)\(^{26}\)\(^{29}\)\(^{31}\)\(^{32}\)\(^{33}\)\(^{35}\)\(^{36}\) used a risk factor free clinical classification (eg, OCSP\(^7\)) to categorise subjects as ‘lacunar’ or ‘non-lacunar’ stroke (38% of studies, 23% of patients). Nine studies\(^10\)\(^{13}\)\(^{15}\)\(^{18}\)\(^{23}\)\(^{27}\)\(^{34}\)\(^{36}\) used the risk factor based TOAST classification (38% of studies, 66% of patients). Six studies\(^9\)\(^{11}\)\(^{20}\)\(^{24}\)\(^{25}\)\(^{30}\) used imaging features alone to subtype the clinically apparent stroke (25% of studies, 11% of patients).

The extent of cognitive testing varied. Sixteen studies\(^9\)\(^{12}\)\(^{14}\)\(^{15}\)\(^{18}\)\(^{21}\)\(^{23}\)\(^{25}\)\(^{28}\)\(^{32}\)\(^{34}\)\(^{35}\) performed detailed cognitive testing of all cognitive domains (67% of studies, 72% of patients). Six studies\(^16\)\(^{24}\)\(^{29}\)\(^{31}\)\(^{33}\)\(^{36}\) used only the MMSE (25% of studies, 18% of patients). One\(^11\) (n=58) used the Rotterdam Cambridge Cognitive Assessment (R-CAMCOG) while another\(^16\) (n=75) used the Alzheimer’s Disease Assessment Scale cognitive subscale (ADAS-COG) and MMSE. One study\(^13\) (n=726) based cognitive assessment on the assessing clinician’s impression of whether the patient was demented. Two studies\(^10\)\(^{20}\) recorded the number of subjects with MCI but no dementia. One study\(^15\) (n=336) reported test results across OCSP subtypes: lacunar patients performed worse on the ‘matrix reasoning’ test of visuoperceptual functioning and anterior (not posterior) circulation cortical stroke patients performed worse on the Stroop test of executive function; neither difference was seen on other tests of the same function.

The timing of the cognitive assessment varied. Four studies\(^9\)\(^{14}\)\(^{29}\)\(^{34}\) assessed cognition in the first month after stroke (17% of studies, 42% of patients). Twelve studies\(^10\)\(^{12}\)\(^{15}\)\(^{16}\)\(^{23}\)–\(^{28}\)\(^{30}\) assessed cognition between 3 months and 1 year (50% studies, 34% patients), and eight\(^13\)\(^{18}\)\(^{20}\)\(^{31}\)–\(^{33}\)\(^{35}\)\(^{36}\) assessed cognition between 1 and 4 years after stroke (33% studies, 24% patients).

We compared the risk of cognitive impairment in patients with lacunar stroke with those with non-lacunar stroke (figure 2). As we were not able to analyse studies that only presented mean cognitive test scores nor those without a control group, only 16/24 studies\(^9\)–\(^{16}\)\(^{18}\)\(^{22}\)–\(^{28}\) of 6478 subjects contributed to this analysis. The proportion of patients with cognitive impairment—either MCI or dementia—did not differ between patients with lacunar and non-lacunar stroke: 638/2222 (29%) subjects with lacunar stroke compared with 1001/4256 (24%) subjects with non-lacunar stroke (OR 0.72; 95% CI 0.43 to 1.20). The substantial heterogeneity between the studies (I\(^2\) 91%; p<0.001) was not apparently due to publication bias (figure 3) or to differences in the time elapsed between the index stroke and assessment of cognition (figure 2).

We performed a sensitivity analyses (figure 4) by sequentially excluding studies with particular characteristics and then repeating the analysis. Among hospital based studies, cognitive impairment was more common in non-lacunar stroke (OR 0.67; 95% CI 0.51 to 0.86); in community based studies, cognitive impairment was more common in lacunar stroke (OR 1.56; 95% CI 0.32 to 7.64) but with considerable heterogeneity (I\(^2\) 98%; p<0.001). There was little difference in the summary OR between studies that only included first stroke

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**Figure 1** Flow diagram of search strategy and results.

