

ISCEV Guide to Visual Electrodiagnostic Procedures

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1. Introduction

Visual electrophysiologic testing incorporates different tests based upon the recording of evoked electrical potentials in response to visual stimuli, using electrodes situated on the surface of the eyes, the peri-orbital skin or scalp. The tests are non-invasive and provide an objective indication of function relating to different locations and cell types within the visual system. This document has been developed by the International Society for Clinical Electrophysiology of Vision (ISCEV) to provide an introduction to visual electrodiagnostic procedures in widespread use, and to suggest some of the common clinical indications for which these tests are applicable. Detailed specifications for each procedure may be found in the appropriate ISCEV Standards^{1,2,3,4,5}. The basic principles of electrodiagnostic testing are outlined in this document, but the document is not intended to be prescriptive or to address every clinical scenario and is not a mandate for specific procedures on individual patients. It is stressed that the utility of electrophysiological testing is greatest when performed in conjunction with clinical assessment by specialist eye care professionals, essential to enable appropriate clinical management, and that the results must always be placed in clinical context.

This guideline describes the basic methods and underlying principles of testing for each of the standard test modalities; the full-field flash electroretinogram (ERG); the multifocal electroretinogram (mfERG), the pattern electroretinogram (pattern ERG or PERG), the electro-oculogram (EOG) and the cortical-derived visual evoked potential (VEP). Common clinical presentations and symptoms are described with illustrative examples and suggested investigation strategies.

2. The Electrophysiological tests

2.1 The full-field ERG

The full-field ERG (Figure 1a) is a global response of the retina and provides an assessment of generalized retinal function under light- and dark-adapted conditions. A Ganzfeld (German for “whole field”) stimulator, which provides a uniformly illuminated field, is used to deliver a range of full-field flash stimuli that evenly illuminate the maximal area of retina. Responses are recorded with corneal electrodes close to the inferior corneal limbus or with skin electrodes attached to the lower eyelids. Several types of corneal electrode may be used including contact lens, fibre, jet and gold foil electrodes. Reliable interpretation of recordings requires comparison with electrode-specific and age-matched normative data. The pupils are dilated to maximize retinal illumination and to minimize inter-subject and inter-visit variability. The normal test-retest variability of ERG parameters is an important consideration if used to monitor disease progression or the safety or efficacy of treatments.

The ISCEV-standard protocol includes scotopic recordings after 20 minutes dark adaptation (DA) to flash strengths of 0.01, 3.0 and 10.0 cd.s.m⁻² (DA 0.01; DA 3.0; DA10.0). The dim flash (DA 0.01) ERG arises in the inner retinal rod bipolar cells and is the only standard testing modality which

selectively monitors rod system function. Abnormality of the DA 0.01 ERG does not specify the locus of dysfunction and can be caused by either rod photoreceptor dysfunction or by selective dysfunction occurring post-phototransduction or at the level of the inner retinal rod bipolar cells. The DA 3.0 and DA 10.0 (strong flash) ERGs are mixed rod and cone system responses but the dark-adapted rod system contribution dominates in a normal subject. Approximately the first 8ms of the cornea-negative a-wave reflects rod hyperpolarizations and as the a-wave in the DA 10.0 ERG is of shorter peak time and larger than in the DA 3.0 ERG, it provides a better measure of rod photoreceptor function. The subsequent cornea-positive b-wave arises largely in the rod On- bipolar cells and reflects function that is post-phototransduction or inner retinal. Thus the DA strong flash ERG enables localization of dysfunction to the rod photoreceptors (a-wave reduction and concomitant b-wave reduction) or to a level that is post-phototransduction or inner retinal (sparing of the a-wave; b-wave reduction). The DA oscillatory potentials (OPs) are small high frequency components normally visible on the rising limb of the DA 3.0 and DA 10.0 ERG b-waves and are thought to arise in the amacrine cells. Reduction in the OPs is often associated with other ERG abnormalities but may occur selectively in some disorders. The cone system contribution to both DA ERG a- and b-waves is minor in a normal retina but can adopt greater significance in patients with disease primarily or exclusively affecting the rod system. Photopic recordings provide two measures of generalized cone system function, both obtained to a flash strength of 3.0cd.s.m^{-2} , after a standard period of 10 minutes light adaptation (LA) in the Ganzfeld with a constant background luminance of 30cd.m^{-2} . A 30Hz flash stimulus, superimposed on the photopic background, is used to elicit the photopic 30Hz flicker ERG (LA 30Hz), generated largely by inner retinal structures, and a single flash cone ERG (LA 3.0), consisting mainly of a photopic a- and b-wave. The LA 3.0 ERG a-wave arises in the cone photoreceptors and Off- bipolar cells; the b-wave is dominated by a combination of inner retinal cone On- and Off- bipolar cell activity.

The full field ERG enables the distinction between generalized outer and inner retinal dysfunction and predominant rod or cone system dysfunction. Symptoms and/or clinical signs may suggest a retinopathy but the presence, severity and nature of retinal dysfunction cannot always be inferred from the clinical findings and in clinical context ERGs can help differentiate between a wide range of disorders when appropriately placed in clinical context (see below and table 1). It is stressed that the full-field ERG is largely generated by the retinal periphery and there is minimal contribution from the macula. Electrophysiologic assessment of macular function requires the use of different techniques such as the pattern ERG or multifocal ERG.

2.2 The Pattern ERG

The ISCEV-standard PERG is recorded to an alternating high contrast checkerboard using a corneal electrode. The transient PERG has two major components of diagnostic value; a positive polarity P50 and a negative polarity N95 (Figure 1a and Figure 2). Both components reflect retinal ganglion cell function but there is an additional more distal retinal contribution to the P50 component. Both P50 and N95 depend on the function of the macular cones and P50 reduction and/or delay can characterize macular dysfunction. Selective reduction in N95 with preservation of P50 suggests dysfunction at the level of the retinal ganglion cells. There may be additional P50 reduction in severe or chronic retinal ganglion cell dysfunction, but in such circumstances P50 usually shortens in peak time, reflecting loss of the retinal ganglion cell contribution to the P50 component. Pattern ERGs are attenuated by poor refraction and ocular media opacity and care must be taken to optimize the optical quality of the checkerboard stimulus; for this reason contact lens electrodes are not suitable. Preservation of P50 helps to establish the effective quality and stimulus contrast of the checkerboard in patients who may have poor visual acuity for reasons other than maculopathy. Comparison of responses to a standard and additional large stimulus field may help characterize the area of macular dysfunction, although spatial resolution is lower than for the mfERG.

The pattern ERG provides a measure of macular and retinal ganglion cell function and complements the full-field ERG, in differentiating between maculopathy and generalized retinopathy. The PERG also enables a more meaningful evaluation of a VEP, to exclude a macular cause of VEP abnormality and to provide an additional assessment of retinal ganglion cell involvement (see below).

2.3 The Multifocal ERG

The ISCEV-standard mfERG (Figure 3a) provides a measure of cone system function over 61 or 103 discrete hexagonal retinal areas, within the central 40-50 degrees of the posterior pole centered on the macula. Each hexagonal stimulus element is modulated rapidly to display white or black frames (each frame is 13.33ms for a frame rate of 75Hz) according to an irregular but predetermined binary sequence known as a “pseudo-random” or “m-sequence”, identical but time-shifted for each hexagonal stimulus element. The signal associated with a particular hexagon is extracted from a continuous recording from each eye, using an automated cross-correlation analysis. The responses can be mathematically stratified into components associated with single illumination events (the first-order kernel), used for ISCEV-standard testing, or with sequences of multiple illumination events (the second-order and higher-order kernels). The hexagons of the ISCEV-standard stimulus array are scaled to elicit comparable response amplitudes from each stimulus region, resulting in larger hexagons in the peripheral portions of the stimulus pattern. Corneal electrodes are required as signals are small. The optical quality of the stimulus is important and patients should be optimally refracted and must fixate accurately on a central target or cross-hairs throughout the recording period. There is a compromise between spatial resolution (smaller, more numerous hexagons) and the recording duration necessary to complete the entire m-sequence to obtain satisfactory responses with a reasonably high signal-to-noise ratio. If the area of dysfunction shows reasonably good radial symmetry, interpretation may be facilitated by averaging waveforms from all the hexagons in each concentric ring in the stimulus pattern (ring-averaging). There are two major response components; an early negative polarity N1 component is derived from cone bipolar cells with a cone photoreceptor contribution and a later positive polarity P1 component that arises in cone bipolar cells.

