Ischaemic optic neuropathy in pulseless disease

T. J. K. LEONARD and M. D. SANDERS

From the Department of Neuro-Ophthalmology, National Hospital for Nervous Diseases, Queen Square, London WC1

SUMMARY A diagnosis of pulseless disease was made in a young European female patient who presented with symptoms confined to the visual system. She demonstrated the classical ocular features of pulseless disease but developed the previously unrecorded complication of ischaemic optic neuropathy. These findings are discussed.

Ocular symptoms may be the presenting feature of occlusive disease in vessels arising from the aortic arch. Reduced perfusion to the upper part of the body leads to diminished or absent peripheral pulses, making the descriptive term 'pulseless disease' a suitable alternative to the many eponymous titles¹ for this condition.

In 1827 Robert Adams² recorded the first case of pulseless disease. Twelve years later an elegant description by Davy³ detailed the symptoms of confusion and syncope in a 55-year-old British Army officer. His illness was attributed to a chest wound sustained at the Battle of Waterloo. At necropsy a thoracic aortic aneurysm was found with involvement of the great vessels suggesting a diagnosis of syphilis. During life diminished pulsation at the neck and wrist had been noted.

In 1908 Takayasu⁴ was the first to describe the retinal complications, and at the same meeting Onishi and Kagashini related the ocular changes to the absence of radial pulses. Comprehensive accounts of the clinical features with pathological correlation followed, and an elaborate review by Pinkham⁵ in 1955 established visual symptoms in at least 70% of the patients. In 1976 Uyama and Asayama⁶ reclassified the retinal changes and found a correlation between microaneurysm formation and the systolic ophthalmodynamometry pressures.

We here report for the first time the association of ischaemic optic neuropathy with pulseless disease.

Case report

A 37-year-old Caucasian female presented to the National Hospital, Queen Square, under the care of Professor J. Marshall in October 1981 complaining of a 6-month history of transient attacks of blurred vision. The attacks were confined to her left eye and lasted for a few seconds at a time. She complained of a 'grey blind' crossing the visual field from left to right, with complete loss of vision followed by rapid resolution. She could not relate these attacks to any activity or time of day. Otherwise she was well, though complaining of general lethargy. The visual obscurations increased in frequency, and attacks could be precipitated by minimal exercise and relieved by rest. She had no relevant previous medical history.

On examination her visual acuity was 6/6 and N5 in each eye, with normal colour vision to Ishihara testing. Pupil reactions were sluggish to light and accommodation, but no relative afferent pupil defect was detected; perimetry with Bjerrum and Goldmann fields was normal. The intraocular pressures were low

![Fig. 1 Flourescein angiogram right eye. Early venous phase. Late filling of the choroid, retinal venous dilatation, and microaneurysms (arrow) can be seen. The retinal peripapillary capillary circulation is also dilated.](image)
slight venous dilatation, and a few microaneurysms in both eyes. Fluorescein angiography showed these microaneurysms and dilatation of the peripapillary capillaries with delayed choroidal filling (Fig. 1).

On general examination she was well, but there were absent radial pulses, an absent left carotid pulse, and a weak right carotid pulse. Pulses in the legs were normal. The blood pressure could be measured only in the legs and was 150/70 mmHg.

The only abnormal results of blood tests were a slight leucocytosis and an ESR of 35 mm/h. VDRL and TPHA tests for syphilis were negative, and the serum lipid profile was normal. Arch arteriography (Fig. 2) showed changes consistent with an occlusive arteriopathy of the vessels arising from the aortic arch. A diagnosis of pulseless disease was made.

Three weeks later she developed episodic faintness, general malaise, and a slight fever. Sudden onset of tunnel vision in the right eye precipitated admission. Her blood pressure had fallen to 90/70 mmHg and the pulse rate was 110/minute. She appeared confused and disorientated without localising neurological signs and felt faint on any change in posture. Visual acuity was 6/9 N5 in the right eye, but the left remained at 6/5 N5. The right visual field had become reduced to the central 20° isoptre, while the left showed mild generalised depression. A right relative afferent pupil defect was present. Intraocular pressures were reduced to 2 mmHg in each eye, but the colour vision, eye movements, and anterior segments were all normal.

Funduscopy showed right optic disc oedema, with pallid swelling at the upper and lower poles and small nerve fibre layer haemorrhages (Fig. 3). The appearances were consistent with an ischaemic optic neuropathy, with sparing of the papillo-macular nerve fibre bundle. Retinal arteries were now constricted and the veins dilated. Small punctate haemorrhages were scattered throughout the fundus, with a marked increase in the number of microaneurysms. The left disc was normal, but the retinal changes were similar to the right.