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### Table 1  Characteristics of included studies

<table>
<thead>
<tr>
<th>Reference</th>
<th>Setting</th>
<th>Total patients consented/total tested</th>
<th>Timing of tests post-stroke</th>
<th>Test done</th>
<th>Subtyping</th>
<th>Primary outcome</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nys 2007</td>
<td>Hospital The Netherlands</td>
<td>190/168*</td>
<td>3 weeks</td>
<td>Imaging</td>
<td>MCI†</td>
<td>29/64 (45%); Lacunar 47/63 (75%); Non-lacunar 74/120 (62%)</td>
<td></td>
</tr>
<tr>
<td>Sachdev et al</td>
<td>Hospital Australia</td>
<td>210/170*</td>
<td>3–6 months</td>
<td>NP</td>
<td>Risk factor</td>
<td>Dementia or MCI</td>
<td>24/46 (52%); Lacunar 4/20 (20%); Non-lacunar 17/38 (45%); Non-lacunar 13/83 (16%)</td>
</tr>
<tr>
<td>De Koning et al</td>
<td>Hospital The Netherlands</td>
<td>130/121*</td>
<td>3–9 months</td>
<td>NP</td>
<td>Imaging</td>
<td>Dementia</td>
<td>29/64 (45%); Lacunar 47/63 (75%); Non-lacunar 74/120 (62%)</td>
</tr>
<tr>
<td>Censori et al</td>
<td>Hospital Italy</td>
<td>121/110**</td>
<td>3–4 Months</td>
<td>NP and MMSE††</td>
<td>Clinical</td>
<td>Dementia</td>
<td>Lacunar 24/46 (52%); Non-lacunar 74/120 (62%)</td>
</tr>
<tr>
<td>Tatemichi et al</td>
<td>Hospital USA</td>
<td>927/726</td>
<td>2 years</td>
<td>Clinical impression</td>
<td>Risk factor</td>
<td>Dementia</td>
<td>Lacunar 25/227 (11%); Non-lacunar 91/499 (18%)</td>
</tr>
<tr>
<td>Bejot et al</td>
<td>Community France</td>
<td>3948/3201*</td>
<td>1 month</td>
<td>NP</td>
<td>Risk factor</td>
<td>Dementia</td>
<td>Lacunar 333/887 (38%); Non-lacunar 289/1960 (15%)</td>
</tr>
<tr>
<td>Lin et al</td>
<td>Hospital Taiwan</td>
<td>353/283</td>
<td>3 months</td>
<td>NP</td>
<td>Risk factor</td>
<td>Dementia</td>
<td>Lacunar 13/136 (10%); Non-lacunar 13/147 (9%)</td>
</tr>
<tr>
<td>Patel et al</td>
<td>Community UK</td>
<td>1454/654*</td>
<td>3 months</td>
<td>MMSE</td>
<td>Clinical</td>
<td>MMSE&lt;24</td>
<td>Lacunar 72/218 (33%); Non-lacunar 124/297 (42%)</td>
</tr>
<tr>
<td>Cordoliani-Mackowiak et al</td>
<td>Hospital France</td>
<td>132/88‡‡</td>
<td>Up to 3 years</td>
<td>NP</td>
<td>Risk factor</td>
<td>Dementia</td>
<td>Lacunar 8/31 (26%); Non-lacunar 24/101 (24%); Non-lacunar 40/57 (70%); Non-lacunar 59/87 (68%); Non-lacunar</td>
</tr>
<tr>
<td>Rasquin et al</td>
<td>Hospital The Netherlands</td>
<td>176/144</td>
<td>1 years</td>
<td>NP</td>
<td>Imaging</td>
<td>MCI or Dementia</td>
<td>Lacunar 4/20 (20%); Non-lacunar 17/38 (45%); Non-lacunar 13/83 (16%)</td>
</tr>
<tr>
<td>Pohjasaava et al</td>
<td>Hospital Finland</td>
<td>451/337</td>
<td>3 months</td>
<td>NP</td>
<td>Risk factor</td>
<td>Dementia</td>
<td>Lacunar 5/21 (24%); Non-lacunar 13/147 (9%)</td>
</tr>
<tr>
<td>Tang et al</td>
<td>Hospital China</td>
<td>484/280*</td>
<td>3 months</td>
<td>MMSE</td>
<td>Imaging</td>
<td>Dementia</td>
<td>Lacunar 33/166 (20%); Non-lacunar 18/90 (20%); Non-lacunar 5/41 (12%); Non-lacunar 541 (12%); Non-lacunar</td>
</tr>
<tr>
<td>Madureira et al</td>
<td>Hospital Portugal</td>
<td>180/165</td>
<td>3 months</td>
<td>NP</td>
<td>Imaging</td>
<td>Dementia</td>
<td>Lacunar 6/139 (4.3%); Non-lacunar 5/41 (12%); Non-lacunar 541 (12%); Non-lacunar</td>
</tr>
<tr>
<td>Klimkowicz-Mrowiec et al</td>
<td>Hospital Poland</td>
<td>173/145‡‡</td>
<td>3 months</td>
<td>NP and MMSE</td>
<td>Clinical</td>
<td>Dementia</td>
<td>Lacunar 2/24 (8%); Non-lacunar 35/149 (23%); Non-lacunar 21/106 (20%); Non-lacunar 31/97 (33%) Non-lacunar</td>
</tr>
<tr>
<td>Dong et al</td>
<td>Hospital Singapore</td>
<td>300/239*</td>
<td>3–6 months</td>
<td>NP, MMSE, and MoCA§§</td>
<td>Risk factor</td>
<td>Moderate MCI</td>
<td>Lacunar 21/59 (36%); Non-lacunar 59/168 (35%); Non-lacunar</td>
</tr>
<tr>
<td>Tatemichi et al</td>
<td>Hospital USA</td>
<td>Not stated/227</td>
<td>3 months</td>
<td>NP</td>
<td>Clinical</td>
<td>MCI</td>
<td>Lacunar 5/21 (24%); Non-lacunar 13/147 (9%)</td>
</tr>
<tr>
<td>Tay et al</td>
<td>Hospital Singapore</td>
<td>216/169</td>
<td>9 days</td>
<td>MMSE</td>
<td>Clinical</td>
<td>NA</td>
<td>Lacunar 21/59 (36%); Non-lacunar 59/168 (35%); Non-lacunar</td>
</tr>
<tr>
<td>Mok et al</td>
<td>Hospital China</td>
<td>86/75</td>
<td>3 months***</td>
<td>MMSE and clinical dementia rating scale</td>
<td>Imaging</td>
<td>Dementia</td>
<td>Lacunar 10/75 (13%); Mean MMSE: 24.8 lacunar, 27.7 healthy controls</td>
</tr>
<tr>
<td>Samuelsson et al</td>
<td>Hospital Sweden</td>
<td>100/81</td>
<td>2 years</td>
<td>MMSE and NP if impaired.</td>
<td>Clinical</td>
<td>Dementia</td>
<td>Lacunar 2/24 (8%), Non-lacunar 21/106 (20%)</td>
</tr>
<tr>
<td>Anderson et al</td>
<td>Community Australia</td>
<td>Not stated/30</td>
<td>1 year</td>
<td>NP</td>
<td>Clinical</td>
<td>MCI</td>
<td>Lacunar 2/24 (8%); Non-lacunar 21/106 (20%)</td>
</tr>
<tr>
<td>Loeb et al</td>
<td>Hospital Italy</td>
<td>Not stated/108</td>
<td>Up to 4 years</td>
<td>NP</td>
<td>Clinical</td>
<td>MCI</td>
<td>Lacunar 25/108 (23%); Non-lacunar 41/71 (58%)</td>
</tr>
<tr>
<td>Fure et al</td>
<td>Hospital Norway</td>
<td>71/64‡‡</td>
<td>2–7 days</td>
<td>MMSE</td>
<td>Clinical</td>
<td>MCI</td>
<td>Lacunar 25/108 (23%); Non-lacunar 41/71 (58%)</td>
</tr>
<tr>
<td>Barker-Collo et al</td>
<td>Community New Zealand</td>
<td>357/336</td>
<td>5 years</td>
<td>NP</td>
<td>Clinical</td>
<td>Cognition test results.</td>
<td>Mean MMSE scores Lacunar 25.6, Lacunar 26.6</td>
</tr>
<tr>
<td>Appelros et al</td>
<td>Community Sweden</td>
<td>253/232*</td>
<td>1 year</td>
<td>MMSE</td>
<td>Risk factor</td>
<td>Lacunar 25/108 (23%); Non-lacunar 41/71 (58%)</td>
<td></td>
</tr>
</tbody>
</table>