The spatial resolution of the mfERG is better than for the PERG and this enables improved characterization of focal central, annular, hemifield or discrete paracentral areas of posterior pole dysfunction, but reliable recording requires good patient fixation. Illustrative examples of mfERG recordings are shown in a case of RP with central macular sparing (Figure 3b), in macular dystrophy (Figure 3c) and in a patient with an enlarged blind spot (Figure 3d). The mfERG is also a useful adjunct to the VEP and is less affected by optical factors than the PERG; there is no retinal ganglion cell contribution to the mfERG and a normal response excludes primary macular photoreceptor dysfunction as cause of VEP abnormality or central visual loss. However, in some conditions such as CME, the mfERG may be preserved or less affected than the PERG.

2.4 The Electro-oculogram

The EOG is used to assess generalized RPE function. There is potential difference between the apical and basal surfaces of the RPE that results in a dipole across the eye, with the cornea being positive with respect to the back of the eye. This potential difference may be recorded using skin electrodes placed at the medial and lateral canthus of each eye during uniform horizontal saccades. During a standard 15-minute period of dark adaptation there is a fall in the recorded standing potential across the RPE and globe, typically reaching a minimum at 10–15 minutes, and this is referred to as the Dark Trough (DT). The dark phase is followed by a 15-minute period of continuous light adaptation to a standard photopic background (100 cd.m^{-2}), provided by a Ganzfeld stimulator. Following light onset there is an increase in the trans-epithelial potential resulting in the EOG light peak (LP). The LP:DT ratio (Arden index) provides an indirect measure of generalized RPE function. The development of a normal EOG light peak requires not only a normally functioning RPE, but also normally functioning rod photoreceptors, with the degree of EOG abnormality broadly corresponding to the degree of rod-

derived ERG abnormality. An EOG assessment of generalized RPE/photoreceptor function is most useful when interpreted in the context of normal or only mildly subnormal rod-mediated ERG findings. In the presence of severe rod dysfunction from any cause, the EOG will be abnormal, and not additionally informative about the function of the RPE. Common causes of generalized RPE dysfunction are outlined below.

2.5 Visual Evoked Potential

The VEP describes the scalp recording derived from electrical currents generated in the visual cortex in response to visual stimulation. The visual cortex is activated primarily by the central visual field and VEPs reflect the function of the whole of the retino-cortical visual pathway. Recording electrodes are positioned on the scalp according to anatomical landmarks using a standardized “International 10/20 system” measurement method. The recording montage includes at least one occipital electrode (Oz) referred to a mid-frontal reference (Fz). Computerized signal averaging is used to extract the time-locked VEP from spontaneous brain activity (the electroencephalogram or EEG).

The ISCEV standard for VEP testing describes three stimulus modalities; pattern reversal, pattern onset-offset and diffuse flash stimulation. A reversing checkerboard, similar to that for PERG, is used to record the pattern reversal VEP, generally most useful for the assessment of optic nerve function, but requiring an adequate level of fixation and compliance. The normal pattern reversal VEP has a prominent positive component at approximately 100ms (P100; Figure 2), although normal ranges differ and are age and laboratory dependent. Multichannel VEPs, in excess of the current ISCEV standard, are needed to detect optic nerve misrouting or to detect and characterize chiasmal or retrochiasmal dysfunction. Pattern onset-offset (pattern appearance) stimulation is less commonly used in the diagnosis of optic nerve disease than pattern reversal, but has the advantage of being less affected by nystagmus, non-optimal fixation and voluntary defocus (common in malingerers). Multichannel onset-offset VEPs are particularly useful in the detection of visual pathway misrouting in albinism. Flash VEPs are generally less sensitive to optic nerve dysfunction than pattern VEPs, but may be used in young children or when patients cannot fixate or comply with testing. They are also useful in the presence of media opacity and the use of stronger non-standard flashes may be helpful to establish the integrity of the visual pathway. There is wider variability than for pattern VEPs and an inter-ocular comparison is often most useful. Flash VEPs may occasionally reveal abnormalities in the presence of normal pattern VEPs and this can occur in rare cases of optic neuritis, in some cases of optic nerve sheath pathology or due to unsuspected retinopathy. Multichannel flash VEPs can also reveal the visual pathway misrouting associated with albinism in children, but flash VEPs are usually normal in adults with albinism.

Visual evoked potentials provide an important objective test in the investigation of suspected optic nerve disease or post-retinal visual pathway dysfunction. The pattern reversal VEP may be assessed in terms of P100 timing, amplitude and waveform shape but abnormalities are not specific and can reflect optic nerve or macular dysfunction, but can also be caused by poor compliance or sub-optimal refraction. Reliable interpretation of pattern VEP abnormality usually requires complementary assessment to exclude a macular cause. Similarly a flash ERG may exclude a retinal cause of flash VEP abnormality. There are numerous causes of optic nerve disease and VEP abnormalities are rarely specific, but may suggest or support a suspected diagnosis when interpreted in clinical context. Common causes of optic neuropathy are outlined below.

3. Clinical indications for visual electrophysiology

Symptoms, signs and circumstances that frequently prompt referral for visual electrophysiology are outlined below, with selected examples illustrated in Figures 1, 2 and 3, chosen to illustrate the underlying principles of testing. Accurate localization of dysfunction within the visual pathway may require complementary testing with different techniques and a suggested test strategy is outlined in

Figure 4. It is stressed that multiple tests may not be needed in all patients and that electrophysiological findings and accurate diagnosis require interpretation in the context of the clinical findings. A comprehensive list of all conditions that may prompt visual electrophysiologic examination is beyond the scope of this guideline, but diagnoses that commonly benefit from testing and typical findings are summarized in table 1.

3.1 Visual acuity loss

Visual acuity loss may be caused by inherited and acquired causes of maculopathy (with or without retinopathy), optic nerve and visual pathway disease but this may not be obvious on clinical grounds alone and the distinction is enabled by electrophysiological testing.

3.1.1 Retinal and RPE disorders

The pattern ERG and mfERG may be used to assess the severity of macular dysfunction (Figures 1 and 3) in the presence of fundus abnormality or used to detect dysfunction in occult cases of maculopathy or macular dystrophy. If there is visible evidence of maculopathy on fundus examination, an ERG will determine whether there is peripheral retinal involvement e.g. differentiating between macular dystrophy (normal full-field ERG; Figure 1b) and cone and cone-rod dystrophy (see below and Figure 1c). Common reasons for referral include bull's eye lesions, which may be associated with macular dystrophy, cone or cone-rod dystrophy, or acquired dysfunction, for example caused by hydroxychloroquine toxicity. In Stargardt disease (ABCA4-retinopathy), the most common monogenic cause of inherited macular/retinal dystrophy, there is usually visible maculopathy and fleck lesions (not always present in children or early disease) and ERGs establish whether dysfunction is confined to the macula or whether there is generalized cone or cone and rod involvement.

