**CONCISE REPORT**

**White matter hyperintensities on brain magnetic resonance in systemic sclerosis**

F Sardanelli, A Iozzelli, B Cotticelli, C Losacco, M Cutolo, A Sulli, F Nobili, G Rodriguez


**Objective:** To evaluate the brain status of patients with systemic sclerosis (SSc).

**Methods:** Fourteen female patients with SSc aged 24–74, with a disease duration of 1–12 years and without other relevant systemic diseases, were enrolled. All patients and an age matched female control group (CG) of 14 clinically normal subjects, underwent brain magnetic resonance examination at 1.5 T; spin echo proton density weighted images were evaluated. Mann-Whitney U and Spearman rank correlation tests were used for statistical analysis.

**Results:** 170 white matter hyperintensities ≥2 mm in diameter were counted in the patient group (range 0–75, mean 12.1, median 4.5), only 13 in the CG (0–2, 0.9, 1, respectively), with a significant difference (p = 0.011). Moreover, 208 white matter hyperintensities <2 mm were found in the patient group (0–38, 14.9, 8, respectively), only 31 in the CG (0–7, 2.0, 1, respectively), with a significant difference (p = 0.006). No statistically significant correlation between the number of hyperintensities and either patient’s age or disease duration was observed.

**Conclusion:** White matter hyperintensities are more common in patients with SSc than in a CG. These findings might be related to oblitative microvascular processes due to the disease. Early brain involvement in patients with SSc may occur.

**METHODS**

We studied 14 female patients with SSc (aged 24–74 years) with a disease duration of 1–12 years, who had no serious diseases other than SSc. All the patients had received a diagnosis of SSc according to the criteria of the American Rheumatism Association. Exclusion criteria included severe or uncontrolled arterial hypertension; uncontrolled diabetes mellitus; relevant renal, respiratory, or hepatic failure; and severe anaemia. Three patients included were affected by mild arterial hypertension and were treated with angiotensin converting enzyme inhibitors or with a calcium blocking agent. One further patient was affected by moderate hypertension and was regularly treated with an angiotensin converting enzyme inhibitor and diuretics. Blood pressure was well controlled in these patients by the present treatment. None of the patients included was diabetic, but one showed impaired glucose tolerance. Renal function, as assessed by routine creatinine and blood urea nitrogen assays, was normal in all cases. These patients had no history of cerebrovascular accidents, head trauma, or other major neurological or psychiatric disease. Fourteen age matched clinically normal women were studied as a control group (CG).

**Results:**

Fourteen female patients with SSc aged 24–74, with a disease duration of 1–12 years, who had no serious diseases other than SSc. All the patients had received a diagnosis of SSc according to the criteria of the American Rheumatism Association. Exclusion criteria included severe or uncontrolled arterial hypertension; uncontrolled diabetes mellitus; relevant renal, respiratory, or hepatic failure; and severe anaemia. Three patients included were affected by mild arterial hypertension and were treated with angiotensin converting enzyme inhibitors or with a calcium blocking agent. One further patient was affected by moderate hypertension and was regularly treated with an angiotensin converting enzyme inhibitor and diuretics. Blood pressure was well controlled in these patients by the present treatment. None of the patients included was diabetic, but one showed impaired glucose tolerance. Renal function, as assessed by routine creatinine and blood urea nitrogen assays, was normal in all cases. These patients had no history of cerebrovascular accidents, head trauma, or other major neurological or psychiatric disease. Fourteen age matched clinically normal women were studied as a control group (CG).

**Conclusion:**

White matter hyperintensities are more common in patients with SSc than in a CG. These findings might be related to oblitative microvascular processes due to the disease. Early brain involvement in patients with SSc may occur.

**S**ystemic sclerosis (SSc) is a progressive multisystem connective tissue disease of unknown aetiology, characterised by increased amounts of collagen in the blood vessels, skin, and visceral organs. Gastrointestinal tract, lungs, heart, and kidneys are mostly affected. The vascular changes begin with an injury to the endothelial cells and an intimal proliferation with fibrous thickening of the media, leading to oblitative microvascular disease.

