CT findings in Fahr’s disease
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DESCRIPTION
A 65-year-old man presented to our services with features of dementia and movement disorder for 2 years. The symptoms had worsened over the past month with development of auditory hallucinations. CT demonstrated extensive calcification in bilateral basal ganglia, grey–white matter junctions and dentate nuclei. Serum calcium, phosphate and parathormone levels were normal. Fahr’s disease was diagnosed based on the clinical and imaging findings.

Fahr’s disease is an autosomal dominant inherited disorder characterised by basal ganglia and extraganglionic calcification, extrapyramidal symptoms and psychosis.¹ Usual age of presentation is 40–60 years with no gender predilection.¹ Psychosis is proportionate to degree of calcification and cerebral atrophy. Calcification typically occurs in the lateral part of the globus pallidus, dentate nuclei and caudate nuclei.¹ ¹ Other sites include grey–white matter junction, white matter tracts and thalami (figure 1). However, the sites of calcification are not specific and a similar pattern may be found in hypoparathyroidism, pseudohypoparathyroidism and hyperparathyroidism.² Normal calcium, phosphate and parathormone levels in Fahr’s disease aid in diagnosis.¹ ¹² Other differentials include senescent calcification (punctate, less extensive, occurring in the medial globus pallidus), HIV encephalopathy (cerebral

Figure 1  Fahr’s disease (Striopallidodentate calcinosis). Axial non-contrast CT depicts bilateral symmetric calcifications in cerebellar foliae (thin black arrows) (A). Also note calcification in both dentate nuclei (white arrowheads) (A). There is extensive bilateral calcification in head of caudate nuclei (thick black arrows) (B and C), putamina and lateral globus pallidi (thin white arrows) with relative sparing of medial globus pallidi and thalami (B). There is calcification at grey–white matter junction as well (black arrowheads) (B). Linear calcification is seen to extend perpendicularly from the caudate nuclei (thick white arrows) (D).

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atrophy, low CD4 counts), Cockayne syndrome (dwarfism, microcephaly, mental retardation, cerebral atrophy), and as sequelae to intrathecal chemotherapy and radiotherapy to the skull. 2

Non-contrast CT is the most sensitive modality and there is lack of enhancement in regions of calcification. 2 Prognosis is poor since the disease is progressive and there is no specific treatment. Antipsychotics may be used; but patients with Fahr’s disease are more susceptible to adverse effects such as malignant neuroleptic syndrome. 3

Competing interests None.

Patient consent Obtained.

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REFERENCES

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