Bitemporal visual field defects associated with anomalies of the optic discs

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Bitemporal visual field defects are usually associated with lesions of the optic chiasm. The field loss may be caused by pressure, primary neoplasia, arachnoiditis, or demyelinating disease. While the findings of a bitemporal defect should always direct attention to these possibilities, a careful inspection of the optic disc may reveal an ocular cause for the field loss.

The purpose of this communication is to emphasize the connection between inferior conus (Fuchs's coloboma) of the optic disc and a temporal or altitudinal field defect. Usually the conus is associated with inversion of the optic disc so that the retinal vessels emerge from the nasal side rather than the temporal side. Frequently depigmentation of the fundus adjacent to the conus is found; this is sometimes best appreciated in a retinal photograph or with indirect ophthalmoscopy. Difficulties in diagnosis can arise as is shown in the following patient.

Case report

Patient 1, a woman aged 62 years, complained of slight deterioration of the visual acuity in the left eye over a period of about 2 months. The corrected visual acuity had been 6/6 - 1 in the right eye and 6/9 part in the left 2 years before. Apart from mild rheumatoid arthritis she enjoyed good health.

Examination

The visual acuity in the right eye was 6/6 - 1, with $-2 \cdot 25$ D sph., -3 D cyl., axis 5°, and in the left 6/18+, with -3 D sph., -2 D cyl., axis 180° .

The fundi showed bilateral Fuchs's coloboma with inversion of the optic disc. The vessels showed congenital tortuosity. The right retina was normal. The central vein pulsated freely at the disc. The left retina showed evidence of a venous stasis retinopathy with scattered haemorrhages at the posterior pole. The blood pressure was 190/100 (on methyl dopa). The carotid arteries were normal.

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Visual fields (Fig. 1)

RIGHT: upper temporal defect to small stimulus. The defect extended slightly across the midline (Fig. 1a).

LEFT: upper defect extending across the midline, seen only with small stimulus (Fig. 1b).

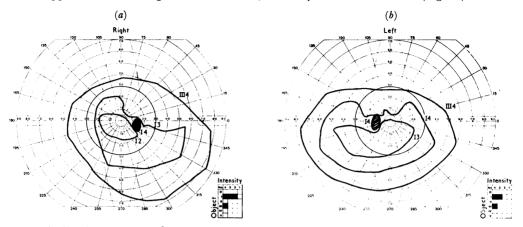


FIG. I (a, b) Case I. Visual fields on November 6, 1969.

Radiography of the skull showed a pituitary fossa at the upper limit of normal size. An air encephalogram showed a doubtful shadow, suggesting a space-occupying lesion beneath both optic nerves anteriorly.

Although it was realized that the visual field defects could be accounted for by the optic disc abnormalities (Fig. 1), the other findings suggested a dual pathology, and an exploratory craniotomy was carried out.

Operation

The chiasm was explored through a right frontal flap. There was extreme tortuosity of both carotid arteries which, with the ophthalmic arteries, had bulged upwards and medially to the optic nerve forming almost an aneurysmal dilatation in each case. The roof of the sella was normal. An attempt was made to separate adhesions between the optic nerve and the carotid artery on both sides to allow the nerve to move away to a limited extent if compression continued.

Result

The post-operative course was uneventful. The venous stasis retinopathy in the left eye cleared completely, and the visual acuity in the left eye improved to 6/9 part. The visual fields remained unaltered.

Material and methods

A series of sixteen patients with conus is summarized in the Table (overleaf, pp. 310 and 311). Two patients (Cases 5 and 8) had a normal disc in the second eye, and one (Case 9) had only a myopic crescent in the second eye. There were thirteen females and three males. Their age ranged between 7 and 72 years. The visual fields of both eyes were examined on the Goldmann kinetic perimeter, if one eye showed conus and/or inversion of the optic disc (Figs 2a, b and 3a, b). Goldmann static perimetry was used on three eyes of two patients who showed field defects, a second curve being constructed using a more myopic correction when retinoscopy suggested an ectatic area of retina in the meridian of the field defect (Fig. 2c). Retinoscopy values tend to vary considerably in an eye with conus on examination slightly away from the fixation axis (Figs 2d, e and 3c, d).