The prevalence and severity of vascular involvement of the central nervous system (CNS) in patients with SSc is still a matter of discussion. It is generally considered to be uncommon or as a secondary consequence of hypertension, uremia, pulmonary dysfunction, and steroid treatment; signs and symptoms of CNS involvement were infrequently reported in the large clinical series of patients with SSc of Tuffanelli and Winkelmann.4 Moreover, cerebrovascular lesions were not found more often in patients with SSc than in an age matched control group at pathological examination.5 However, several authors have reported CNS abnormalities in patients with SSc, dating from the pathological observations of Steven in 1898.6 7 Cases of carotid and intracranial arteritis8 have been reported, also. Neuropsychiatric manifestations, such as loss of memory, disorientation, depression, delusions, hallucinations, and reduced mental acuity were reported in the late phase of the disease, while transient ischaemic attacks, ischaemic stroke, and haemorrhage9 have been described as a primary consequence of SSc in recent years.

Magnetic resonance (MR) imaging is considered to be the most sensitive diagnostic technique for detecting symptomatic and asymptomatic lesions in the brain, as demonstrated in multifocal diseases such as ischaemic encephalopathy and multiple sclerosis. Thus, this study aimed at evaluating the brain status of patients with SSc by MR imaging.

**Abbreviations:** CG, control group; CNS, central nervous system; MR, magnetic resonance; SSc, systemic sclerosis
density and T2 weighted sequences in detecting supratentorial periventricular inflammatory lesions; however, taking into account the low sensitivity of fluid attenuated inversion recovery in detecting subcortical brain stem inflammatory lesions, proton density weighted sequences were finally preferred and used in this study.

Two readers counted the white matter hyperintensities in a randomised order, without knowing the patient/control status of each subject, using a remote console (Magic View, Siemens, Erlangen, Germany). All the abnormalities were electronically measured along the longest diameter to distinguish those ≥2 mm from those <2 mm in diameter.

For statistical analysis, the Mann-Whitney U test and Spearman rank correlation coefficient were used.

RESULTS

One hundred and seventy white matter major hyperintensities (≥2 mm in diameter) were counted in the patient group (range 0–75, mean 12.1, median 4.5), only 13 in the CG (range 0–2, mean 0.9, median 1), with a significant difference (p = 0.011, Mann-Whitney). Two readers counted the white matter hyperintensities in a randomised order, without knowing the patient/control status of each subject, using a remote console (Magic View, Siemens, Erlangen, Germany). All the abnormalities were electronically measured along the longest diameter to distinguish those ≥2 mm from those <2 mm in diameter.

For statistical analysis, the Mann-Whitney U test and Spearman rank correlation coefficient were used.

Figure 1 Proton density weighted brain MR image with major hyperintensities (≥2 mm in diameter, white arrows) and minor hyperintensities (<2 mm in diameter, black arrows) in the deep white matter of a patient with SSc.

DISCUSSION

Our study demonstrated that white matter hyperintensities are more common in patients with SSc than in the CG. The number and size of the hyperintensities suggest abnormalities in the territories supplied by the smaller branches and the perforating vessels. These findings confirm the hypothesis of a previous direct microangiopathic injury of the brain and are in accordance with previously described brain involvement in four cases of a particular cutaneous form of SSc, the so-called “en coup de sabre” linear scleroderma (in all four cases MR imaging demonstrated brain hyperintensities), even though Stone et al performed brain biopsy and found “evidence of an inflammatory process which may be amenable to immunosuppressive treatment”.

We know that the underlying pathogenesis of these brain abnormalities could be explained by a range of different mechanisms, which might act individually or, together, in producing the lesions. The first of these proposed pathways are non-immune reactions, with cell mediated activity and autoantibody production likely to account for the isolated cases of vasospasm (and associated autonomic dysfunction). The second mechanism is vascular, with changes in endothelial cells and basement membrane, causing a non-inflammatory microangiopathy of the vasa nervorum, a tendency for platelets to aggregate, and a final ischaemic CNS injury—although the initial precise causal event remains unclear. The third pathway consists of alterations of the collagen metabolism, proliferation of specific fibroblast subpopulations, and invasion into tissue and fibrosis, though this seems unlikely to produce CNS abnormalities because of the paucity of connective tissue in the brain. The role of cerebral vasculitis in SSc remains unknown, as previous necropsy reports have rarely described such alterations. All these reports and hypotheses deserve further investigation, both histologically and clinically, to elucidate better the pathophysiology of CNS involvement in SSc.

In conclusion, our study shows that brain MR hyperintensities are more common in patients with SSc than in the CG, supporting the hypothesis of an early and frequent brain involvement in patients with SSc.

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