Rapid loss of visual acuity may occur in acquired disorders such as paraneoplastic (carcinoma associated retinopathy; CAR), or autoimmune retinopathy (AIR), which are often without fundus abnormality at presentation, and are typically associated with pronounced rod and cone photoreceptor dysfunction, evident on ERG testing. In cases of vitelliform macular lesions an ERG and EOG are indicated; Best vitelliform macular dystrophy is characterized by a severely reduced EOG light peak to dark trough ratio in the absence of ERG abnormality, confirming generalized RPE dysfunction and largely excluding other disorders that may resemble Best disease on fundus examination, including some pattern dystrophies such as adult onset vitelliform macular dystrophy (table 1).

3.1.2 Optic nerve/post-retinal disorders

In the absence of obvious fundus abnormality the pattern VEP in combination with a PERG or mfERG distinguishes optic nerve dysfunction from occult macular disease. The pattern VEP is usually abnormal in macular disease and PERG P50/central mfERG preservation largely excludes macular dysfunction as a cause of pattern VEP abnormality.

Acute visual acuity loss with pain on eye movement is typical of optic neuritis and VEPs are typically delayed in keeping with demyelination (Figure 2), with or without amplitude reduction; the VEP abnormality usually persists even if visual acuity improves. VEP abnormalities may occur in an asymptomatic eye and in visually asymptomatic patients with multiple sclerosis, consistent with subclinical demyelination. Approximately 35% of patients with optic nerve demyelination manifest a reduced PERG N95:P50 ratio, in keeping with retrograde involvement of the retinal ganglion cells and occurring 4-6 weeks after presentation, although this can occur in any form of optic neuropathy. A sudden painless and irreversible loss of vision is typical of non-arteritic anterior ischemic optic neuropathy (NAION) and unlike demyelination, pattern VEPs typically show amplitude reduction without significant delay (Figure 2). In arteritic ischemic optic neuropathy (AION), there is usually severe visual loss and gross VEP abnormality. Leber hereditary optic neuropathy (LHON) typically

presents with sudden sequential, painless visual loss and pattern VEPs are usually undetectable or severely abnormal at presentation; PERG P50 amplitude is typically normal providing fixation is adequate, but there may be marked reduction in N95 in the acute stages, in keeping with primary ganglion cell dysfunction.

Compressive lesions of the visual pathways are associated with progressive or insidious visual acuity loss, although if unilateral this may be noticed suddenly by the patient. If a unilateral optic nerve lesion is anterior to the optic chiasm, there will be unilateral pattern VEP abnormality. Localization of dysfunction posterior to the optic nerves requires multichannel VEP recordings. Chiasmal dysfunction results in a “crossed asymmetry”, such that the VEP from each eye is abnormal over a different hemisphere. Retrochiasmal dysfunction results in an “uncrossed” asymmetry such that monocular VEPs from both eyes are abnormal over the same hemisphere. Progressive visual loss is also a feature of dominant optic atrophy and nutritional optic neuropathies such as that caused by vitamin B12 deficiency. Toxic etiology includes ethambutol, methyl-alcohol poisoning (also associated with retinopathy) and rare cases of tobacco toxicity. Visual loss may also result from injury to the occipital cortex usually resulting in both pattern and flash VEP waveform degradation or distortion.

3.1.3 Non-organic visual loss

In cases of unexplained or suspected “functional” visual loss normal electrophysiology helps to exclude an organic cause. A well-formed pattern reversal VEP is incompatible with a visual acuity of approximately 6/36 or worse, although care must be taken to ensure adequate patient compliance during testing. Flash VEPs are usually normal and even if there is dysfunction with non-organic overlay, it is difficult to reconcile a detectable flash VEP with “no perception of light” vision. The significance of pattern VEP abnormality depends on the results of macular testing with PERG P50 or mfERG and the importance of flash VEP abnormality may similarly depend on the absence of significant full-field ERG abnormality. Normal visual electrophysiology does not preclude the presence of underlying organic disease and particular caution must be exercised if there is a possibility of higher cortical dysfunction.

3.2 Night blindness

Night blindness (nyctalopia) can result from generalized rod system dysfunction and this may be confirmed or excluded using a full-field ERG. The DA 3.0 and DA 10.0 ERGs enable localization of dysfunction to the rod photoreceptors (a-wave reduction and concomitant b-wave reduction) or to a level that is post-phototransduction or inner retinal (sparing of the a-wave; b-wave reduction).

3.2.1 Night blindness due to rod photoreceptor dysfunction

In progressive retinal degenerations such as retinitis pigmentosa (RP), which in mild cases may be associated with a normal or near-normal fundus appearance, there is ERG evidence of a rod-cone dystrophy (Figure 1d). The severity of generalized retinal dysfunction in RP varies, but there may be preserved visual acuity and relative preservation of macular function until the late stages in many cases, as revealed by PERG P50 (Figure 1d) or mfERG (Figure 3b). Progressive degeneration encroaching upon the macula may lead to eventual blindness and it is important to distinguish from other causes of rod system dysfunction. In RP, the DA 0.01 ERG is typically reduced and the bright flash (DA 3.0 and DA 10.0) ERGs show a-wave reduction. The reduction in the a-wave confirms rod photoreceptor dysfunction; there is concomitant b-wave reduction because the b-wave is generated “downstream” from the abnormal rod photoreceptors. The LA 30Hz and LA 3.0 ERGs are typically delayed and/or reduced but dysfunction is milder than in the rod system. The reduction in the a-wave makes the distinction from the two common forms of congenital stationary night blindness (complete and incomplete CSNB; see section 3.2.2). There are other rare forms of CSNB that cause severe rod-driven ERG abnormalities (strong flash ERG a-wave reduction) but with spared cone system function and these include “Riggs-type” CSNB, Oguchi disease and fundus albipunctatus. In the latter two

disorders there are usually characteristic fundus abnormalities and improvement or recovery of rod system function after prolonged DA (see table 1 for a summary). The fundus appearance in fundus albipunctatus may be similar to patients with retinitis punctata albescens (Bothnia dystrophy); patients with Bothnia dystrophy may also show partial ERG recovery following prolonged dark adaptation but the phenotype is more severe than in fundus albipunctatus and evolves to a progressive rod-cone dystrophy.

Acquired night blindness with a normal fundus can occur in Vitamin A deficiency and CAR although in rare cases of CAR there may be an electronegative ERG (see below). The ERGs in vitamin A deficiency are characterized by severe rod-system dysfunction and normal or near-normal cone system function, similar to the ERGs in “Riggs-type” CSNB. However the ERGs in vitamin A deficiency usually return to normal following treatment.

3.2.2 Night blindness due to dysfunction occurring post-phototransduction

Complete and incomplete CSNB are associated with a normal fundus and generalized retinal dysfunction that is post-phototransduction (Figure 1e and 1f), with normal (or near-normal) a-waves and electronegative DA 3.0 and DA 10.0 ERG waveforms (b:a ratio <1). In complete CSNB the DA 0.01 ERG is undetectable. The LA 30Hz ERG, although often of normal amplitude may have a slightly broadened trough and often shows borderline or mild peak time delay. The LA 3.0 ERG has normal a-wave amplitude but with a broadened bifid trough and a b-wave with a sharply rising peak with no oscillatory potentials; the b:a ratio varies but is usually mildly subnormal. The shape of the DA and LA ERG waveforms are characteristic of loss of On-pathway function with Off-pathway preservation, also evident in the long-duration On-Off ERG, which reveals an electronegative On response and a normal Off response. Complete CSNB is caused by a defect in 1 of 5 genes (table 1), expressed by On-bipolar cells and consistent with the ERG abnormalities. In incomplete CSNB the DA0.01 ERG is present but subnormal. LA 30Hz ERGs are markedly reduced and have a bifid shape. The LA3.0 ERG is markedly subnormal with a low b:a ratio. Long duration stimulation reveals reduction in both the On b-wave and Off d-wave. It is noted that the 2 genes implicated in incomplete CSNB (table 1) are involved in neurotransmitter release from the photoreceptor pre-synaptic membrane, consistent with ERG evidence of both On- and Off bipolar cell dysfunction.