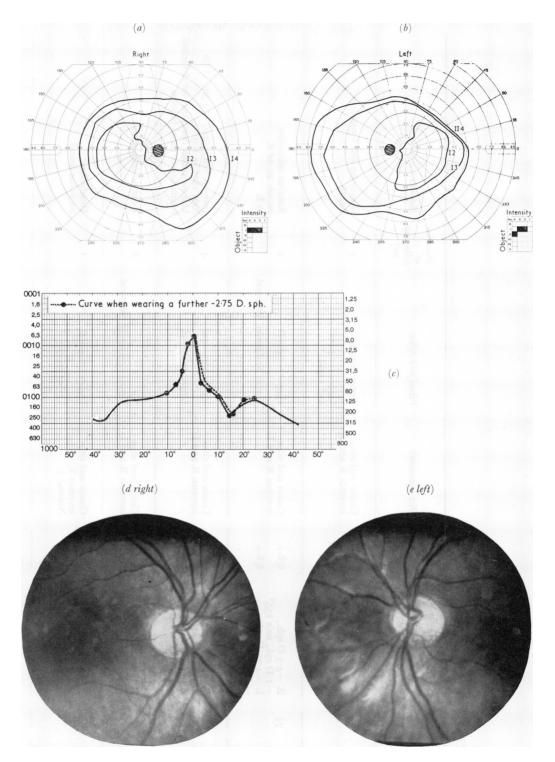


FIG. 2 (a, b) Case 2. Visual fields on April 3, 1971. (c) Case 2. Static perimetry. (d, e) Case 2. Retinal appearances, right and left.

Table		inical f	Clinical particulars of sixteen patients	nts				
Patient No.	Sex	Age (Jrs)	Refraction	Visual acuity	Disc appearance	Visual field defects	Duration of follow-up (Jrs)	Remarks
-	Гц	62	R –2.25 D sph., –3 D cyl., axis 5° L –3 D sph., –2 D cyl., axis 180°	6/6 pt 6/9 pt	Conus and situs inversus R and L	Upper temporal Upper temporal and nasal	3	Headache and visual loss 1a, 1b
8	F4	71	R -4.5 D sph., -1 D cyl., axis 155° L -4.75 D sph., 1 D cyl., axis 10°	6/6+	Conus and situs inversus R and L	Upper temporal Upper and lower temporal	a	Routine refraction 2a, 2b, 2c, 2d R 2e L
en L	۲щ	68	R -2.5 D sph., -0.5 D cyl., axis 180° L +0°5 D sph., -2°5 D cyl., axis 60°	6/9 pt 6/9 pt	Conus and situs inversus R and L	Upper temporal Upper temporal	4	(pre-op. refraction) $-3.50/-1.50$ Axis $180^\circ = 6/9$ Left successful operation for detached retina upper nasal balloon
4	۴ı	19	R + 75 D cyl., axis 90° L -0·5 D sph.	6/6 pt 6/9 + 1	Conus and situs inversus R and L	Nil Upper and lower temporal	æ	Routine refraction
Ω	۲L	64	R –2:5 D sph., –2 D cyl., axis 95° L +0.75 D cyl., axis 170°	6/7-5 pt 6/5	Conus and situs inversus R Normal L	Upper temporal Upper temporal	г	Headaches: no cause found
Q	Γ	46	R -5 D sph., - 2:5 D cyl., axis 85° L -5 D sph., -0:5 D cyl., axis 40°	6/6 pt 6/5 pt	Conus and Situs inversus R and L	Upper temporal Upper and lower temporal and lower nasal	I	Routine refraction 34, 36 R 3d L
7	ъ	33	R -2.75 D sph., -2 D cyl., axis 145° L -0.75 D sph., -1 D cyl., axis 55°	6/6 pt	Inferior nasal conus situs inversus R and L	Upper temporal Upper temporal	H	Routine refraction

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ы.	88	R +9 D sph., +2 D cyl., axis 135° L +8 D sph., +1 D cyl., axis. 180°	6/5 6/5	Inferior conus R Normal L	Upper temporal Nil	-	R and L aphakic refraction
ч	58	R –2 D sph., – 1D cyl., axis 40° L –7.75 D sph., –0.5 D cyl., axis 10°	6/24 +1 6/9	Myopic temporal crescent Inferior nasal conus Situs inversus	Nil Upper temporal defect	61	R detached retina replaced
Ŀч.	72	Not known	6/9 6/18	Conus	Bitemporal and upper nasal	۲	L glaucoma
М	20	R –6.75 D sph., –0.25 D cyl., axis 90° L –2.5 D sph., –1.75 D cyl., axis 90°	6/9 6/12	Conus	Bitemporal	٢	Glaucoma
ř.	30	R -0.25 D cyl., axis 165° L +0.25 D sph., -0.5 D cyl., axis 15°	6/53 6/5	Conus	Upper bitemporal loss	12	Referred originally with conjunctivitis
M	52	R – 1 • 5 D sph., – 0• 75 D cyl., axis 180° L – 0• 75 D cyl., axis 90°	6/12 6/5	Conus Situs inversus	Bitemporal	4	Lens opacities
z	20	Not known	6/12 6/12	Conus Pseudo-papilloedema	Bitemporal	8	Lens opacities
۲u	20	R —0.75 D sph., —1.5 D cyl., axis 70° L —0.25 D sph., —2: D cyl., axis 110°	6/6 6/9 +2	Conus	Bitemporal	6	Suspected glaucoma, 1962
H	46	R -3°25 D sph., -2 D cyl., axis 115° L -1°25 D sph., -1°5 D cyl., axis 135°	6/6 6/9	Conus	Bitemporal	25	Routine refraction