*Includes subjects with haemorrhagic stroke/transient ischaemic attack.
††Full neuropsychological (NP) testing.
‡‡Mild cognitive impairment.
§§Subtype reported only in patients who had a visible lesion on CT.
**Six subjects with specific cognitive impairment excluded.
†††Mini-Mental State Examination.
‡‡‡Data reported on subjects who could not have full tests.
§§§Calculated from reported OR.
¶¶¶Calculated from χ² statistic.
***The 61 non-demented patients were followed up 2–3 years later; no information on outcome of those who did have dementia at 3 months.
§§§MoCA, Montreal Cognitive Assessment.
(OR 0.88; 95% CI 0.3 to 2.58), first or recurrent stroke (OR 0.69; 95% CI 0.54 to 0.9), where subjects with prior dementia were excluded (OR 0.63; 95% CI 0.43 to 0.91) or included (OR 0.88; 95% CI 0.35 to 2.53), between studies that tested smaller or larger numbers of participants (OR 0.63; 95% CI 0.46 to 0.86), for studies testing <80% (OR 0.63; 95% CI 0.46 to 0.86) against studies testing >80% of patients (OR 0.72; 95% CI 0.33 to 1.57), or according to patient age, whether limited or full neuropsychological testing was performed, or by the proportion of patients who had MRI scanning. Additionally, there was little difference in the summary OR between studies that used clinical, imaging or risk factor based subtyping to diagnosis lacunar stroke (see online supplementary figure). We did not have enough information to

