Bilateral central and centrocaecal scotomata due to mass lesions

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SUMMARY Unilateral central or centrocaecal scotoma may result from optic nerve compression. However, such defects bilaterally usually indicate non-compressive optic neuropathy of toxic or nutritional, hereditary, or demyelinating origin. Three cases are reported of patients who presented with somewhat atypical bilateral central or centrocaecal scotomata and were found to have suprasellar mass lesions demonstrated by CT scan and confirmed neurosurgically.

Unilateral central or centrocaecal scotoma may result from optic nerve compression as well as optic neuritis.1 Bilateral central or centrocaecal scotomata (which must be distinguished from bitemporal hemianopic scotomata of chiasmal origin) usually indicate optic neuropathy of toxic or nutritional,2 hereditary,3,4 or demyelinating origin and have only rarely been associated with mass lesions. We report 3 patients who presented with somewhat atypical bilateral central or centrocaecal scotomata and were found on computed tomography (CT) scan to have a suprasellar mass, confirmed by neurological exploration and pathological examination.

Case report

CASE 1 A 45-year-old male had noted decreased visual acuity to 20/30 OU in 1970, with clearing over the subsequent several months. In 1975 there was recurrence, with visual acuity 20/30 OU and central scotoma to 16/300 red object only OU (Fig. 1). The discs were minimally pale. Fluorescein angiography was normal. General physical and neurological examinations, cerebrospinal fluid analysis, skull, orbit and optic

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channel x-rays, and brain scan were normal. The visually evoked response (VER) to pattern stimulus showed prolonged latency to 140 ms OU.

His vision continued to fluctuate, and in December 1976 visual acuity was 20/50 OU. External examination and motility were normal. Four of six Hardy-Rand-Rittler pseudoisochromatic (AOHR) colour plates were interpreted correctly OU. Visual fields showed centrocaecal scotomata to 10 mm red (not to 3 mm white) at 1 meter and a left upper homonymous hemianopic defect (Fig. 2). The pupils were sluggish without relative afferent defect, and there was mild temporal disc pallor OU.

Skull and optic canal x-rays and optic canal polytomies were normal. CT scan showed a suprasellar mass. At right frontal craniotomy a cystic chiasmal mass was found, presumed a glioma and not biopsied.

The patient was followed up without radiotherapy and did well, with improvement in visual acuity OD to 20/20-1 (OS 20/60). CT scan in March 1979 was unchanged. In May 1979 he complained of severe headache. Papilloedema was found, and CT showed expansion of the tumour, displacement of the third ventricle, and obstruction of the foramen of Monro (Fig. 3). A second craniotomy was done, and a biopsy taken from a chiasmal mass was considered to show cerebral histiocytosis. Radiotherapy was begun. His vision remained unchanged at 31 March 1981.

case 2

A 31-year-old male had noted painless central blurring of vision with dyschromatopsia OS since March 1979, fluctuating clearly in response to prednisone. In June 1979 he developed a similar but less severe involvement OD. CT scan was negative without high resolution orbital views. In July 1979 bilateral central scotomata were found (Fig. 4) to 2/1000 white OD and 10/1000 white OS, with visual acuity 20/20 OD, 20/200 OS, with mild general depression of field (5 white not seen OS).

In August 1979 visual acuity was 20/20-1 OD and 20-70 OS. External examination and motility were normal. AOHR colour plates were interpreted correctly 2½ of 6 OD and none OS. Visual fields showed temporal nonhemianopic paracentral relative scotoma to 10/1000 red OD and centrocaecal scotoma OS to 3/1000 white and relative to 10/1000 red (Fig. 5). There was a mild relative afferent pupillary defect

Fig. 2  Case 1. Central visual fields show centrocaecal scotomata to 10/1000 red and left upper homonymous hemianopic defect to 3/1000 white, 8 December 1976.

Fig. 3  Case 1. Coronal CT shows a suprasellar enhancing mass (black arrows) with intrasellar invasion. The mass indents the 3rd ventricle (white arrows) with resulting hydrocephalus.
OS. There was mild left and questionable right disc pallor. General physical and neurological examinations were negative, as were complete blood count, erythrocyte sedimentation rate, lupus erythematosus cell test, and cerebrospinal fluid analysis. VER indicated left optic neuropathy.

Fig. 4  Case 2. Bilateral central scotomata to 2/1000 white OD and 10/1000 white OS, July 1979 (by Dr Thomas Kuhns).

Fig. 5  Case 2. Temporal non-hemianopic paracentral scotoma to 10/1000 red OD and centrocaecal scotomata to 3/1000 white and 10/1000 red OS, 23 August 1979.