Acquired night blindness with a normal fundus and electronegative ERG can occur in melanoma-associated retinopathy (MAR) and rarely in CAR (see above). MAR is rare but is associated with malignant melanoma and the ERG findings are identical to those in complete CSNB (Figure 1e). The ISCEV-standard ERG features in MAR, CAR and vitamin A deficiency are different to each other, but are indistinguishable from some of the inherited disorders mentioned above, highlighting the importance of clinical context in the interpretation of ERGs.

3.3 Photophobia

Photophobia is commonly associated with generalized cone system dysfunction and can be an early symptom in cone and cone-rod dystrophies. In cone dystrophies the LA 30Hz and LA 3.0 ERGs show delay and/or amplitude reduction and cone-rod dystrophy there is additional abnormality of the DA ERGs (Figure 1c). In both conditions there is usually severe macular dysfunction evident on PERG (Figure 1c) or mfERG testing. Rod monochromacy (achromatopsia) is characterized by severe cone system dysfunction from early infancy; the LA ERGs are typically undetectable but the DA 0.01 ERG, selective for the rod system, is normal and the DA 10.0 ERG is normal or shows mild reduction in the a- and b-waves, due to a loss of the normal dark-adapted cone system contribution. The ERG findings in S-cone monochromacy (a form of “X-linked incomplete achromatopsia”) are similar but DA ERGs may be additionally attenuated due to high myopia; there may be a markedly abnormal (but detectable) LA 3.0 ERG and the short-wavelength (“blue”) flash ERG is relatively preserved. Congenital photophobia may also be a feature of albinism. Photophobia is rarely caused by dysfunction confined to the macula. Acquired causes of photophobia include retinal inflammatory

disease such as uveitis and birdshot retinochoroidopathy (BRC) both associated with a high incidence of generalized cone system dysfunction, AIR and CAR. Photophobia is a rare feature of optic nerve disease but can also occur in neurological disorders such as migraine, meningitis and in carotid artery or vertebral artery disease.

3.4 Visual field loss

Peripheral visual field constriction is a common feature of rod-cone dystrophy (RP) and this can occur without classical intra-retinal pigment deposition, particularly in children. Cone and cone-rod dystrophies may present with visual field defects include central scotomata, generalized depression of sensitivity, ring scotomata and peripheral field loss if there is relative sparing of central macular function. Peripheral visual field loss may also occur in inflammatory retinal disorders such as BRC, associated with variable retinal dysfunction but often characterized by generalized cone system dysfunction, manifest as delay in the LA 30Hz ERG, and sometimes associated with additional inner retinal rod system involvement (reduction in DA 10.0 ERG b:a ratio) which may be reversible following treatment (Figure 1g and 1h). In Acute zonal occult outer retinopathy (AZOOR) there is usually field loss disproportionate to visible fundus changes and persistent photopsia within the scotoma. Full-field ERG abnormalities are common and some may show a reduction in the EOG light peak to dark trough ratio, not explained by abnormalities in rod function. Autoimmune disorders, such as CAR and AIR may also present with rapid visual field constriction and marked ERG abnormality (see above). The pattern ERG is sensitive to ganglion cell dysfunction and nerve fiber layer loss in glaucoma, and can be of value in the evaluation of “glaucoma suspects” with glaucomatous risk factors such as elevated intraocular pressure, or optic nerve head changes, prior to the measurable loss of visual field. Homonymous hemianopic visual field defects usually reflect chiasmal or retrochiasmal brain lesions and these may be detected by multichannel VEP recordings, and require prompt further investigation. Field loss may also be seen in shallow retinal detachments and retinoschisis with concomitant full-field ERG changes, and clinical or ultrasound eye examination is essential

3.5 Disc pallor

Disc pallor may be a feature of optic neuropathy or retinopathy, including cone and cone-rod dystrophies. In central retinal artery occlusion (CRAO) there may be unilateral retinal edema and a “cherry red” spot at the fovea in the acute phase, but after a few weeks this resolves as disc pallor develops. The sub-acute and chronic phases may be mistaken for ischemic optic neuropathy and the electrophysiology enables the distinction. The ERG in CRAO has an electronegative DA 3.0 or DA 10.0 ERG and there is usually marked involvement of the LA ERGs, in keeping with generalized inner retinal dysfunction. There are several other potential masquerades of optic neuropathy including occult maculopathy (inherited or acquired) and central serous chorioretinopathy (CSR); both may manifest PERG P50 or central mfERG abnormalities. In acute idiopathic blind spot syndrome (AIBSS) the mfERG may characterize the nasal area of reduced function (Figure 3d). Occult retinopathy (including acute zonal occult outer retinopathy; AZOOR), autoimmune and paraneoplastic retinopathies (see below) typically show marked ERG abnormalities and in posterior scleritis (which like optic neuritis may be accompanied by pain on eye movement) there may be inflammatory changes affecting the retina that cause ERG abnormality.

3.6 Nystagmus

Congenital nystagmus is a feature of several ocular and neurological disorders. Isolated idiopathic congenital motor nystagmus (CMN) is not associated with other ocular abnormalities; pattern reversal

VEP and PERG may be difficult or impossible to record due to eye movements but flash VEPs and full-field ERGs are normal. Common retinal causes of nystagmus include Leber congenital amaurosis (LCA), associated with severe generalized photoreceptor dysfunction (DA and LA ERGs are severely reduced or undetectable), cone and cone rod dystrophy, rod and S-cone monochromacy and complete and incomplete CSNB, characterized by different ERG phenotypes (see above). Nystagmus is also associated with ocular and oculo-cutaneous albinism (see above) and diagnosis in the former may be difficult in the absence of obvious skin depigmentation.

Acquired nystagmus may result from drug toxicity or medication that impairs the function of the labyrinth, thiamine or vitamin B12 deficiency, head injury, stroke, multiple sclerosis or any disease or injury of the brain that affects neural centers that control eye movements. Exclusion of afferent visual pathway dysfunction with electrophysiology may provide an important contribution to the management of such cases.

3.7 Vascular Retinopathies or Ischemic status of Retina

The full field ERG is sensitive to retinal ischemic disorders affecting the inner retina. There may be reductions in the DA 3.0 and DA 10.0 ERG b:a ratios, the DA oscillatory potentials are usually abnormal or extinguished and LA 30Hz ERGs show prolonged peak times and waveform distortions. The ERG may be invaluable in detecting ischemic CRVO, progression of non-ischemic to ischemic CRVO and in the diagnosis of Ocular ischemic syndrome especially when the carotid Doppler scans are normal or equivocal. The ERG has advantages over commonly used fluorescein angiography in being safe and non-invasive, providing information on deeper layers and peripheral areas of retinal blood supply and may be informative in patients with systemic co-morbidities or pregnancy, in patients allergic to fluorescein dye or in cases of retinal hemorrhage obscuring the fundus view.

3.8 Ocular media opacity.

Full-field ERGs and flash VEPs can provide valuable information in patients with suspected retinal or visual pathway disease when the fundus is obscured or when the use of retinal imaging techniques is precluded by an opaque ocular media. Integrity of retinal and visual pathway may be important considerations prior to treating patients with corneal lesions, cataracts or vitreous hemorrhage, particularly if there is a history of retinal detachment, retinal or neurological involvement. A normal or relatively preserved ERG or flash VEP may suggest a better prognosis for improved vision. An abnormal full-field ERG may suggest generalized retinal dysfunction but may also occur in vitreous hemorrhage. An abnormal ERG does not exclude central visual recovery because it does not assess macular function. It is noted that the ERG is usually abnormal in the presence of intra-ocular silicone oil tamponade (for retinal detachment) but interpretation confounded because the oil impedes conduction of the electrical signals from the retina to the corneal surface.