**Figure 2** Odds of cognitive impairment in lacunar stroke versus cortical stroke.

**Figure 3** Funnel plot demonstrating that the heterogeneity was not due to publication bias. This figure is only reproduced in colour in the online version.
assess the impact of SVD on imaging. No studies provided a multivariate analysis of the risk of cognition in lacunar stroke adjusted for imaging evidence of SVD, depression, age or other confounders.

Heterogeneity was due to one large study14 (n=2847) in which cognitive impairment was more common in lacunar stroke (OR 3.48, 95% CI 2.89 to 4.18). When the meta-analysis was repeated after omitting this study, the OR changed from 0.72 (95% CI 0.43 to 1.20) with significant heterogeneity (as above) to 0.67 (95% CI 0.54 to 0.84) without significant heterogeneity (I² 25%, p=0.17).

For the second meta-analysis, we aimed to establish the incidence and prevalence of cognitive impairment and dementia after a lacunar stroke. We included data from all studies that had measured cognition in patients with a lacunar stroke (a total of 21 studies of 2606 lacunar strokes) whether or not there was a control group. We were not able to include those studies that reported only average cognitive test results and did not dichotomise patients into ‘impaired’ and ‘not impaired’. A pooled proportion of eight9 10 20 27 28 32–34 studies of 541 patients found that 38% (95% CI 24 to 52, I² 92%) had MCI or dementia at any stage after a first or recurrent lacunar stroke. Fifteen studies provided data on dementia at any time after lacunar stroke (16%, 95% CI 10 to 24, I² 93.8%)10–16 18 22–26 30 31.

We then calculated the incidence (table 2) of dementia or MCI from studies that excluded patients with these conditions prior to stroke. One small study12 (n=21) found that the incidence of dementia was 10% (2/21; 95% CI 2 to 30) after a first lacunar stroke and another study10 (n=46) found that the incidence of MCI was 16% (95% CI 10 to 24). Six studies10 12 15 18 25 26 (n=397) found that the incidence of dementia after a first or recurrent lacunar stroke was 12% (95% CI 6 to 18). No studies of first and recurrent lacunar stroke recorded the incidence of MCI alone.

We planned to calculate the prevalence of dementia or MCI from studies that had included patients who had these conditions prior to the index stroke but no studies did this for MCI. Six studies provided these data for patients with dementia, two14 31 (n=987) for first stroke (pooled percentage 21%; 95% CI 10 to 55) and six11 13 14 23 24 31 (n=1421) for first or recurrent lacunar stroke (pooled percentage 20%; 95% CI 9 to 33).

