Other Optic Nerve Conditions

Dr Simon Barnard
Phd FCOptom FAAO DipCLP DipClinOptom DipTh(IP)

Contents
Other Optic Nerve Conditions ................................................................. 1
Introduction ............................................................................................. 2
  The nerve .............................................................................................. 2
  Blood supply .......................................................................................... 2
General clinical features of disorders of the optic nerve. ......................... 2
Classification of diseases affecting the optic nerve ................................. 3
  Developmental anomalies .................................................................... 3
    Aplasia & hypoplasia. .......................................................................... 3
    Colobomas.......................................................................................... 3
    Morning glory syndrome ................................................................. 3
    Optic nerve pit.................................................................................... 4
  Myelinated nerve fibres ................................................................. 4
  Retarded myelination ........................................................................ 4
  Glial remnants .................................................................................... 4
  Congenital swelling of the optic disc (pseudo - papilloedema) ........ .... 5
  Congenital and myopic crescents (conus) ........................................ 5
  Tilted disc........................................................................................... 6
Vascular anomalies ................................................................................... 6
  Aneurysms & subarachnoid haemorrhage ........................................ 6
  Ischaemia & infarction of the nerve tissue ......................................... 6
  Toxic including iatrogenic and nutritional disorders .......................... 7
  Tumours & cysts ............................................................................... 7
  Drusen of the Optic Disc .................................................................. 7
  Trauma ............................................................................................... 8
  Optic Atrophy .................................................................................... 8
  "Choked disc" ................................................................................... 8
Example referral letter ............................................................................. 9
Further reading ....................................................................................... 9
Introduction

The nerve

The O.N. is a tract consisting mainly of the axons of the ganglion cells of the retina. These axons converge on the optic disc, which is approximately 1.5mm in diameter, pierce the sclera at the lamina cribosa, a sieve-like structure, then form bundles of myelinated nerve fibres separated by connective tissue septa. Largely because of the presence of the myelin sheaths and the connective tissue septa behind the level of the lamina cribosa, the optic nerve has a greater diameter at the point at which it leaves the globe than at its head (the optic disc).

Each optic nerve is encased in sheaths continuous with and similar to the meninges of the cranium (pia, arachnoid, and the dura).

The optic nerve may be considered as consisting of four parts:-
(1) the head & ocular portion which traverses the sclera
(2) the orbital portion, which is about 3cm in length and has an S-shaped course so that the globe is able to move freely
(3) the portion in the optic canal, which is some 5-7mms long
(4) an intracranial portion which extends from the optic canal to the anterior part of the optic chiasm

Blood supply

The arterial supply to the optic nerve anterior to the lamina cribosa is derived from the short ciliary arteries. Immediately behind the lamina cribosa vessels derived from the Circle of Zinn, which is itself supplied by the short ciliary arteries, enter the optic nerve. The orbital portion of the optic nerve derives its blood supply from the pial circulation and perhaps also to some extent from the ophthalmic artery and its branches, including the central retinal artery. That portion of the optic nerve lying in the optic canal derives its arterial blood supply from the ophthalmic artery, whilst the intra-cranial part of the optic nerve is supplied centripetally through the pial vessels. Venous drainage from the ocular and orbital portions of the optic nerve is chiefly into the central retinal vein.

General clinical features of disorders of the optic nerve.

Unilateral visual failure unassociated with ophthalmoscopic signs may be due to optic nerve disease. However when the cause is due to toxic or systemic disease, the visual loss may be bilateral.

Consider visual acuities, contrast sensitivity, colour vision, visual fields and pupil reactions. If visual failure is complete the direct pupil response to light will be lost and the pupil may be dilated. If visual loss is less severe, the pupil
may show loss of sustained direct response to light stimulus and a rhythmical contraction of abnormal amplitude (hippus).

**Classification of diseases affecting the optic nerve**

May be based on aetiologic factors and/or clinical appearance:-

Developmental anomalies  
Vascular disorders  
Glaucoma  
Demyelinating and inflammatory disease  
Toxic including iatrogenic disorders  
Tumours & cysts  
Drusen  
Trauma

**Developmental anomalies**

Aplasia & hypoplasia.

Aplasia (rare) and hypoplasia is a failure of the optic nerve to develop. Hypoplasia is bilateral in about 60% of cases. May be a sporadic developmental anomaly or hereditary. Small grey disc often surrounded by yellow halo of hypopigmentation due to choroidal and RPE abnormality, reduced V.A. and a variety of visual field defects.

**Colobomas**

Incomplete closure of foetal cleft thus appearing inferiorly. May be associated with choroidal/iris coloboma. Field defect corresponds to the area of projection of the missing fibres. Visual acuity may be affected.

