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Temporal Lobe Atrophy and White Matter Lesions are Related to Major Depression Over 5 Years in the Elderly

Running title: Brain Changes Related to Major Depression in the Elderly

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ABSTRACT

The influence of organic brain changes on the development of depression in the elderly is uncertain. Cross-sectional studies, most often from clinical samples, report associations with brain atrophy and cerebrovascular disease, while longitudinal population studies have given mixed results. Our aim was to investigate whether cortical atrophy and white matter lesions on computed tomography (CT) predict occurrence of depression in the elderly. This is a prospective population-based study with 5 years follow-up. The baseline sample included 525 elderly, age 70–86 years, without dementia or major depression and with a score on the Mini-Mental State Examination above 25, and without dementia at follow-up. Cortical atrophy and white matter lesions were evaluated at baseline using CT. The main outcome measure was development of major or minor depression at follow-up according to *DSM-IV*, as evaluated using neuropsychiatric examinations and hospital discharge registers. Logistic regression was used to estimate risk. Over 5 years, 20 individuals developed major and 63 minor depression. Presence of temporal lobe atrophy (odds ratio [OR] = 2.81, 95% confidence interval [CI] 1.04–7.62) and moderate-severe white matter lesions (OR = 3.21, 95% CI 1.00–10.26) independently predicted major, but not minor, depression after controlling for various confounders. Other brain changes did not predict occurrence of depression. Our findings suggest that temporal lobe atrophy and white matter lesions represent relatively independent and complementary pathways to major depression in the elderly. This may have implications for prevention, as both neurodegeneration and cerebrovascular disease has been related to preventable factors.

Keywords: depression, elderly, brain atrophy, white matter lesions, population, longitudinal

INTRODUCTION

Depression is one of the most common mental disorders in the elderly and has a significant impact on quality of life, mortality, and cognitive function (Palsson and Skoog, 1997). It is therefore important to learn more about its risk factors. Reported risk factors include female gender, adverse life events, other psychosocial factors, and somatic disorders, such as stroke and cardiovascular disorders (Djernes, 2006; Palsson and Skoog, 1997). In addition, it has been suggested that organic brain changes, such as neurodegeneration and cerebrovascular disease, may be important in the etiology of geriatric depression.

Most studies on the association between brain atrophy and depression in the elderly have been cross-sectional and performed in clinical samples. The most consistent findings from cross-sectional studies are that depression in the elderly is associated with atrophy in the frontal and temporal lobes, especially hippocampal atrophy and ventricular enlargement (Schweitzer *et al*, 2001). Only one longitudinal population-based study examined the association between brain atrophy and occurrence of depressive symptoms or syndromes during follow-up (Ikram *et al*, 2010). This study reported a cross-sectional association between temporal and parietal lobe atrophy at baseline and concurrent depressive symptoms, and no association with depression occurring during follow-up. Another longitudinal study in elderly without dementia examined the association between mean depressive symptom scores over time and change in brain atrophy on magnetic resonance imaging (MRI) over nine years (Dotson *et al*, 2009). Cross-sectional associations between temporal and frontal lobe atrophy and depressive symptoms, and a longitudinal association between volume decline in frontal white matter and depressive symptom score over time were reported (Dotson *et al*, 2009). This study did not

examine baseline brain atrophy in relation to incidence of depressive symptoms during follow-up.

Regarding cerebrovascular disease, cross-sectional studies, often from patient samples, consistently show associations between white matter lesions (WMLs) and depression (Greenwald *et al*, 1998; Ikram *et al*, 2010; Kumar *et al*, 2000; Kumar *et al*, 2002). The results from longitudinal population-based studies are conflicting. Three studies using MRI reported that WMLs increased the risk of subsequent depression (Godin *et al*, 2008; Steffens *et al*, 2002; Teodorczuk *et al*, 2007) while two could not confirm these findings (Ikram *et al*, 2010; Versluis *et al*, 2006).