**DISCUSSION**

The available literature suggests that approximately 30% of patients will be cognitively impaired in the 4 years following a lacunar stroke, a similar proportion to non-lacunar stroke (23%). However, precision and reliability of the estimate were
There is a lack of information on long term prognosis yet the chances of continuing cognitive decline after lacunar stroke may be higher than in non-lacunar stroke due to the differences in cognitive impairment and fatigable lacunar stroke commonly affects patients in their 50s and 60s with many years of life and employment ahead. One study of 10833 patients performed a year after the index stroke, yet lacunar stroke commonly affects patients who had dementia prior to the stroke. Importantly, there were few data on cognition beyond subtyping supported by MRI that when impairment does occur, it is generalised rather than domain specific.

The limitations of the literature included the fact that many studies were small, few assessors were blinded to the stroke subtype, most studies did not account for important confounders, and the diagnostic methods and reporting varied considerably. Importantly, there were few data on cognition beyond 1 year after the index stroke. lacunar stroke is related to cerebral SVD which is hampered by several factors. Thus despite SVD being the commonest vascular cause of cognitive impairment and lacunar stroke often affecting people in middle not just old age, data to calculate the true incidence or prevalence of cognitive impairment after lacunar stroke and associated risk factors in order to advise patients and plan services are lacking.

A recent review summarised domain specific cognitive impairment after lacunar stroke but did not provide information on the proportion of lacunar stroke patients who are cognitively impaired, nor on whether this differs from non-lacunar stroke, or on the outcome of patients who were not able to have detailed cognitive assessment, nor the effect of depression, premorbid IQ or imaging evidence of SVD. However, it did highlight that when impairment does occur, it is generalised rather than domain specific.

The estimate of cognitive impairment after lacunar versus non-lacunar stroke was strongly influenced by one study in which dementia was much more common in lacunar than in non-lacunar stroke (OR 3.48 vs OR 0.67 in all other studies combined). We should not discount the findings of this large population based study that contained 38% of all patients. However, we noted two problems: cognition was assessed less than 1 month after stroke and the OR of cognitive impairment in lacunar stroke changed over the study period, from 10.1 in 1991–1996 to 1.51 in 2003–2008, implying a change in methodology over time.

Our study has limitations. We lacked the resources to contact the authors to obtain original patient data on risk factors or missing values. However, it appeared that most of the missing information had not been collected.

This study’s strengths include a comprehensive literature review amassing data relating to 2860 lacunar stroke patients and 4275 non-lacunar controls, including studies published in Asia, Europe and America. We did not exclude any studies because we could not obtain the papers or translate them into English.

No studies

Prevalence of MCI or dementia (including studies where the methods were unclear)

<table>
<thead>
<tr>
<th>No of patients</th>
<th>Pooled risk or proportion (95% CI) (%)</th>
<th>12 (95% CI) (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1st stroke</td>
<td>1262</td>
<td>20 (8 to 36)</td>
</tr>
<tr>
<td>1st or recurrent stroke</td>
<td>1777</td>
<td>18 (9.6 to 30)</td>
</tr>
<tr>
<td>1st or recurrent stroke</td>
<td>464</td>
<td>45 (34 to 58)</td>
</tr>
<tr>
<td>1st or recurrent stroke</td>
<td>275</td>
<td>37 (23 to 53)</td>
</tr>
<tr>
<td>1st or recurrent stroke</td>
<td>275</td>
<td>38 (13 to 66)</td>
</tr>
<tr>
<td>1st or recurrent stroke</td>
<td>668</td>
<td>96 (93.3 to 97.3)</td>
</tr>
</tbody>
</table>

MCI, mild cognitive impairment.
underlying pathology: SVD affects the brain diffusely whereas non-lacunar stroke often has an extracranial cause. The prognosis is important to patients who are cognitively impaired after lacunar stroke who wish to know if they will recover, to plan stroke services and to enable the planning of sample size and length of follow-up in future interventional studies.

Future research should focus on previously cognitively intact patients with a first lacunar stroke, extend follow-up for as long as practicable, subtype the stroke on clinical grounds aided by early MRI–DWI, account for confounding factors including differences in vascular risk factors, premorbid IQ, pre-stroke cognitive decline and depression, and validated imaging biomarkers of SVD. Analysis should calculate the odds of dementia free survival, which would reduce survivorship bias.

Contributors SDJM drew up the protocol, carried out the search, extracted the data, performed the meta-analysis and drafted the manuscript. ST performed data extraction, checking and editing. JMW discussed the protocol, reviewed papers that were uncertain, advised on the meta-analysis, interpreted the data, and reviewed the manuscript. All authors approved the manuscript before submission.

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Competing interests None.

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