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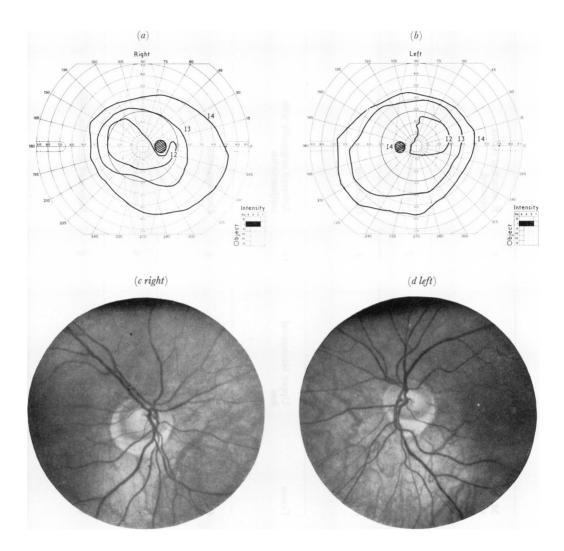


FIG. 3 (a, b) Case 6. Visual fields on February 9, 1971. (c, d) Case 6. Retinal appearances.

Two patients presented with retinal detachment, subsequently replaced: in one the affected eye did not show inversion or conus at the disc (Case 9); in the second the detached retina occurred in an eye with conus and inversion (Case 3). As the detachment in this case was an upper left nasal balloon, the field has been included in the results. Other reasons for seeking medical advice were:

Routine refraction	6	(Cases 2, 4, 6, 7, 8, 16)
Glaucoma	2	(Cases 10, 11)
Lens opacities	2	(Cases 13, 14)
Headache and visual loss	I	(Case 1)
Suspected glaucoma	I	(Case 15)
Headache	I	(Case 5)
Conjunctivitis	I	(Case 12)

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In 28 out of 29 eyes which showed conus at the optic disc depression of the field was found to a small stimulus. In one patient the eye with a normal disc showed the characteristic upper temporal defect. Most of the defects were found in the upper temporal quadrant, but some extended to the lower quadrant or across the midline into the upper nasal quadrant.

Retinoscopy in the meridian of the defect at about 20° to the optic axis invariably showed a slightly more myopic state than close to the fovea, suggestive of some degree of ectasia corresponding to the site of the field depression. Static perimetry across this meridian (Fig. 2c), with first the conventional correction and then with more myopic correction, only showed a very slight improvement in sensitivity. Peripheral fields were normal in all cases. All phakic eyes showing conus were associated with myopic astigmatism or compound myopic astigmatism.

Diminished retinal pigmentation was a marked feature in the lower nasal quadrants. This was well shown in Cases 3, 5, and 6, in which the fundus pallor extended close to the disc margins. The field defects in each case corresponded approximately to the pale area.

None of the patients was aware that there was any field defect present. No change in the field defect was noted in any case during follow-up periods which varied from 1 to 12 years.

Discussion

Szili (1883) showed that two phenomena were associated with inferior conus of the optic disc; the lower half of the fundus showed less pigmentation than the upper, and the error of refraction in the lower half was frequently greater than at the macula or temporal region. Rucker (1946) showed that temporal field defects occurred with situs inversus alone, and with conus alone. Fuchs (1882) emphasized the frequent association of situs inversus with inferior conus. Caccamise (1954), Berry (1963), and Odland (1967) described bitemporal field defects associated with inferior crescents, situs inversus, and thinning of the choroid. Schmidt (1955) showed that ectasias may give rise to field defects which disappear when higher myopic corrections are used. Maguire (1971) emphasized the value of static perimetry in this respect. That some degree of ectasia is present in the lower nasal quadrants of the retina in the present series is suggested by the higher myopic refraction in this area compared with the macula. All these cases were examined with correcting lenses. When higher minus lenses were used, a slight decrease of the field defects would seem, therefore, to be due to two possible factors:

(a) Some degree of ectasia of part of the globe. This is supported by the difference of the refraction in the affected area compared with that at the macula.

(b) Hypoplasia of the retina and choroid with fewer receptors per unit area. The decrease in pigmentation of the fundus in the lower nasal quadrants is advanced to support this theory.

In differentiating between bitemporal field defects due to ocular and chiasmal lesions, the behaviour of the field relative to the vertical meridian should be noted. Chiasmal lesions tend to respect the midline and end in a sharp edge, which in the early stages does not cross to the nasal side. Defects due to ocular lesions tend to slope obliquely across the midline and frequently involve part of the upper nasal quadrants.

Summary

Bitemporal visual field defects are described in a series of sixteen patients showing inferior conus of the optic disc with situs inversus. The nature of the field defect helps to distinguish an ocular from a chiasmal aetiology. The cause of the defect is postulated to be a combination of local retinal ectasia with hypoplasia.

References

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