It has been suggested that cortical atrophy and WMLs represent two relatively independent pathways to major depression in the elderly (Kumar *et al*, 2000; Kumar *et al*, 2002). This hypothesis was based on a cross-sectional clinical study, and has not been tested in longitudinal population-based studies. We examined brain atrophy and WMLs on computed tomography (CT) in a population-based sample of elderly without depression or dementia who were followed for five years for the occurrence of major and minor depression.

MATERIALS AND METHODS

Participants

The study sample has been described in detail (Simoni *et al*, 2008), and was derived from the Prospective Population Study of Women (PPSW) (Bengtsson *et al*, 1973) and the Gerontological and Geriatric Population Studies (H70) (Skoog, 2004) in Gothenburg,

Sweden. The participants were obtained from the Swedish Population Register based on birth date and were born in 1908, 1914, 1918, 1922 and 1930. Persons living in private households and institutions were included. In total, there were 1495 eligible individuals in 2000–2001, and 1051 accepted to participate (response rate 70.3%). Of the 1051, 1018 participated in a psychiatric examination and were invited for a CT scan. Of these, 642 (63.1%) accepted to have a CT scan. The 642 individuals who participated in the CT examination were younger (mean age 74 versus 77 years, $p<0.001$), less often women (71.3% versus 88.0%, $p<0.001$), and had a lower mortality rate from baseline to November 2003 (3.1% versus 11.5%, $p<0.001$) compared to the 376 non-participants (Simoni *et al*, 2008). According to the Swedish Hospital Discharge Register, participants in CT experienced a lower frequency of dementia (0.8% versus 7.7%, $p<0.001$), any psychiatric diagnosis (5.9% versus 13.9%, $p<0.001$) and stroke (5.6% versus 12.3%, $p<0.001$) compared to non-participants. There was no difference in the prevalence of major (3.0% versus 3.5%, $p=0.66$) or minor (9.7% versus 9.6%, $p=0.97$) depression between participants and non-participants.

A follow-up was performed in 2005–2006. Among those surviving, 459 accepted a new psychiatric examination (response rate 84.7%). Those lost to follow-up (83 refusals and 21 who died) were traced for dementia, major and minor depression in the Swedish Hospital Discharge Register. Thus, the sample comprised 563 individuals.

Individuals with dementia ($n=16$; 2.5%), major depression ($n=19$; 3.0%) or a score on the Mini-Mental State Examination (MMSE) less than 26 ($n=44$; 6.9%) at baseline were excluded. Those who developed dementia during follow-up ($n=38$) were also excluded from further analyses, leaving 525 individuals.

Informed consent was obtained from all participants and/or their relatives. The study was approved by the Ethics Committee for Medical Research at the University of Gothenburg.

Neuropsychiatric Examinations and Interviews

Both in 2000–2001 and in 2005–2006, trained psychiatric research nurses performed the semi-structured neuropsychiatric examinations that included ratings of the past month's psychiatric signs and symptoms according to Comprehensive Psychopathological Rating Scale (Asberg *et al*, 1978), which is valid and reliable in elderly populations (van der Laan *et al*, 2005), and Mini-International Neuropsychiatric Interview (Sheehan *et al*, 1998), history of depression and assessment of current medications. Ratings of common signs and symptoms of dementia were also done (e.g. assessments of memory, orientation, general knowledge, apraxia, visuospatial function, understanding proverbs, following commands, naming ability and language) and has been described in detail (Guo *et al*, 2007; Skoog *et al*, 1993).

Cognitive function was also measured with the MMSE (Folstein *et al*, 1975). The psychiatric nurses who performed the examinations were supervised and trained by psychiatrists. In training sessions, nurses and psychiatrists rated symptoms and signs among both participants without dementia and those with dementia. Inter-rater reliability between psychiatrists and nurses was studied in 50 individuals. Kappa values for the presence versus absence of signs and symptoms necessary to diagnose depression were between 0.62 and 1.00 indicating “good” (reference range kappa=0.61-0.80) or “excellent” (kappa=0.81-1.00) agreement.

