REVIEWS

VISUAL DISTURBANCES – A BORDERLINE BETWEEN OPHTHALMOLOGY AND NEUROLOGY

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ABSTRACT

Visual acuity loss can be due to eye ball diseases or to lesions of the optic nerves, visual pathway or cortical projection of vision. The differential diagnosis can be challenging and though patients go first to the ophthalmologist, there is not always the ophthalmologist who can diagnose and treat the patient. For a great number of causes/diseases, a neurologist should also see the patient, even if there is a transient or persistent monocular or binocular visual acuity loss a disturbance of the visual field or of the perception of colours. Visual acuity loss can be more than this. It can be a symptom or a sign of a neurological or systemic disease and a rapid diagnosis and specific treatment are mandatory in order to treat the symptoms and improve the patient’s quality of life.

Keywords: visual acuity loss, retinopathies, optic neuritis, blindness, visual field disturbances

INTRODUCTION

The visual system assures the most important integration of the body in the surrounding world, and this is proven by its cortical projections and connections with the majority of the areas into the central nervous system. A visual control is needed for movement, coordination, praxia, reading, writing, playing an instrument, painting (recognising colours, the form of the objects), driving a car, spatial orientation, executive functions and even for the behaviour.

The visual system is represented by the eyeball, the optic nerve and the visual pathway with its connections and the cortical projection. The extrinsic muscles of the eyeball innervated by the cranial nerves (nerves III, IV and VI – comune oculomotor, trochlear and abducens) assure a clear binocular vision. The pathological lesions affecting the eyeball (and there are not few) are diagnosed and treated by the ophthalmologist. Lesions affecting any other part of this pathway can be subject of a team-work or a problem to be solved by the neurologist, as well as diplopia while using both eyes (monocular diplopia beeing the subject of an ophthalmologic examination).

Functional visual loss (a decrease in visual acuity and/or visual field) is usually sending the patient first to the ophthalmologist but, if lenses do not correct the low vision, if the intraocular pressure is not high enough to explain concentric loss of the visual field or if no other ocular organic causes can explain the pathological changes, then, the patient has to be referred to a neurologist, as the cause is not an ophthalmological one, but can be part of a neurological or a systemic disease (1).

Visual acuity loss is a frequent complaint. Children and teenagers can be diagnosed with traumatic or non-traumatic conditions, as congenital disturbances, amblyopia, myopia or hypermetropia, retinitis pigmentosa, prematurity retinopathy, hypoplasia of the optic nerve, tumours of the optic nerve or suprasellar tumours. In adults, ophthalmologists are used to diagnose cataract, glaucoma, diabetic retinopathy or macular degenerescence, but vascular causes (as the occlusion of the central retinal artery or vein) or inflammatory and demyelinating lesions of the optic nerve are not rare at all (1).

The visual acuity changes can be caused by refractive errors, opacification of the transparent media of the eye (cornea, aqueous humour, crystalline...
lens, vitreous humour), by lesions of the retina (detachment, inflammation, infection, vascular occlusion, vasculitis, tumours) or by lesions of the optic nerve or optic pathways and can be monocular or binocular, with an acute or gradual onset, episodic or permanent.

OVERVIEW OF DISEASES

Transient monocular blindness has most frequent a vascular cause, and is known as amaurosis fugax. There is an acute loss of vision that can be partial or complete, lasting for some seconds to minutes. The most frequent causes are the stenosis of the internal carotid artery with secondary retinal/choroidal hypoperfusion, giant cells arteritis, dissection of the internal carotid artery. In this case, the patient has to be referred as soon as possible to a neurologist for further investigation, as this can be a sign of a severe vascular systemic disease, announcing a future stroke (2).

