Cerebral amyloid angiopathy-related infarcts imitating high-grade differentiation of a benign meningioma

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DESCRIPTION

Meningiomas are divided, by the WHO, into three groups: benign (grade I), atypical (grade II) and malignant (grade III). Although some overlapping features can occur, atypical meningiomas are more likely to exhibit heterogeneous density or intensity, heterogeneous contrast enhancement and relatively large perifocal oedema on CT scan and MRI.1

Meningiomas typically receive their blood supply from dural arteries, although larger meningiomas can also have a blood supply from leptomeningeal or parenchymal branches. Cerebral amyloid angiopathy (CAA) results from β-amyloid deposition in the walls of leptomeningeal and cortical arteries.2 It usually causes lobar intracerebral haemorrhage and often occurs in association with Alzheimer’s disease.

Rarely, CAA has been documented to be associated with inflammatory changes, sometimes thought to be co-existent primary central nervous system vasculitis or giant cell arteritis, although most literature has focused on probable inflammatory changes secondary to β-amyloid deposition, sometimes described as amyloid β-related angiitis.3–6 We could not find any specific association of CAA with Wegener’s granulomatosis in the literature.

We present serial brain imaging findings describing the evolution of changes from a stable benign meningioma to a lesion imitating an atypical meningioma, which was surgically removed and subsequently shown to be affected by amyloid angiopathy.

A 59-year-old woman presented with a short history of right-sided tinnitus, hearing loss and sinusitis. Her other significant known condition was Wegener’s granulomatosis involving the lungs, diagnosed in another institution. Brain MRI in January 2011 (figure 1) was negative for vestibular schwannoma, but demonstrated an incidental 14 mm parafalcine meningioma in the left frontal region on the background of small vessel cerebrovascular disease. MRI performed in March 2013 (figure 2), due to acute neurological symptoms, demonstrated extensive abnormalities in the right parieto-occipital lobe with further haemorrhage in the left frontal lobe; these were assumed to be due to the patient’s known Wegener’s granulomatosis, rather than from potential CAA. There was some increase in the size of the meningioma, which continued to show homogeneous enhancement. A follow-up MRI in April 2014 (figure 3) showed further increase in size of the meningioma to 28 mm, still with homogeneous enhancement and without any significant surrounding brain changes.

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In November 2014, the patient had further clinical deterioration with the new-onset right-sided neurological symptoms. MRI (figure 4) showed the meningioma as having a heterogeneous appearance, with necrosis and peripheral enhancement. The adjoining brain also appeared grossly heterogeneous and necrotic, with peripheral enhancement and significant mass effect. A neurosurgical review suspected the changes to be those of a high-grade tumour progression, and the meningioma was resected. Neuropathological examination found CAA involving vessels in the meninges, cortex and within the meningioma, resulting in infarcts in a WHO grade I meningioma and adjacent cortex without any invasion or other features of an atypical meningioma (figure 5). No evidence of vasculitis or Wegener’s granulomatosis was found. A follow-up MRI in February 2015 (figure 6) demonstrated residual blood products with minimal postoperative enhancement and resolution of most of the surrounding changes.

The current case shows unusual appearances in a grade I meningioma resulting from leptomeningeal, cortical and...
meningioma vessel involvement in amyloid angiopathy. These radiological appearances can easily be misinterpreted as an atypical meningioma. Involvement of blood vessels in a meningioma has not been previously described in CAA; it is otherwise a common condition in the elderly and should be kept in mind if other associated brain changes, such as lobar haemorrhages, are present in patients with a meningioma.

Competing interests None declared.
Patient consent Obtained.
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REFERENCES

Learning points
- Infarcts in a WHO grade I meningioma can mimic high-grade differentiation, prompting neurosurgical concern.
- Cerebral amyloid angiopathy (CAA) can affect not only normal brain but also a pre-existing meningioma.
- Presence of brain changes consistent with CAA should prompt the suspicion of similar changes in a necrotic meningioma.

Figure 6  MRI in February 2015 (postoperative). T2 axial (A and B), fluid-attenuated inversion recovery coronal (C), postcontrast T1 axial (D and E) and coronal (F) images showing residual changes with some haemosiderin deposition and minor residual enhancement (white arrow).