Measurements of Covariates

At baseline, participants underwent a physical examination and blood tests. Body mass index (BMI) was calculated as kg/m^2 . Systolic and diastolic blood pressure (SBP and DBP) was measured in the right arm in the seated position after 5 minutes' rest using a mercury

manometer. DBP was defined as Korotkoff phase 5. Hypertension was defined as SBP \geq 140 mm Hg and/or DBP \geq 90 mm Hg and/or taking antihypertensive medication. Blood samples were taken after overnight fasting. Plasma glucose and serum cholesterol concentrations were measured. The diagnosis of stroke was based on information from self-reports, close informants and the Swedish Hospital Discharge Register. Diagnoses of myocardial infarction, angina pectoris and claudicatio intermittens were based on self-report and key informant interviews. Diabetes mellitus was defined based on a physician's diagnosis, being on antidiabetic therapy or having two fasting venous or capillary whole blood glucose values \geq 7.0 mmol/L.

Diagnosis of Dementia and Depression

Dementia was diagnosed by geriatric psychiatrists according to the Diagnostic and Statistical Manual of Mental Disorders (DSM), third edition, revised (American Psychiatric Association, 1987) based on symptoms rated by the examiners during detailed neuropsychiatric examinations and close informant interviews, as described (Skoog *et al*, 1993). Diagnoses of major and minor depression according to DSM-IV (American Psychiatric Association, 1994) were determined from algorithms based on symptoms rated by the examiners during the psychiatric examinations. Depression was only diagnosed in the absence of dementia. Major depression excluded a diagnosis of minor depression. According to DSM-IV, major depression includes \geq 5 symptoms of depression during the same 2-week period while minor depression includes 2-4 symptoms during the same 2-week period. The symptoms "depressed mood" or "loss of interest" is required for diagnoses.

The diagnosis of depression for individuals lost to follow-up was based on information from the Swedish Hospital Discharge Register. In Sweden, doctors registers primary and secondary

diagnoses according to the International Classification of Diseases, version 10 (ICD-10) (World Health Organization, 1993) on all patients discharged from hospitals. ICD-10 codes for major depression were F31.4-5/32.2-3/33.2-3 and for minor depression F31.3/32.0-1/32.8-9/33.0-1/33.8-9/34.1/34.8-9/38.0-1/41.2.

CT Scan Rating Methods

The same CT scanner and scanning procedure were used for all scans. Ten millimetre continuous slices were obtained. No contrast enhancement was used. The scans were evaluated by a neurologist (MS) experienced in CT and MRI ratings of WMLs and other cerebral lesions. The neurologist was blind to all clinical data.

The Gothenburg scale was used to rate WMLs (Skoog *et al*, 1994). According to this scale, WMLs are defined as periventricular or subcortical areas of decreased attenuation below that expected for normal white matter. The changes were always diffusely distributed within the white matter. The scale gives a global measure of WMLs in the brain, and has been used in our epidemiological studies since 1986 (Skoog *et al*, 1994). It is a 0–3 point scale which takes into account the severity of the hypodensity of WMLs, including 0 (absence of any attenuation), 1 (mild signal attenuation), 2 (moderate signal attenuation), and 3 (severe signal attenuation).

A subjective scale with three grades (absent, mild or moderate and severe) was used to evaluate cortical atrophy. Location of atrophy was categorized as frontal, temporal, parietal or occipital regions, according to the anatomical subdivision (Gustafson *et al*, 2004), and severity was scored according to the extent of sulcal widening (De Leon *et al*, 1980).

The intra-rater Kappa values for the CT assessment were between “fair” ($\kappa = 0.21\text{--}0.40$) and “good” ($\kappa = 0.61\text{--}0.80$) for both WMLs and atrophy (Simoni *et al*, 2008). Inter-observer agreement between the rater and a neuroradiologist was “fair” ($\kappa = 0.30$) for WMLs, “moderate” for temporal lobe atrophy ($\kappa = 0.43$) and “fair” for the other cortical regions ($\kappa = 0.29\text{--}0.36$).