Fig. 6  Case 2. Bitemporal hemianopic scotomata with scotomatous nasal central depression OS, 18 September 1979.
After tapering of the prednisone his vision worsened, and one month later his visual acuity was 20/20 OD, 20/800 OS, with bitemporal hemianopic scotoma and scotomatous nasal central depression as well OS. (Fig. 6). CT scan showed a suprasellar mass (Fig. 7A and 7B). At craniotomy there was a ‘large mass in the suprasellar area between the optic nerves, touching both optic nerves and most likely originating in the optic chiasm . . . grossly a typical glioma . . . grayish in color with vascularity resembling nerves over it.’ Biopsy confirmed optic glioma. Radiotherapy was started. He was not followed up.

**Case 3**

A 22-year-old male had noted fluctuating progressive visual blurring over 2 months since September 1981. On 13 November visual acuity was 20/25-1 (slowly) OD and 20/50-2 OS. External examination and motility were normal. The visual fields showed generalised constriction with bilateral centrocaecal scotomata to 3/1000 white, relative to 10/1000 red (Fig. 8). AOHR colour plates were interpreted correctly 0 of 6 OU. The pupils were slightly sluggish with a possible right relative afferent defect. The posterior poles were normal. General physical and neurological examinations were otherwise unremarkable.

The complete blood count, ESR, antinuclear antibody test, serology, and vitamin B₁₂ levels were all within normal limits. X-rays of skull and optic canals were normal. CT scan showed a suprasellar mass (Figs. 9, 10). At craniotomy a cystic cranio-pharyngioma was found (Fig. 11) and drained. Four days later visual acuity was 20/20 OU, with normal visual fields.

**Discussion**

These 3 patients presented initially with somewhat atypical bilateral central or centrocaecal scotomata. There was no history suggestive of non-compressive bilateral optic neuropathy of toxic or nutritional, hereditary, or demyelinating origin, and suprasellar mass lesions were demonstrated (intrinsic in cases 1 and 2 and extrinsic in case 3). Such bilateral central or centrocaecal scotomata (which must be distinguished from bitemporal hemianopic scotomata) have only rarely been reported in patients with mass lesions. In 1923 in his Bowman lecture de Schweinitz reviewed past studies of visual field defects occurring with pituitary tumours. He mentioned only one report of bilateral centrocaecal scotomata by Lohman (p. 39). In 1939 Henderson reviewed the visual field changes in 243 patients operated on by Cushing for pituitary adenoma. In 6 cases bilateral central scotomata were recorded but only diagrammatically represented. Hirsh and Hamlin reported central scotomata in 11% of 59 cases with pituitary tumours without giving further details of the visual field defect and probably unilateral. Hollenhorst and Younge in an analysis of 1000 cases of pituitary adenoma did not record...
bilateral central or centrocaecal scotomata, although such cases may have been included in 27 cases with central or temporal scotoma (both eyes).'

Single cases of bilateral centrocaecal scotomata due to mass lesions were reported by Kelly⁹ (his Fig. 7), Nettleship¹⁰ (his Figs. 49, 50), Schlezinger et al.¹¹ (their Fig. 6, with irregular shape), Lloyd¹² (pp. 177–8, where smoking was a possible factor and bitemporal hemianopia supervened with pituitary disease 3 years later), and most recently Kline¹³ (his Fig. 1, with constriction of peripheral isoptres as well, with the finding at surgery of compression by a

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**Fig. 8** Case 3. Bilateral centrocaecal scotomata to 3/1000 white, relative to 10/1000 red, with peripheral depression, 3 November 1981.

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**Fig. 9** Case 3. Axial CT shows the suprasellar cistern enlarged and deformed by a low density mass (black arrows).

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**Fig. 10** Case 3. Coronal CT with metrizamide shows deformity of the lower part of the 3rd ventricle (black arrows) by a suprasellar mass (white arrows).
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As in the 3 cases reported here, it would appear that one must consider intracranial mass lesion(s) as a cause of not only unilateral but also occasionally bilateral central or centrocaecal scotoma. Such an aetiology should be considered in the absence of a history suggesting a toxic or nutritional or hereditary basis or other indication of demyelinating disease. Other atypical features prompting such consideration include a relentlessly progressive course, unusual pattern of scotomata, peripheral visual field depression, especially if hemianopic, or steroid responsiveness, as suggested in our second case. Little or no disc pallor is characteristic of early optic nerve or chiasm compression and does not militate against its consideration or appropriate investigation. A high resolution CT scan should be obtained in such cases of unexplained or atypical central or centrocaecal visual field defects.

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