**Morning glory syndrome**

Rare dysplastic coloboma of optic disc. ON head enlarged and cupped with hyaloid remnants within its base. Usually unilateral. Vision grossly impaired.
Optic nerve pit

Not related to coloboma. Rare (1 per 11,000). Bilateral in 15% of cases. Occasionally more than one hole. Usually positioned inferotemporal quadrant of papilla. Size varies from 0.1 to 0.7 disc diameter (average 0.3). Depth varies from 0.5D to 20D. Usually grey in colour. Visual field defects of various types. Cause not certain. Possible sequel is serous retinal detachment.

Myelinated nerve fibres

Myelination of the optic nerve develops embryologically from a posterior to anterior direction. The fibres are grossly myelinated in the intracranial and intracanal portions of the optic nerve by the seventh month but myelin does not usually reach the lamina cribosa until full term. In some instances the myelin does not stop short at the lamina cribosa but continues into the eye, appearing on ophthalmoscopic examination as bright white flame-shaped streaks, usually contiguous with the margin of the optic disc. These streaks follow the normal course of the retinal nerve fibres and are situated superficially on the disc and retina and hence may cover some of the retinal blood vessels. They give rise to field defects corresponding to their area of projection into the visual field.

Retarded myelination

A number of observers have reported the appearance of the optic disc as being grey in premature and some full-term infants. Possible cause of strabismus?

Glial remnants

The posterior portion of the hyaloid artery may persist extending into the vitreous from the disc surrounded by glial tissue (Bergmeister's papilla). Sometimes sheets of persistent glial tissue are seen covering the optic disc in part or as a whole and producing a slightly grey filmy (gossamer) appearance.
**Congenital swelling of the optic disc (pseudo - papilloedema)**

This is a physiological variation from the norm seen fairly frequently in patients with high **axial hypermetropia**.

Margins of disc appear blurred and disc tissue somewhat elevated. Major points in favour of diagnosis of pseudo-papilloedema as opposed to pathological "choked-disc" include the absence of exudates or haemorrhages and a normal visual field with a **small-sized** physiological blind spot. Fluorescein angiogram will be normal.

A tilted disc because the nerve enters the back of the eye obliquely can give a sectorial crowding.

The eyes are often small (i.e., hypermetropic) so less room for the nerve fibres which are therefore crowded into a small space. The cup will often be small.

Buried drusen will also or due a crowded appearance. Disc drusen are composed of small spheres of protein and mucopolysaccharides. material that become calcified with advancing age.

Pseudopapilledema may be caused by hyaloid remnants and glial tissue.

**Congenital and myopic crescents (conus)**

The ophthalmoscopic appearance of the margin of the optic disc is determined to a large extent by the distances from the disc margin at which the RPE, Bruch's membrane and the choroid terminate. In axial myopes these tissues are often absent over a crescent-shaped area on the temporal side of the disc (myopic crescent). This is probably due to chorioretinal atrophy and/or traction of the retina and choroid on the nasal side due to elongation of the globe at the posterior pole.
Tilted disc

Occasionally the optic nerve leaves the sclera more obliquely than usual giving rise to an inferior or superior conus. Visual field anomalies may be present.

Vascular anomalies

Aneurysms & subarachnoid haemorrhage

Aneurysms involving ON may arise from internal carotid, anterior cerebral, anterior communicating, and occasionally from ophthalmic artery. Because aneurysms exert their effect by slow growth they may give rise to the same symptoms as tumours i.e. compression $\Rightarrow$ visual field defects and primary optic atrophy. When they bleed they give rise to the same symptoms as subarachnoid haemorrhages. Symptoms suggestive of an aneurysm are those of sudden onset which vary from day to day. Apart from an aneurysm, subarachnoid haemorrhage may result from trauma, tumours, the breakthrough of an intracerebral haemorrhage, or spontaneously. Always check that the limiting edge of a subconjunctival haemorrhage is visible.

Ischaemia & infarction of the nerve tissue

Oclusive disease (either inflammatory or degenerative in origin) of the arteries supplying the optic nerve is a frequent cause of damage. Aetiologies
include giant cell arteritis, CRA occlusion by embolism or thrombus, thrombosis of CRV, and occlusion of ophthalmic artery or internal carotid, most commonly by atheromatous plaque formation.

**Temporal arteritis** may be accompanied by giant cell arteritis of posterior ciliary arteries resulting in ischaemic optic neuropathy. Giant cell arteritis usually affects patients over 50 years of age. Where it includes temporal arteritis the symptoms may include unilateral headache and tenderness of the temple. At first there may be dimming of the vision followed by complete amaurosis.