Statistical Analysis

The statistical analyses on the outcome major depression were based on participants without major depression or dementia at baseline, and who did not develop dementia during follow-up. In the analyses on minor depression, also those with minor depression at baseline and those who developed major depression at follow-up were excluded. WMLs were dichotomized as none or mild versus moderate or severe lesions. Cortical atrophy was dichotomized as absence versus presence of atrophy in each brain region. The comparison group was free from both depression and dementia during follow-up. Sex and age-category (70 versus >70 years) were preselected covariates. Additional covariates that were evaluated included level of education (0–7, 8–10 or ≥ 11 years), baseline stroke, myocardial infarction, angina pectoris, claudicatio intermittens, hypertension and diabetes mellitus, and levels of plasma glucose, serum cholesterol, BMI, SBP and DBP.

The statistical analyses were performed with logistic regression models in two steps: First, we selected the covariates to be included in the final models by analysing the association of each covariate with major depression adjusting for gender. Factors were included in the final models if they were associated at a level of $p \leq 0.10$. This resulted in hypertension and serum cholesterol being included as covariates. Second, multivariate logistic regression analyses were performed estimating the odds of major or minor depression during follow-up in relation

to cortical atrophy or WMLs at baseline and including age, sex, and the covariates selected in step one. Odds ratios (ORs) and 95% confidence intervals (CI) were calculated.

We present the results for three models: Model I without adjustments, Model II adjusting for age, sex and the covariates selected in step one; and Model III including both cortical atrophy and WMLs in the same regression model. As a secondary analysis, we also present the results for Model III performed on a subsample of individuals with MMSE above 25 at both baseline and follow-up.

Statistical analyses were performed using SPSS version 16.0. Results were considered statistically significant at a level of $p < 0.05$.

RESULTS

Baseline characteristics are given in Table 1. Of the 525 participants who were free from major depression and dementia at baseline, 20 developed major depression (19 were diagnosed at the follow-up examination and one from the hospital discharge register). In the analyses on minor depression we also excluded 46 participants with minor depression at baseline as well as those who developed major depression, leaving 471 individuals. Among these, 63 developed minor depression (58 were diagnosed at the follow-up examination and 5 from the hospital discharge register). Results are presented for unadjusted (Table 2, Model I) and adjusted (Table 2, Model II and III) models.

Individuals with any temporal lobe atrophy at baseline more often developed major depression (6.4% versus 2.2%, OR=2.81, 95% CI 1.04–7.55) during follow-up than those with no atrophy (Table 2, Model II) adjusted for confounders (age, gender, baseline hypertension and serum cholesterol). Similarly, individuals with moderate to severe WMLs more often developed major depression (9.6% versus 3.2%, OR=3.21, 95% CI 1.02–10.10) during follow-up (Table 2, Model II) compared to those with no or mild WMLs at baseline. There was no association between brain atrophy or WMLs and development of minor depression.

A logistic regression analysis, adjusted for confounders, and including both temporal lobe atrophy and WMLs at baseline showed that presence of temporal lobe atrophy (OR=2.81, 95% CI 1.04–7.62) and moderate to severe WMLs (OR=3.21, 95% CI 1.00–10.26) independently predicted major depression at follow-up (Table 2, Model III). None of the confounders were significantly associated with major depression in this model. Excluding participants with MMSE less than 26 at follow-up ($n=35$; two with major depression) did not change the main results. Including baseline history of depression ($n=143$) as confounder in the multivariate models did not change the findings.

DISCUSSION

We found that temporal lobe atrophy and moderate to severe WMLs predicted occurrence of major, but not minor, depression during a 5-year follow-up of a population-based sample of elderly. Our findings support the hypothesis that brain atrophy and WMLs represent relatively independent and complementary pathways to the development of major depression in the

elderly (Kumar *et al*, 2000; Kumar *et al*, 2002). The only previous study that has explored this hypothesis is a clinical cross-sectional study, and depression was related to frontal but not temporal lobe atrophy (Kumar *et al*, 2000). Our study adds to this previous study by providing longitudinal data from a population-based sample. Both neurodegenerative and cerebrovascular processes may thus be important for the development of major depression in the elderly.