3.9 Family history of visual pathway disease

Visually asymptomatic patients with a family history of retinal or optic nerve disease or suspected cases of syndromic retinal dystrophy may require electrophysiological testing for evidence of subclinical disease. For example visually asymptomatic obligate carriers of X-linked RP usually manifest abnormal and asymmetrical ERG abnormalities, irrespective of whether there is fundus abnormality, whereas the ERGs in carriers of X-linked choroideremia are usually normal until late in life. Carriers of X-linked ocular albinism and patients with rubella retinopathy may also have abnormal fundus pigmentation; the ERGs are normal in the former and normal or near-normal in the latter. There is variable expressivity in (autosomal dominant) Best disease such that some heterozygotes have a normal fundus and an EOG may be needed to confirm the diagnosis. Similarly VEP and PERG N95 abnormalities may indicate optic nerve and retinal ganglion cell dysfunction in cases of suspected dominant optic atrophy.

3.10 Monitoring of disease progression, treatment efficacy and safety

Serial testing may assist the distinction between stationary and progressive conditions, important for diagnosis and patient counseling. Pattern and flash VEPs have diverse applications and may be used to monitor visual pathway maturation in infants with poor vision or amblyopia or to monitor optic nerve function in patients with known neurological disease. In inflammatory retinal diseases such as BRC, the ERGs can be used to monitor efficacy of treatment objectively (Figure 1g and 1h) thus informing clinical management and titration of potentially toxic medication. Worsening VEPs may prompt the need for surgical intervention in dysthyroid eye disease or in neurological disorders, irrespective of stable neuroradiology.

Several medications commonly administered systemically for non-ocular conditions are potentially toxic to the macula, retina or optic nerves (table 1) and pre-treatment assessment and monitoring may be considered. Intraocular drugs, intraoperative dyes and bright lights of ophthalmic surgical equipment have become another source of toxic/phototoxic maculopathy that may need retinal and macular electrophysiology testing for monitoring, for clinical study or for diagnosis. ERG evaluations are also becoming an integral part of various clinical trials comparing outcome efficacies of various surgical or medical procedures involving the macula such as macular holes, epiretinal membranes, anti-VEGF treatments, macular detachments and central serous chorioretinopathy. Similarly ERGs may be used to monitor retinal safety of new treatments and as objective outcome measures in clinical trials that aim to restore visual function or arrest disease progression.

4.0 Special considerations and indications for ERG and VEP testing in infants and children.

Accurate diagnosis may be difficult in young children who are unable to describe their visual symptoms or who are difficult to examine. The objective data provided by electrophysiological testing are fundamental to the management of the child with suspected visual pathway dysfunction but there are important considerations relating to maturation of responses, ability to comply with testing and causes of visual pathway dysfunction more specific to the pediatric population. Both ERG and VEP responses show profound developmental changes during infancy and childhood and although all visual electrophysiological values are considered in relation to age, it is even more important in young patients. Infants up to the age of about two years can frequently undergo successful ERG testing without general anesthesia, while being held in a parents' lap, either using only topical anesthetic eye drops and corneal electrodes or by using surface electrodes on the lower eyelids. It may be appropriate to shorten the standard ERG protocol and many practitioners start with light-adapted ERGs and perform limited dark adaptation, dependent upon the compliance and comfort of the child. VEP testing in infants is equally feasible, but may require simple flash stimulation, if steady fixation on the center of the VEP pattern stimulus cannot be induced with a moving toy, jangling keys or similar to encourage central fixation.

Examination under anesthesia may enable the use of corneal electrodes in the non-compliant child, but anesthesia usually alters ERG timing and amplitudes, and interpretation requires caution. Similarly the use of skin electrodes limits sensitivity since the signal:noise ratio is lower, but in this age group there is rarely a need to detect subtle abnormalities and most clinically appropriate questions may be easily addressed. For example is there a detectable ERG, is there a functioning cone system, is there a response after dark adaptation and is there an electronegative ERG waveform? The cortical neurons which drive the VEP are much more susceptible to general anesthesia than the retina, precluding reliable VEP recordings.

4.1.1 Unexplained visual loss

Absent or impaired visually-mediated behavior may indicate a disorder affecting any level of the visual system. Babies who not fix and follow and presumed amblyopic patients that fail to respond to

treatment may require testing to confirm or exclude pathology. Early diagnosis of retinal dystrophy may be essential to identify young candidates who are potentially amenable to future experimental treatments. A normal ERG may also prompt the need for further investigations such as VEPs or neuroradiology. Non-organic visual loss is relatively common in older children and in such circumstances the electrophysiological data are usually normal even though there may be reported profound visual loss.

4.1.2 Congenital nystagmus.

The differential diagnosis includes several retinal disorders such as Leber congenital amaurosis, congenital stationary night blindness, and rod and S-cone monochromacy. The ERG will help differentiate these conditions. Young children with albinism show multi-channel flash VEP evidence visual pathway misrouting, although with increasing age (above about 5 years) this may be best demonstrated with pattern onset-offset VEPs. Flash VEPs and ERGs are normal in idiopathic motor nystagmus. Clinical examination is also needed to investigate or exclude TORCH infections like viral retinitis that result in nystagmus and variable ERG abnormalities.

4.1.3 Known or suspected hereditary disorders.

The ERG may be helpful in advising families with patients at risk of hereditary retinal disorders. The extent to which the various retinal dystrophies are detectable in early infancy is frequently not known, but a normal ERG at age 7 or 8 years of age largely excludes X-linked RP. Night blindness may be associated with RP or CSNB and ERGs help differentiate between progressive and stationary disorders. In young cases of suspected Best disease children may be unable to comply with EOG testing but testing of the parents will almost invariably identify the parent carrying the mutation, irrespective of whether the fundi are normal.

4.1.4 Perinatal infections.

Perinatal infections, particularly the “TORCH” agents may attack ocular tissues, with possible profound associated dysfunction. The most common perinatal infection is probably rubella retinopathy, which frequently results in mottled RPE pigmentation and a “salt and pepper” appearance but in such cases the ERG is usually normal or near-normal.

4.1.5 Perinatal brain injury.

Perinatal brain damage may lead to severe visual impairment. VEPs enable objective determination of the nature of the deficit and may help grade the severity of cortical dysfunction. However, it is important to recognize that VEPs do not reflect higher processing required for normal vision.

4.1.6 Trauma

In children who have suffered head/orbital trauma or suspected visual pathway injury, complementary retinal and VEP testing may localize dysfunction and help to confirm, exclude or distinguish between retinopathy and optic nerve or post-retinal dysfunction, particularly in those unable or too young to communicate verbally.

4.1.7 Delayed visual maturation.

Infants often present with “visual indifference”, showing little or no reaction to visual stimuli for several months. If the eye exam, ERG, and VEP are normal or near-normal, this provides reassurance, and the prognosis for development of normal or near-normal vision is reasonably good.

4.1.8 Monitoring for retinal drug toxicity.

The most common indication in this category is vigabatrin, which is used for the treatment of infantile spasms (West syndrome). The drug causes peripheral visual field constriction in approximately 30% of adults, and a post-marketing monitoring program is in effect in the USA. The ERG is helpful in monitoring patients who are too young or lack the ability to perform visual field testing.

4.1.9 Amblyopia

Children suspected of having amblyopia are often referred for electrophysiology to exclude other causes of poor vision. For example when visual acuity has not improved with patching and the fundi are normal or when visual acuity is reduced bilaterally. In amblyopic eyes, pattern reversal VEPs may show amplitude reduction. Delays in the major positive (P100) component can occur, but this tends to be more prominent in strabismic rather than anisometropic amblyopia. Pattern VEPs may also be used to monitor the efficacy of occlusion therapy in amblyopic and fellow eyes, but subjective assessment of vision (if possible) should generally take priority.