Other inflammatory diseases affecting the arteries supplying the optic nerve include *polyarteritis nodosa, thromboangiitis obliterans, optic disc vasculitis* and *syphilitic meningitis*. *Ischaemic optic neuropathy* is not always the result of giant cell arteritis (e.g. it may be caused by arteriosclerotic occlusive disorders of the posterior ciliary arteries.

**Toxic including iatrogenic and nutritional disorders**

Many substances can affect the function of the optic nerve and may give rise visual field defects and colour vision anomalies.

Examples include: Tobacco, ethyl and methyl alcohol; sedative drugs such as morphia and phenyprazine, anti-infective drugs such as chloramphenicol, and other drugs such as digoxin, thyroxin, nicotinic acid, quinine derivatives and salicylic acid; metals such as lead, thallium and arsenic.

Optic neuritis of nutritional origin include beri-beri and pernicious anaemia.

**Tumours & cysts**

Intrinsic - e.g. gliomas and meningiomas originating from nerve tissue
Extrinsic  e.g. meningiomas of sphenoidal ridge or olfactory groove, pituitary adenomas and some metastatic tumours. Tumours in the orbit may interfere with ocular motility and cause proptosis and additionally interfere with vision if they affect the ON conductive system

**Drusen of the Optic Disc**

This degenerative condition consists of a build up of spheres of hyaline material in front of the lamina cribosa. As they enlarge they give rise to an elevation of the optic disc and a blurring of the margins. When they are situated superficially the ophthalmoscopic appearance is of white or white-yellow spheres which have a sheen or appear translucent if illuminated from one side. If they are deep they may not be observable with the ophthalmoscope and cause a misdiagnosis of papilloedema. Can be progressive. Visual field defects. Usually bilateral.
Trauma

A small percentage of head injuries can produce damage to the optic nerve which do not have the Schwann sheaths necessary for effective regeneration of the nerves.

Optic Atrophy

Observed clinically as pallor of the disc

**Primary** - disc margins are distinct  
**Secondary** - disc margins indistinct - such cases result from previous oedema or inflammation of the nerve head  
**Consecutive** (or ascending) optic atrophy is that which occurs secondary to retinal disease in which ganglion cells have been destroyed e.g. CRA occlusion  
**Descending** optic atrophy may result from lesions affecting the orbital, optic canal portion, or intracranial portion of ON

"Choked disc"

May be due to papilloedema or papillitis. Ophthalmoscopic appearance indistinguishable.

Papilloedema is a non-inflammatory swelling of optic nerve head with abnormal elevation of disc and blurring of margins. As condition progresses haemorrhages (usually flame shaped) may appear on the nerve head and surrounds. The oedema may spread into the surrounding retina causing concentric folds. There is a loss of spontaneous venous pulsation. Commonest causes include Grade 4 vascular hypertension and raised intracranial pressure.

In all but the late stages central vision and colour vision unaffected. Enlarged blind spot.

Papillitis is an active inflammation of nerve head (an optic neuritis). Central vision affected including field loss, colour vision disturbances, and reduced vision. Pain on movement of eye. Possible causes include MS.
Example referral letter

Professor Chris Bentley  
2nd Floor  
The Wellington Hospital (South)  
Wellington Place  
London NW8 9LE  

29th October 2012  

Dear Chris  

Re: Miss LS (age 12)  
London HA  

I should be grateful for your opinion on L who presented reporting that her eyes and head have been hurting intermittently, about every three weeks, mostly right sided for a number of months. She had been seen by my colleague MS for a routine examination on 3rd September 2012 and did not appear to have mentioned this at that time. Last night she had experienced pain in her left eye which woke her up and earlier this morning the pain was quite bad. On questioning she described the pain as being sharp. There is no associated nausea or visual disturbance. Her general health is good, she takes occasional Tynelol and there is no family history of ocular disease.

Refraction gave R -3.00/-1.25 x 5 = 6/75 L -2.75/-1.00 x 5 = 6/9-1. I also confirmed this under cycloplegia. This is very similar to the spectacles prescribed by Menachem Salasnik in September.

External eyes, pupillary reflexes, ocular motility and ocular motor balance are normal. Anterior chamber angles are wide and quiet. The ocular media are clear. The ocular fundi appear normal but I did note bilateral mild crowding of the nasal optic nerve heads which I have explained to Mrs S is probably physiological. This was noted by M in September. Previous records going back to 2009 only alluded to small cup disc ratios.

In view of the symptoms, which I do not believe have a visual cause, I have suggested to Mrs S that a routine ophthalmological opinion would be wise.

With kindest regards  

Dr Simon Barnard PhD FCOptom FAAO DipCLP DipClinOptom DipTh(IP)  

Cc Dr MW  
Mrs S  

Further reading  

Alexander L. (1995) Primary Care of the Posterior Segment. Appleton & Lang, Norwalk  