Only one previous longitudinal population-based study has examined the association between brain atrophy at baseline and depressive symptoms or syndromes during follow-up (Ikram *et al*, 2010). This study reported cross-sectional associations between temporal and parietal lobe atrophy and depressive symptoms, but no association with depressive syndrome or depressive symptoms during follow-up. A longitudinal MRI study on elderly without dementia examined the association between mean depressive symptom scores over time and change in brain atrophy over nine years (Dotson *et al*, 2009). In that study (Dotson *et al*, 2009), cross-sectional analyses indicated associations between temporal and frontal lobe atrophy and depressive symptoms, and longitudinal analyses indicated an association between volume decline in frontal white matter and mean depressive symptom scores over time. However, in contrast to our study, brain atrophy at baseline was not reported in relation to occurrence of depression or depressive symptoms during follow-up. The somewhat inconsistent findings between these previous longitudinal studies and our study may partly be explained by the fact that we report separately on major and minor depression rather than on depressive syndrome or depressive symptoms.

Presence of temporal lobe atrophy, but not atrophy in other brain areas, predicted major depression. Support for the importance of the temporal lobe in major depression is found in

cross-sectional hospital-based MRI studies reporting that temporal lobe atrophy is associated with major depression in elderly patients (Dolan *et al*, 1986; Rabins *et al*, 1991; Rainer *et al*, 2006; Wurthmann *et al*, 1995). However, major depression in cross-sectional MRI studies has also been associated with other brain regions (Almeida *et al*, 2003; Dolan *et al*, 1986; Rabins *et al*, 1991; Rainer *et al*, 2006; Schweitzer *et al*, 2001; Wurthmann *et al*, 1995), including atrophy in the frontal and parietal lobes. Differences between these cross-sectional studies and our longitudinal study, which may explain the inconsistent findings include the estimate of atrophy (i.e. measures of sulcal widening, visual rating of cortical atrophy, estimates of CSF volume etc), the severity of atrophy (e.g. any atrophy, severe atrophy), the choice of brain regions studied, the diagnostic criteria for depression, and the age of the samples.

It is well-known that fronto-subcortical circuits are important for depression (Drevets *et al*, 2008; Ressler and Mayberg, 2007). These circuits include limbic structures such as the amygdala which has substantial projections to the temporal cortex (Price and Drevets, 2010). Damage to fronto-subcortical circuits may be caused by subcortical lesions or atrophy in cortical structures. Thus, one explanation for the associations between WMLs and temporal lobe atrophy and major depression could be that these alterations in the brain have caused damage to fronto-subcortical circuits.

The cellular mechanisms that underlie the association between brain atrophy and depression in the elderly remain largely unknown. Due to the few post-mortem studies performed, the results should be interpreted cautiously. Nevertheless, in relation to temporal lobe involvement, changes in myelination, axonal growth and synaptic function have been reported in the temporal cortex of patients with major depression (Aston *et al*, 2005). Atrophy in the temporal lobe may also reflect hippocampal alterations. Post-mortem studies have reported on

cellular alterations such as neuronal death, neuronal shrinkage, decrease in dendritic and glial density, and cell death in the frontal cortex and hippocampus of those with major depression (Hercher *et al*, 2009; Khundakar and Thomas, 2009). Further support for a role of the temporal lobe in major depression is findings from a functional MRI study showing that the temporal lobes are involved in regulation of emotions (Beauregard *et al*, 1998). Hippocampal atrophy in depression (Pittenger and Duman, 2008) and during normal aging (Lupien *et al*, 1998) has been associated with increased cortisol levels, potentially reflective of physiological and environmental stress. One animal study indicated that psychosocial stress could be related to cell death in the temporal cortex (Lucassen *et al*, 2004).

WMLs predicted major depression in addition to the effect of temporal lobe atrophy. Cross-sectional associations between WMLs and depression have been shown repeatedly, especially for MRI and mainly from clinical studies (Greenwald *et al*, 1998; Ikram *et al*, 2010; Kumar *et al*, 2000; Kumar *et al*, 2002), although not all studies support this association (Rainer *et al*, 2006; Skoog *et al*, 1994). Our finding that WMLs preceded major depression is in agreement with three previous longitudinal studies using MRI (Godin *et al*, 2008; Steffens *et al*, 2002; Teodorczuk *et al*, 2007). However, two longitudinal population-based studies did not support this association (Ikram *et al*, 2010; Versluis *et al*, 2006). WMLs most likely reflect ischemic demyelination (Pantoni and Garcia, 1997). The association between depression and changes in the white matter has given rise to the “vascular depression” hypothesis (Alexopoulos *et al*, 1997) which postulates that depression in the elderly may have a vascular origin. Our results support this view, especially regarding major depression.