5. Complementary testing

Electrophysiologic testing complements routine ophthalmic examination, subjective tests of visual function and retinal imaging methods commonly employed in the assessment of patients with visual impairment. Electrophysiological methods are objective and uniquely assess aspects of function and dysfunction. Ophthalmic examination and imaging techniques may be normal in the presence of retinal and visual pathway dysfunction or may reveal abnormalities that do not correlate with the nature or severity of dysfunction. Optimal assessment is obtained with judicious use of widely used techniques including those outlined below.

5.1 Subjective assessment of function

5.1.1 Visual acuity.

Visual acuity (VA) testing is a long established method of assessing central visual function in almost any form of visual system pathology, from ptosis of the eyelids and corneal epithelial edema to retinal degenerations and optic neuropathies. However, VA loss is non-specific and cannot be used to localize dysfunction within the visual pathway. The VA does not give an indication of peripheral retinal function and may also be relatively or completely preserved in the presence of macular or optic nerve dysfunction. VA may be normal for example in paracentral and peripheral retinal derangements, nerve fiber bundle defects (as in glaucoma), in sub-acute optic neuritis and lesions of the posterior visual pathways which spare the projections of the central retina.

5.1.2 Visual fields.

Visual field testing is widely available, and with the advent of automated static perimetry, highly standardized and reproducible. Visual fields allow localization of visual impairment, with classic patterns of visual field loss associated with localized and generalized retinal disorders, macular and optic nerve disease, chiasmal disruptions, lesions of the lateral geniculate body and optic radiations, and cortical lesions. Pattern ERG and multifocal ERG can be of great value in distinguishing between macular and optic nerve disease, often associated with similar visual field abnormalities and often indistinguishable by VEP alone. Full-field ERG abnormalities are a “leading indicator” of

degenerative retinal disorders such as retinitis pigmentosa. Peripheral visual fields are important in the adequate assessment of degenerative retinal diseases such as retinitis pigmentosa, in which the extent of scotomas, and the presence of residual temporal islands of vision, are of great importance to the patient but cannot be adequately assessed by central Humphrey visual fields and peripheral automated visual field programs. It is highlighted that visual fields do not always correlate with objective suprathreshold electrophysiological measures of function.

5.1.3 Contrast sensitivity.

Loss of contrast sensitivity is readily documented with special eye charts designed for the task, or CRT-based vision testing devices, and can occur in the absence of significant VA reduction. The causes of reduced contrast sensitivity include optical problems such as corneal haze or cataract and complementary use of different electrophysiological tests (Figure 4) can differentiate these from a wide range of visual pathway disorders.

5.1.4 Color Vision Testing.

Color vision is an important visual faculty, and abnormalities may derive from retinal, optic nerve, or (rarely) cortical pathology. Commonly used Ishihara plates are highly sensitive to even minor dyschromatopsias, but detect only red-green (protan or deutan axis) abnormalities. Other sets of test plates, such as the H-R-R plates, also detect tritan axis problems. The common X-linked protan and deutan color vision defects are rarely associated with abnormalities in the ISCEV-standard ERG, but can be detected with non-standard chromatic stimuli. Absence or severe loss of normal color vision suggests more severe pathology, such as achromatopsia or optic nerve disease, which are readily detected by ERG or VEP.

5.1.5 Dark adaptometry.

Abnormalities of dark adaptation are difficult for patients and physicians to assess without formal testing, as normal difficulties seeing in dim light may be reported as abnormally impaired night vision. Formal dark adaptometry can be performed with specialized instruments, such as the Goldmann-Weekers Dark Adaptometer. Qualitative assessment can be readily obtained with much simpler materials, such as the Hyvarinen cone adaptation test, in which an examiner with normal dark adaptation, compares his/her adaptation with that of the patient, who is asked to sort colored plastic tiles in a very dim room. Abnormalities of dark adaptation generally imply retinal pathology, including CSNB, vitamin A deficiency, paraneoplastic retinopathies and degenerative disorders such as retinitis pigmentosa, usually readily differentiated by full-field ERG in clinical context.

5.2 Retinal imaging

Fundus photography has been available as a clinical tool since 1926 and fluorescein angiography was introduced in 1959. More recently, advances in fundus imaging have appeared with increasing frequency, not only documenting ophthalmoscopic findings, but extending the range of clinical perception in depth (ICG angiography) and resolution; spectral domain OCT now approaches the resolution of low-power microscopy, without the need to remove tissue from the eye for histologic processing. However, the enhanced capability of fundus imaging has not displaced physiologic methods in the armamentarium of the practicing ophthalmologist. The need to complement anatomic methods with studies of visual function is as keen as ever and perhaps more so as increasing detail in fundus imaging allows ever finer diagnostic distinctions to be made, for which the functional consequences must be determined.

5.2.1 Fundus photography

Fundus photography documents the appearance of the retina, and allows ready estimation of the size of fundus lesions. Digital photography has improved resolution and enabled more objective

assessments as well as multi-spectral imaging. Newer cameras provide a wider-field image far greater than the 30–40 degree fields of traditional fundus cameras, revealing important pathology of the peripheral retina which was previously unappreciated or more difficult to assess, especially in children.

5.2.2 Fluorescein angiography

Fluorescein angiography documents the extent and integrity of the retinal vascular compartment, and remains an important tool even in the era of advanced OCT imaging, which lacks the dynamic aspect of the evolving fluorescein angiogram. ICG angiography extends the range of angiographic imaging deeper into the choroid, demonstrating vascular structures and abnormalities hidden to traditional imaging with visible light.

5.2.3 Fundus autofluorescence

Fundus autofluorescence imaging (FAF) can reveal otherwise invisible manifestations of disrupted RPE metabolism. To short wavelength excitation the main fluorophore is lipofuscin, derived from the phagocytosis of shed photoreceptor outer segments in the RPE. The distribution of FAF across the posterior pole and abnormal accumulations or depletions of the FAF signal can detect or accentuate the appearance of lesions in a wide range of disorders, and the technique has largely replaced fluorescein angiography in the assessment of inherited retinal and macular dystrophies. Since the technique was introduced in the early 1990s, methods such as PERG and mfERG have helped establish the functional significance of common FAF abnormalities and the value of FAF in monitoring disease progression.

5.2.4 Optical coherence tomography

Optical coherence tomography (OCT) has revolutionized retinal evaluation. It is far superior to even the most careful ophthalmoscopy at detecting anatomic disruptions of the posterior pole, such as cystoid edema, vitreomacular traction, or shallow serous detachments of the retina or RPE. Moreover, the recognition of the role of the line of photoreceptor inner segment ellipsoid (or inner segment/outer segment junction) as an indicator of the integrity of the photoreceptors has clarified the diagnosis of many retinal disorders. For example, in many cases of “occult macular dystrophy” OCT may expose subtle or localized outer retinal loss. Focal OCT abnormalities do not always correlate with the severity of dysfunction or the function of surrounding retinal tissues.

5.2.5 Adaptive optics

Adaptive optics (AO) techniques use active optical elements to compensate for the optical aberrations of the eye and provide a non-invasive method for extending spatial resolution and studying the micro-morphology of the retina in-vivo. Clinical implications are only beginning to emerge but otherwise invisible disruptions in the photoreceptor mosaic have been documented in different retinal and macular disorders.

5.3 Genetic testing

Electrophysiology has a pivotal role to characterize disorders and the phenotypic variability associated with a known genotype or to guide the screening of genes associated with a known electrophysiological phenotype. Advances in molecular biology have enabled genotyping of many inherited retinal and macular dystrophies but the functional consequences remain difficult to predict due to allelic heterogeneity and other factors. In rare retinal dystrophies ERGs can be used to identify the gene responsible e.g. in enhanced S-cone syndrome (NR2E3), “cone dystrophy with supernormal ERG” (KCNV2) and RGS9-retinopathy, as outlined in table 1. It is more usual for the ERGs to suggest a range of disorders or possible genotypes e.g. in complete CSNB the ERG phenotype is

common to X-linked and autosomal recessive forms with mutations in 1 of 5 different genes and ERGs are additionally identical to those in melanoma associated retinopathy, highlighting the importance of interpretation in clinical context. The emergence of unbiased whole exome and whole genome sequencing may reveal novel or unexpected genetic alterations and electrophysiological interrogation likely to prove increasingly important to establish the functional consequences and genotype-phenotype correlations.