Another cause of episodes of monocular transient blindness, with duration of seconds, precipitated by postural changes or Valsalva manoeuvres is the optic disc edema (papilledema). The papilledema can be bilaterally, asymmetric, secondary to intracranial hypertension, even if the visual loss is transient monocular, and excepting these episodes, the visual acuity is normal. The optic disc is elevated, hyperaemic, imprecisely delimited and the veins are turgescent. The visual field can be normal, the blind spot can be larger or it can be concentric reduced. The patient should be investigated by cerebral Magnetic Resonance Imaging (MRI) in order to diagnose tumours, hydrocephalus, cerebral sinus vein thrombosis. If this imaging test shows no pathological findings, an infectious pathology has to be excluded by examination of the cerebrospinal fluid (meningitis, encephalitis) (2).

Other causes of transient monocular blindness are: repeated angle closure glaucoma attacks, discal abnormalities (drusen), coloboma iridis, orbital tumours. In the demyelinating optic neuropathies, the Uhthoff phenomenon is described, with transient visual loss during physical exercises or exposure to excessive heat (3).

Cause of transient binocular blindness are described in Table 1.

Monocular persistent visual acuity loss can have an ocular cause or can be related to the optic nerves, extending to the junction with the optic chiasm (optic neuropathies) (2).

The most frequent causes of ocular disturbances are presented in Table 2.

The borderline between ophthalmology and neurology is represented by the differential diagnosis between macular and optic nerve disturbances (Table 3).

The most frequent retina diseases are:

1. Serous retinopathy – leakage of serous fluid in the sub-retinal space, causing detachments of the retinal layers. It mostly affects males in the decades 5 and 6 of age, is considered idiopathic, but can be exacerbated by stress or use of corticosteroids. The onset is quite acute, not painful, with central vision loss and metamorphopsia. The optic disc is normal and ocular computed tomography (OCT) is showing retinal changes. The symptoms may disappear within 1 to 6 months without treatment, but sometimes intraocular injections with Avastin or LASER treatment are required to seal the detached retinal areas (1);

2. Age related macular degeneration (AMD) – affects both eyes and is progressive. The drusen (white-yellowish deposits with unclear limits) is an early sign and it is associated with hypo- or hyperpigmentation of the retinal pigment epithelium. The central vision is initially distorted and then visual acuity is lost progressively. Two types are described: the wet and the dry type (1);

3. Retinal detachments – central visual acuity loss and afferent pupillary defect if the central part of the retina is affected. Photopsia can frequently

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**TABLE 1. Causes of binocular transient blindness (2)**

<table>
<thead>
<tr>
<th>Migraine aura</th>
<th>Scintillant scotoma which can progress to complete congruous hemianopia Followed or not by pulsatile (throbbing) headache</th>
</tr>
</thead>
<tbody>
<tr>
<td>Transient ischemic attack in the basilar or posterior cerebral arteries</td>
<td>Older patients Acute onset If present, headache is concomitantly with the visual field disturbance</td>
</tr>
<tr>
<td>Edema of the optic disc</td>
<td>Causes less frequent</td>
</tr>
<tr>
<td>Occipital focal attacks</td>
<td></td>
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<tr>
<td>Head Trauma</td>
<td></td>
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<tr>
<td>Posterior reversible encephalopathy (lasting hours-days)</td>
<td>Hyptensive encephalopathy Eclampsia/pre-eclampsia Toxic - cyclosporine</td>
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</tbody>
</table>
precede the detachments. The risk factors are severe myopia, recent intraocular surgery, eye trauma, family history of retinal detachments (2);

4. Central retinal artery occlusion (CRAO) – suddenly, not painful, onset of unilateral blindness (if the cause is a dissection of the internal carotid artery or an arteritis with giant cells). The retina is white, opaque, edematized and sometimes the thrombus can be seen (“cherry red spot macula”). Sometimes, the local intra-arterial thrombolysis can have good therapeutic results (4);

5. Central retinal vein occlusion (CRVO) – subacute monocular variable loss of visual acuity that sometimes can spontaneously recover. Retinal veins are dilated, tortuous and cotton-wool spots, macular edema and optic disc edema may be associated to extensive haemorrhages in the posterior pole giving the “blood and thunder appearance” (4).