Brain atrophy and WMLs did not predict occurrence of minor depression during follow-up, suggesting that this less severe form of depression may have a different etiology than major

depression (Beekman *et al*, 1995). Our findings suggest that structural neuropathological changes in the temporal lobes and WMLs are related to more severe depression rather than to the milder forms. This is supported by cross-sectional studies suggesting that temporal lobe atrophy (Kumar *et al*, 1998; Shah *et al*, 1998) may be related to severity of depression rather than to depression *per se*. However, minor depression was related to frontal lobe atrophy in the study by Kumar *et al* (1998). Differences may be explained by differences between population-based and clinical studies, cross-sectional and longitudinal studies and the use of somewhat different diagnostic criteria. We diagnosed minor depression based on DSM-IV criteria (n=58) or ICD codes (n=5), while Kumar *et al* (1998) only diagnosed according to DSM-IV with the additional requirement that the symptoms should be present for at least one month. Thus, they may have looked at a more severe form of the disease compared to our study. Further, they included only those who had their first onset of depression after age 60.

The strengths of the study include the population-based sample, the longitudinal design and the comprehensive examinations performed by experienced psychiatric nurses. Furthermore, to minimize the risk that brain changes reflected preclinical dementia rather than depression, we excluded participants with a MMSE score less than 26 both at baseline and at follow-up. We will discuss some limitations and methodological issues. First, participants in the CT scans had a lower mortality rate, less often stroke, dementia and other mental disorders according to the Swedish Hospital Discharge Register compared to non-participants (Simoni *et al*, 2008) and may thus be healthier than the general population. However, there were no differences in hospitalization with major or minor depression between participants and non-participants. Second, we cannot exclude the possibility that those who developed depression would have developed dementia if we had included a longer follow-up. Third, some of the subgroups were rather small and results should be interpreted cautiously. Future studies on

larger samples are necessary to confirm our findings. Fourth, attrition is an issue that must be considered in follow-up studies. However, our attrition rate at follow-up was only 15.3%. To further reduce the influence of attrition, we used the Swedish Hospital Discharge Register to obtain diagnoses of depression in those lost to follow-up. However, this probably still underestimates the true incidence, as most cases of depression are not hospitalized. Fifth, visual rating of cortical atrophy and WMLs on CT is a rather crude method. The inter-rater Kappa values for agreement between the rater and a neuroradiologist was “fair” ($\kappa = 0.30$) for WMLs, “moderate” for temporal lobe atrophy ($\kappa = 0.43$) and “fair” for the other cortical regions ($\kappa = 0.29\text{--}0.36$). If anything, the lack of perfect agreement most likely attenuates the observed relationship. CT is less sensitive than MRI in detecting WMLs and is more severely affected by bone hardening artifacts, which is particularly severe in the region of the medial temporal lobe (Frisoni, 2001). Nonetheless, visual rating of the temporal lobe has been found to be highly predictive for the discrimination of Alzheimer’s disease versus control patients (Wahlund *et al*, 2000). Further, CT is better in delineating clinically relevant WMLs (Lopez *et al*, 1995) and comparable to MRI in detecting brain atrophy (Frisoni, 2001) and may be more suitable than MRI for the elderly, as it is less sensitive to motion artefacts. CT is also the most used brain imaging tool world-wide. Sixth, all scans were rated by a single person which may increase the risk for systematic error.

The results from this longitudinal population-based study suggest that temporal lobe atrophy and WMLs represent independent pathways to major depression in the elderly, even if they may have shared risk factors. The finding that neurodegeneration and cerebrovascular disease are related to major depression may have implications for treatment and prevention, as these changes have been related to preventable factors, such as hypertension and obesity.