Figure captions

Figure 1. Representative full-field and pattern ERGs in a normal subject (a), in a case of macular dystrophy (b), cone-rod dystrophy (c), rod-cone dystrophy with relative sparing of macular function (d), complete CSNB (e), incomplete CSNB (f) and birdshot retinochoroidopathy (BRC) before treatment (g) and after treatment illustrating full recovery of the ERG and PERG (h). Recordings showed a high degree of inter-ocular symmetry except in BRC (data from other eye is not shown). Note the 20ms pre-stimulus delay in all single flash ERG recordings. Two responses for each stimulus condition are superimposed in all but the normal example (d) to illustrate reproducibility. Broken lines replace large blink artefacts occurring after the ERGs, for clarity.

Figure 2. Representative pattern reversal VEPs and PERGs in the affected (a, c) and fellow (b, d) eyes in a patient with non-acute optic neuritis (Subject 1; a, b) and in an elderly patient with a severe non-arteritic anterior ischemic optic neuropathy (Subject 2; c, d). The P100 component of the pattern VEP in optic neuritis shows a 35ms delay compared with the normal fellow eye, without significant amplitude reduction, consistent with optic nerve conduction delay; pattern ERGs are normal in this case and reveal no evidence of macular or retinal ganglion cell dysfunction. The pattern VEP P100 component in (c) is undetectable and PERG shows a reduced N95:P50 ratio and shortening of P50 peak time (inter-ocular difference 7ms) compared with the fellow eye (d), indicating severe optic nerve dysfunction with retinal ganglion cell involvement. Two responses for each stimulus condition are superimposed to illustrate reproducibility.

Figure 3. Multifocal ERGs recorded to a 103-element stimulus array in a representative normal subject (a), in a case of retinitis pigmentosa showing relative sparing of central macular function (b), in a case of macular dystrophy (c) and in a patient with an eccentric nasal area of retinal dysfunction consistent with an enlarged blind spot extending inferiorly in the right eye (d). MfERGs in cases a, b and c showed a high degree of inter-ocular symmetry; abnormalities were unilateral in d. Traces are shown in retinal view.

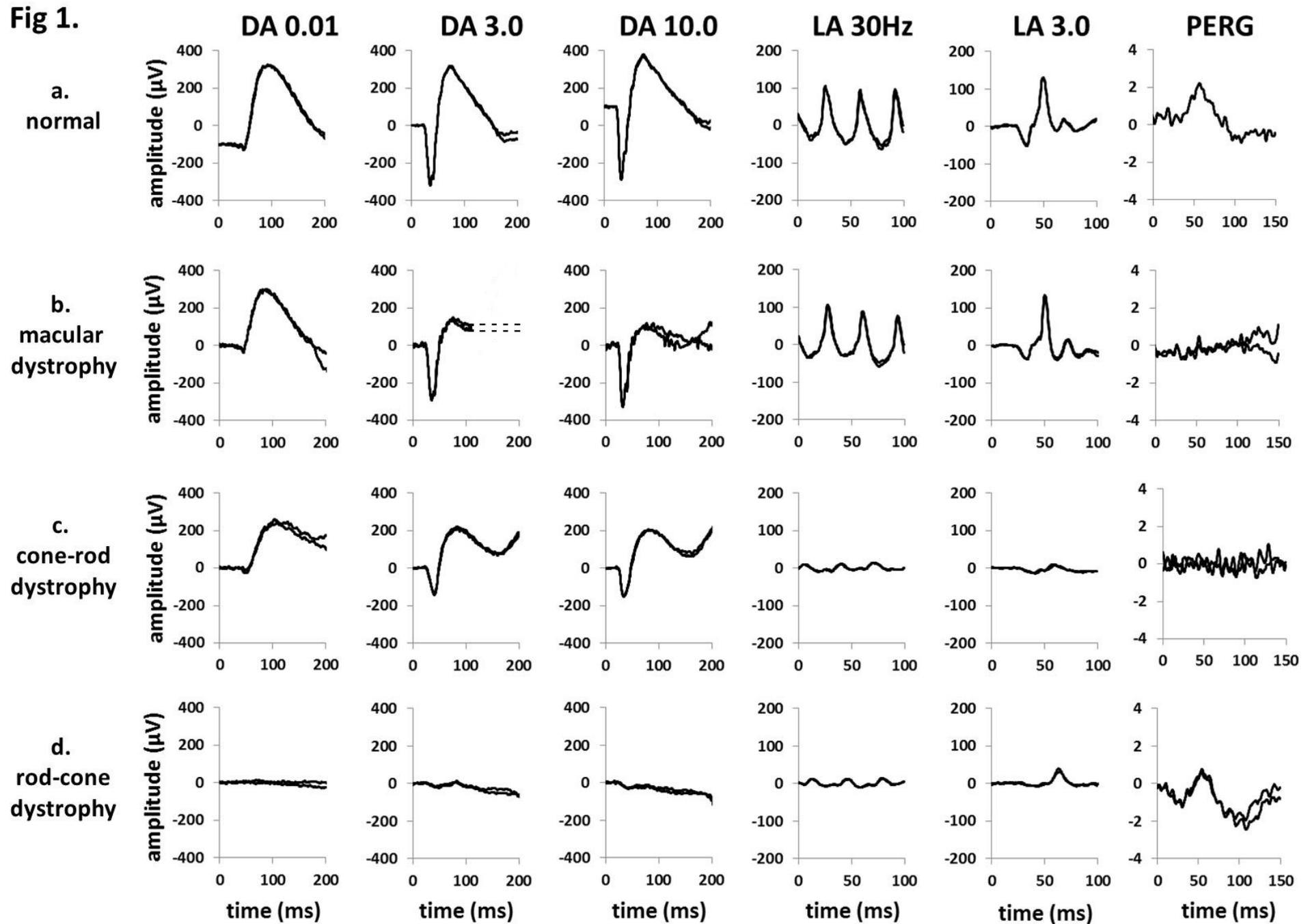
Figure 4. Suggested test strategy for cases of suspected visual pathway dysfunction, illustrating how complementary tests can localize dysfunction within the visual system. ¹In cases of retinal ganglion cell dysfunction the PERG N95:P50 ratio is subnormal but in severe disease P50 may additionally show reduction with shortening of peak time. ²Bestrophinopathies; Best disease is associated with a normal ERG and abnormal EOG; autosomal recessive bestrophinopathy causes severe EOG reduction and later onset progressive retinopathy with relatively mild ERG abnormality. In ADVIRC the EOG is abnormal and the ERG abnormal. See table 1 for details. After; Carr and Siegel (1990) and Holder (2001).

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Fig 1.