The retinal vessel occlusions are affecting the layer of ganglionic cells and this can generate afferent pupillary defect;

6. Diabetic Retinopathy – is a frequent cause of blindness; it is also common in patients with stroke. The retinal ischemia is diffuse, including the macula, which leads to central vision loss. Secondary to ischemia, there is a process of neovascularisation that will eventually be responsible for haemorrhages and retinal detachments. The treatment can be done using laser photocoagulation or intravitreal injections with anti-VEGF agents, but also by maintaining glycaemic control and reducing the cardiovascular risk factors (5);

7. Hypertensive retinopathy – is usually the result of chronic arterial hypertension; it is usually asymptomatic, but the retinal damage may produce vision loss. The retinal vascular changes are usually bilateral, but asymmetric, and include arterial narrowing, irregularity of the vessels, microaneurysms, cotton wool spots, arteriovenous nicking. However, in patients with acute hypertensive retinopathy, there are retinal haemorrhages, macular edema, retinal vessels spasms, cotton wool spots, optic nerve edema. Sometimes, there is isolated op-

<table>
<thead>
<tr>
<th>TABLE 2. Most frequent disturbances of the transparent media of the eye (1)</th>
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<tbody>
<tr>
<td><strong>Cornea</strong></td>
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<tr>
<td><strong>Anterior chamber</strong></td>
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<tr>
<td><strong>Crystalline</strong></td>
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<td><strong>Vitreous body</strong></td>
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<table>
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<tr>
<th>TABLE 3. Differential diagnosis between macular disturbances and optic neuropathy (2)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Macular disturbances</strong></td>
</tr>
<tr>
<td>Central vision loss</td>
</tr>
<tr>
<td>Central scotoma</td>
</tr>
<tr>
<td>Colour vision disturbances</td>
</tr>
<tr>
<td>Pain</td>
</tr>
<tr>
<td>Distorted vision</td>
</tr>
<tr>
<td>Afferent pupillary defect</td>
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</tbody>
</table>
tic nerve edema, thus it’s important to assess arteri-
mal blood pressure in patients with headache and bi-
lateral optic nerve edema. There can be choroidal
ischemia, ischemic optic neuropathy which can
lead to permanent vision loss. In such cases, intra
vitreal injection with bevacizumab is helpful, im-
proving the visual outcome (6).

Unilateral optic nerve neuropathies – can be
generated by inflammation, ischemia or compres-
sion:

A. Inflammatory optic neuropathies

1. Demyelinating – are more frequent in young
adults, especially women. The demyelinating optic
retrobulbar neuritis (NORB) is rapidly progressive,
unilaterally, (sometimes bilaterally, during the
same episode or most frequent at some intervals of
time) and can be inconstantly associated with orbit-
al pain induced by eye movement or eye compres-
sion. The Uhthoff phenomenon representing the
increase of visual loss due to physical exercise or
exposure to heat is characteristic. The afferent pu-
illary defect is present. The fundoscopic eye exam
shows a normal optic disc and only in the case of an
associated papilitis the disc shows edema (7). The
vision prognostic is good, spontaneous remission
can be seen in some weeks to months. Intravenous
pulse-therapy with Solumedrol may help recovery.
NORB can be the first clinical sign of multiple scle-
rosis, that’s why these patients are obliged to be
seen by a neurologist and complete the investiga-
tions with a cerebral and spinal cord MRI, in order
to look for other demyelinating lesions of the cen-
tral nervous system and with a cerebrospinal fluid
examination to look for oligoclonal bands and cal-
culate the index immunoglobulins/albumin. De-
spite the visual loss recovery, in most of the pa-
tients, some degree of optic atrophy rests and the
optic disc is pale in its temporal region. The Visual
evoked potentials are a sensitive tool to diagnose a
NORB, even if it is asymptomatic (7). In the cases
when the optic neuritis is recurrent, bilaterally and
is associated with transverse myelitis and the visual
loss is not recovered during a month, the diagnosis
of optic neuromyelitis (Devic) must be taken into
account, having a poorer prognostic than the optic
neuritis in multiple sclerosis. Optic neuromyelitis
can also be seen in systemic infectious diseases (vi-
ral hepatitis, tuberculosis, syphilis, borreliosis) and
in autoimmune diseases (systemic erythematous
lupus, myasthenia gravis, autoimmune thyroiditis).
Antibodies anti-aquaporin 4 can be found in the se-
rum of up to 80% of the patients with optic neuro-
myelitis (8).