DISCLOSURE

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Table 1 Baseline Characteristics of the Study Population and Those Who Developed Major or Minor Depression Over 5 years Follow-up

	Baseline sample (<i>n</i> =525 ^a)	Major depression during follow-up (<i>n</i> =20 ^a)	Minor depression during follow-up (<i>n</i> =63 ^a)
Characteristic	<i>n</i> (%) or mean (SD)	<i>n</i> (%) or mean (SD)	<i>n</i> (%) or mean (SD)
Women	360 (68.6%)	17 (85%)	49 (77.8%)
Age (years)	72.7 (4.6)	73.2 (4.6)	72.7 (4.4)
Age distribution (years)			
70	381 (72.6%)	13 (65%)	45 (71.4%)
78	86 (16.4%)	5 (25%)	13 (20.6%)
82	47 (9.0%)	2 (10%)	4 (6.3%)
86	11 (2.1%)	0	1 (1.6%)
Education			
0-7 years	156 (30.8%)	6 (33.3%)	21 (34.4%)
8-10 years	182 (35.9%)	9 (50%)	16 (26.2%)
≥ 11 years	169 (33.3%)	3 (16.7%)	24 (39.3%)
MMSE	28.4 (1.1)	28.1 (1.2)	28.1 (1.1)
Blood pressure, SBP/DBP (mmHg)	156.7(22.4)/82.7(11.5)	156.2(15.5)/81.2(11.1)	151.2(24.1)/80.1(11.9)
Serum cholesterol (mmol/l)	5.90 (1.10)	5.52 (1.00)	6.06 (1.32)
Antidepressant treatment	37 (7.0%)	11 (55.0%)	3 (4.8%)
Cardiovascular drugs ^b	148 (28.2%)	8 (40.0%)	17 (27.0%)
Moderate-severe WMLs	52 (9.9%)	5 (25%)	5 (7.9%)
Cortical atrophy	296 (56.4%)	14 (70%)	40 (63.5%)

DBP, diastolic blood pressure; MMSE, Mini-Mental State Examination; SBP, systolic blood pressure; WMLs, white matter lesions.

^aDue to missing values, the group sizes of the following measures were reduced: baseline sample

- education ($n=507$), blood pressure ($n=520$), serum cholesterol ($n=522$); major depression - education ($n=18$); minor depression - education ($n=61$), serum cholesterol ($n=62$).

^bAntihypertensives, anticoagulants, cholesterol- and triglyceride lowering, and antidiabetics

Table 2 Cortical Atrophy and WMLs at Baseline in Relation to 5-year Occurrence of Major Depression.^a

	<i>n</i> / <i>N</i> ^b (%)		OR (95% CI)		
	No atrophy	Atrophy	Model I (unadjusted)	Model II ^c	Model III ^d
Atrophy in any cortical area	6/229 (2.6%)	14/296 (4.7%)	1.85 (0.70–4.88)	-	-
Atrophy in cortical areas					
Temporal	7/322 (2.2%)	13/203 (6.4%)	3.08 (1.21–7.85) ^e	2.81 (1.04-7.55) ^e	2.81 (1.04-7.62) ^e
Frontal	10/290 (3.4%)	10/235 (4.3%)	1.24 (0.51–3.04)	-	-
Parietal	13/383 (3.4%)	7/142 (4.9%)	1.47 (0.58–3.78)	-	-
Occipital	16/440 (3.6%)	4/85 (4.7%)	1.31 (0.43–4.02)	-	-
	No/mild WMLs	Moderate/severe WMLs			
WMLs	15/473 (3.2%)	5/52 (9.6%)	3.25 (1.13–9.34) ^e	3.21 (1.02-10.10) ^e	3.21 (1.00-10.26) ^e

CI, confidence interval; ORs, odds ratios; WMLs, white matter lesions.

^aThe analyses are based on participants with a baseline status of no dementia or depression and MMSE score >25.

^b*n*/*N* (within each group of cortical atrophy or WMLs) = (participants with major depression during follow-up)/(total number of participants)

^cAdjusted for age, gender, baseline hypertension and serum cholesterol. Results for Model II are presented only where Model I is significant.

^dTemporal lobe atrophy and WMLs in the same regression analysis. Adjusted for the same factors as in Model II.

^e*p*<0.05, Cox regression model.