**SURGICAL TREATMENT**

Twenty-four hours after the patient’s admission reconstructive surgery was performed by Professor Norman Browe.- At operation the left common carotid was seen to be obliterated. On the right the innominate artery was patent, leading to an attenuated common carotid, which gave rise to a narrowed internal carotid artery serving as the only main vessel to the head, while the external carotid was fibrosed. An extensive collateral circulation had replaced the subclavian and vertebral arteries. A Dacron graft was placed between the ascending aorta and the right internal carotid.
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Fig. 3 Fundus photograph of the right optic disc 24 hours after reconstructive surgery. Optic disc infarction is visible sparing the papillomacular fibres. Postoperative haemorrhages are present at all retinal levels with extension to the subhyaloid space. The extensive haemorrhages are due to a sudden increase in perfusion pressure through a chronically dilated retinal circulation.

Her general condition dramatically improved after operation. The visual acuity remained at 6/9 N5 right and 6/5 N5 left. The left visual field appeared less depressed but the right remained as an asymmetric tunnel-like field due to large upper and lower arcuate loss. The intraocular pressure had risen to 10 mmHg in each eye, and ophthalmodynamometry readings were 50/35 right and 30/25 left. A good right carotid pulse was palpable.

There was marked haemorrhagic retinopathy in the right eye, with haemorrhages at all retinal levels extending into the subhyaloid space (Fig. 3). The left fundus showed only a few new retinal haemorrhages.

Figure 4 Histological section of right internal carotid artery wall. Disorganisation of the media and elastic tissue can be seen with fibrosis (arrow). Lymphocytic infiltration was present without giant cells. (Elastin van Giesen, ×22).

Pathological study
Histological examination of a biopsy taken from the right common carotid showed a focal lymphocytic infiltrate in the media with subintimal fibrosis (Fig. 4). Destruction of elastic tissue was visible, but there were no giant cells. There was no evidence of atheroma or syphilis.

Discussion
The symptoms and signs of pulseless disease are due to cerebral and ocular hypoperfusion caused by gradual obliteration of the great vessels arising from the aortic arch. The aetiology may be an atheroma, syphilis, or a chronic nonspecific arteritis.

Cerebral Features
The cerebral symptoms include faintness, confusion, and psychiatric disturbance, while the signs are of a transient focal or general neurological disturbance including hemiplegia and dysphasia.

Ocular Features
Pulseless disease produces changes in the retina, choroid, and anterior segment together with marked lowering of the intraocular pressure.

The retinal changes have been well documented by Uyama and Asayama and other authors. The chronic nature of the disease produces progressive dilatation...
of the retinal capillaries leading to the formation of fusiform or saccular microaneurysms. The capillaries show a loss of endothelial cells and preservation of mural cells. Exudate and haemorrhage are rare in pulseless disease, since the intraluminal pressure in these dilated vessels is very low. Progressive retinal hypoxia causes arteriovenous communications to form. These begin in the mid peripheral retina and then progress to surround the disc in a whorl-like arrangement as detailed by Takayasu. Choroidal hypoperfusion, as shown by fluorescein angiography, leads to pathological changes in the outer retinal layers, with loss of receptor cell nuclei.

The intraocular pressure is very low in most cases of pulseless disease. Factors include hypoperfusion of the ciliary body with reduced aqueous secretion, and less aqueous is produced by ultrafiltration. The low intraocular pressure may in part be a homoeostatic response to reduce resistance to flow in the intraocular circulation, which would help to protect the eye from vascular occlusion.

Changes in the anterior segment also occur as a result of ocular hypoxia. Dilated tortuous conjunctival vessels are seen, and eventually the eye becomes painful, with corneal clouding, ruberosis iridis, cataract, and an ischaemic uveitis. The case reported here had the classical early ophthalmological features of pulseless disease with retinal microaneurysms, choroidal hypoperfusion on fluorescein angiography, and marked conjunctival vessel tortuosity in both eyes.

The exceptional feature of this case was the ischaemic optic neuropathy due to a sudden reduction in perfusion pressure to the optic nerve head. Failure of perfusion pressure causing ischaemic optic neuropathy is occasionally seen in occlusive carotid artery disease or due to a sudden drop in systemic blood pressure (e.g., haemorrhage, surgical hypotension, severe cardiac failure).

Although optic atrophy has frequently been described in pulseless disease, acute ischaemic optic neuropathy has not. The chronic nature of the disease allows a collateral circulation to evolve, and despite choroidal hypoperfusion the compensatory changes ensure an adequate blood supply to the optic nerve head. The extremely low intraocular pressures in pulseless disease create very little resistance to perfusion, thus protecting the optic nerve head.

This case showed marked asymmetry in the circulation to the 2 eyes. The right eye was supplied by an ophthalmic artery of good calibre allowing higher ophthalmodynamometry readings and a higher intraocular pressure than its fellow. When obliteration of the only main patent vessel to the head caused ocular perfusion to drop to a critical level, the right eye was unable to compensate and suffered an ischaemic optic neuropathy, while the left, supplied by an adequate collateral circulation, survived.

In conclusion, this case illustrates an unusual cause of acute visual loss in a disease that is exceedingly rare in Europeans. Reconstructive surgery, however, has been very successful in restoring the patient’s general health and maintaining her present level of vision.

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T J Leonard and M D Sanders

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