2. Autoimmune – recurrent episodes of optic
neuritis responsive to corticosteroid therapy, most
frequently in women. During the attack, the optic
nerve disc is slightly edematiated. If this disease is
not properly diagnosed from the onset of symptoms
it can evolve to optic nerve atrophy and severe
visual loss (9).

3. Infectious

a. Neuroretinitis – is affecting young adults
who can recall a viral prodrome. The visual
loss can be variable and may associate affer-
ent pupillary defect. The patients complain
of central or centrocecal scotoma and at fun-
doscopy they present optic disc edema (that
can last 3 months) and stellate maculopathy
(that can last 1 year), sometimes associating
peripapillary retinal detachments (10). The
recovery of visual loss normally starts some
weeks after the onset of the disease, and can
be incomplete if there are recurrent episodes
leaving permanent sequellae (11). A frequent
cause of neuroretinitis is the infection with
*Bartonella henselae* (12);

b. Syphilis – caused by *Treponema pallidum*,
is a disease with sexual transmission or can be
transmitted from mother to her foetus. The
optic nerve is affected in the late phases (sec-
ondary or tertiary) of syphilis, called neu-
rosyphilis. The patients complain of central
visual loss and dyschromatopsia. The optic
disc at fundoscopy can have a normal ap-
pearance, can show atrophy or oedema and
may associate stellate maculopathy, charac-
teristic for neuroretinitis. Cerebral MRI may
be normal, but cerebrospinal fluid analysis
using treponemal and non-treponemal tests
can establish the diagnosis. Penicillin is the
specific treatment and sometimes corticoster-
oids may be associated (13);

c. *Borrelia burgdorferi* – the optic neuropathy
is usually characterised by painless visual
loss, bilateral papilitis and central scotoma
(14). Improvement is possible with specific
antibiotherapy (15);

d. *Mycobacterium tuberculosis* – the optic in-
volvement may be due to tuberculomas com-
pressing the optic chiasm or causing opti-
co-chiasmatic arachnoiditis with secondary
optic nerve inflammation. There may be pain,
the vision loss may be gradual or rapid and
the optic disc is normal, edematous, or
atrophic. The diagnosis is confirmed by bac-
teria detection. A new diagnostic technique is
the nested PCR assay. The treatment includes
TABLE 4. Optic nerve neuropathies associated to systemic diseases

<table>
<thead>
<tr>
<th>Syndrome</th>
<th>Symptoms and Associated Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sarcoidosis</td>
<td>Systemic granulomatous inflammation, recurrent visual disturbances, optic nerve disc is normal,</td>
</tr>
<tr>
<td></td>
<td>atrophic or oedematous, central scotoma, altitudinal defect of the visual field, homonymous hemianopia,</td>
</tr>
<tr>
<td>Wegener granulomatosis</td>
<td>Patients 40-50 years of age, male, BUT optic neuritis is more frequent in women. Orbital signs are</td>
</tr>
<tr>
<td></td>
<td>associated – proptosis, pain, disturbances in oculomotricity, lung and kidney disturbances, c-ANCA</td>
</tr>
<tr>
<td></td>
<td>present in the serum. Treatment – corticotherapy in big doses, cyclophosphamide, rituximab</td>
</tr>
<tr>
<td>Sjögren syndrome</td>
<td>Keratoconjunctivitis, xerostomia, Sicca syndrome, associated optic neuromielitis</td>
</tr>
<tr>
<td>Systemic erythematous lupus</td>
<td>Many other organs are affected, not painful, subacute, severe visual loss, Antibodies anti ds-DNA,</td>
</tr>
<tr>
<td></td>
<td>anti-Smith, antiphospholipid, corticotherapy, cyclophosphamide, cyclosporine, methotrexate, azathioprine</td>
</tr>
<tr>
<td>Paraneoplastic</td>
<td>Associated with small cell lung cancer, B cell lymphoma, prostate cancer, renal carcinoma</td>
</tr>
</tbody>
</table>

4. Optic nerve neuropathies associated to systemic diseases.

B. Acute ischemic optic neuropathy – diagnosed in patients over 50 years of age. It has an acute onset, without pain and progresses in some hours or days. Two types are described: anterior (90% of cases), with oedematous optic nerve disc, and posterior (arteritic), with normal optic nerve disc. There is also a classification of the ischemic optic neuropathies taking into account the presence of arteritis with giant cells. The risk factors are blood pressure drop during heart surgery, systemic haemorrhage, renal dialysis, severe anaemia, hypercoagulability, radiotherapy. The treatment of the non-arteritic form is based on the control of the risk factors and exclusion of the presence of arteritis with giant cells. The arteritic form, present in older patients, associates masticatory claudication, myalgia, weight loss, headache. Laboratory tests will show an inflammatory syndrome with high values of the protein C and the biopsy of the temporal artery that will establish the diagnosis. Corticotherapy consists of metilprednisolone iv 1g/day 3-5 days followed by prednisone 1mg/kg/day for with progressive lowering of the dose is helpful (4).

C. Compressive/infiltrative optic neuropathy – meningioma in adults, glioma in children. The onset is progressive, not painful, colour vision is affected and can be associated with other malignancies or neurofibromatosis (2).

Persistent binocular visual acuity loss is caused by lesions of the retina in both eyes, of both optic nerves, optic chiasma, retrochiasmatic visual pathways (Table 5).

TABLE 5. Causes of persistent binocular visual acuity loss (2), (18)

<table>
<thead>
<tr>
<th>Causes and Pathways</th>
<th>Bilaterally optic neuropathies</th>
<th>Retrochiasmatic visual pathway and cortical lesions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Retinopathies to vitamin A deficiency</td>
<td>Toxic-nutritional (bilaterally, symmetrically, progressive, centrocecal scotoma) (18)</td>
<td>Neurological signs and symptoms associated</td>
</tr>
<tr>
<td>Degenerative retinopathy affecting the cone cells (cones) (tested by electroretinogram ERG)</td>
<td>Hereditary (bilaterally, progressive, centrocecal scotoma, optic atrophy)</td>
<td>Bitemporal hemianopia due to compression on the optic chiasma (Hyphophysial adenoma, meningioma, craniopharyngioma, aneurysms) – cerebral MRI is necessary for the diagnosis</td>
</tr>
<tr>
<td>Toxic retinopathies</td>
<td>Papillar edema secondary to intracranial hypertension</td>
<td>Homonymous hemianopia (affecting the contralateral optic tract pathway, corpus geniculatum laterale, optic radiations or the visual cortex)</td>
</tr>
<tr>
<td>Retinopathies associated to carcinoma</td>
<td>Open angle glaucoma (peripheral reduction of the visual field, high intraocular pressure, optic atrophy)</td>
<td>Cortical blindness (damage of occipital cortex with normal eye – known as Anton Babinski syndrome)</td>
</tr>
<tr>
<td>Retinopathies associated to malignant melanoma</td>
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</tbody>
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Note: All listed conditions and symptoms are associated with visual disturbances and require medical intervention.
CONCLUSIONS

Visual acuity loss has different causes and pathophysiological mechanisms, with a great burden in everyday’s life, severely affecting patients and their quality of life. Early diagnosis and specific treatments are without any doubt the only way to reduce this discomfort. Ophthalmologists and neurologists should work together to overcome many problems of differential diagnosis.

REFERENCES