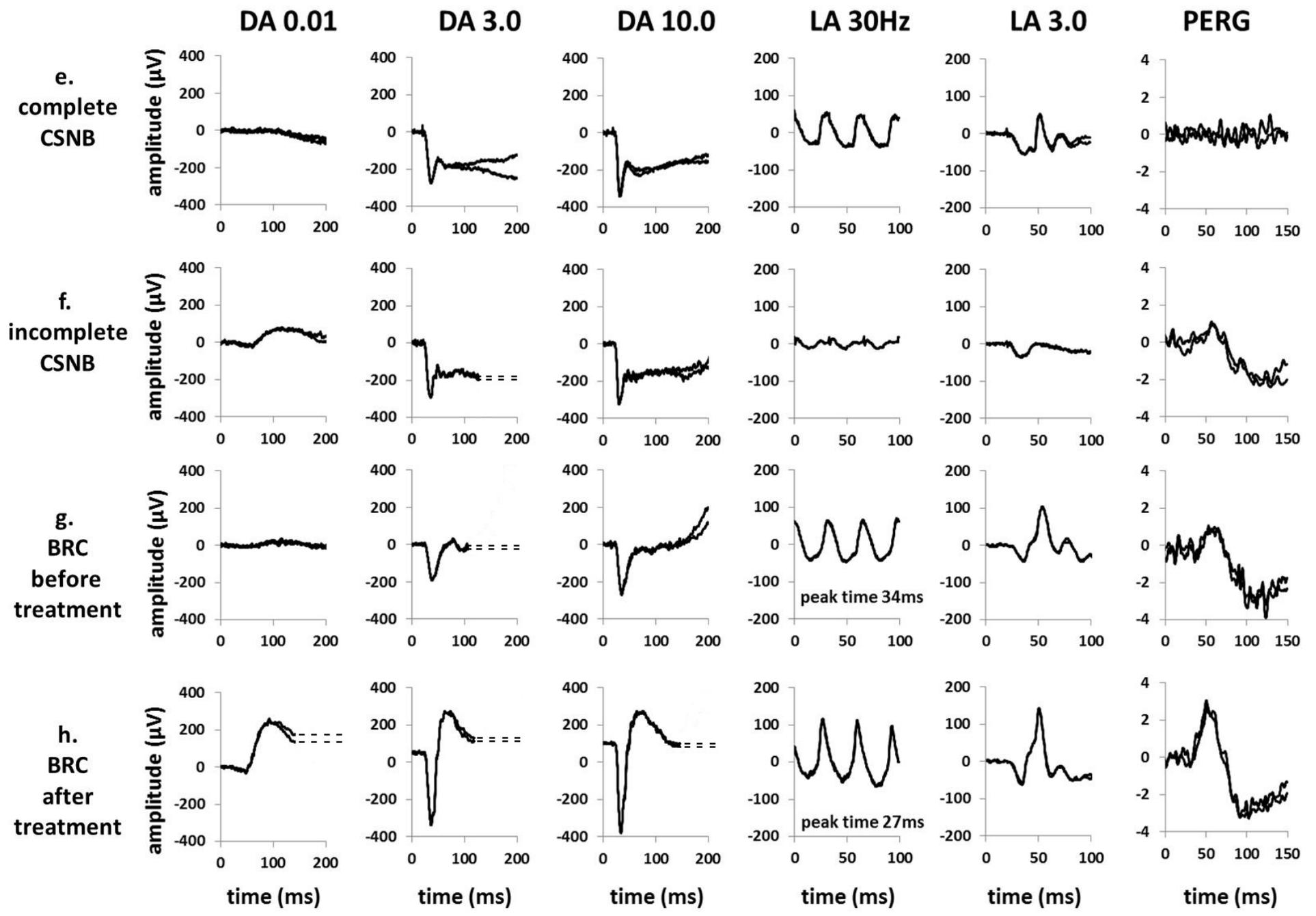
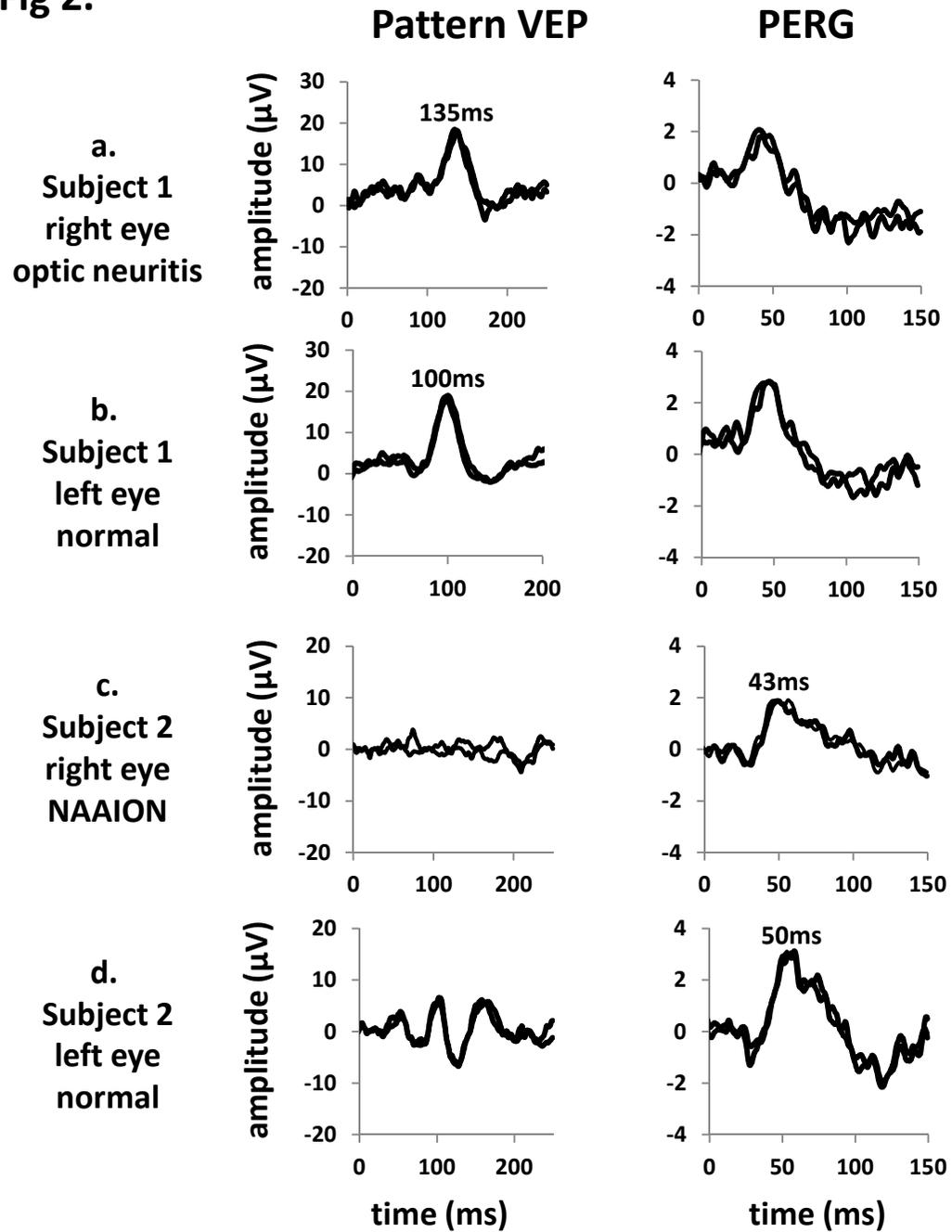
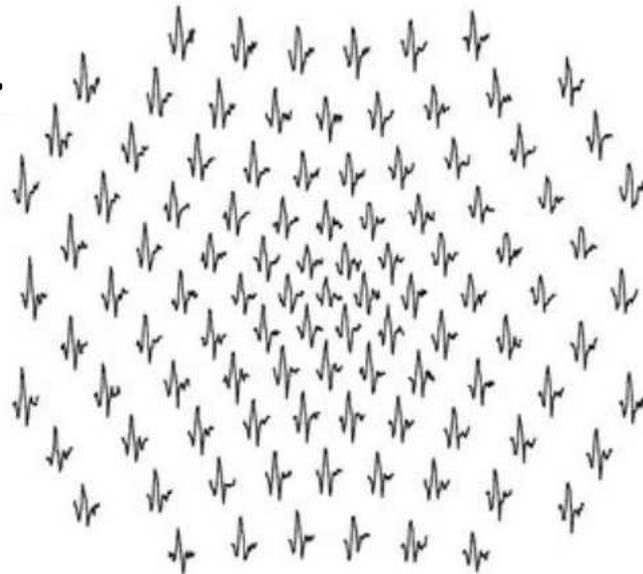


Fig 2.

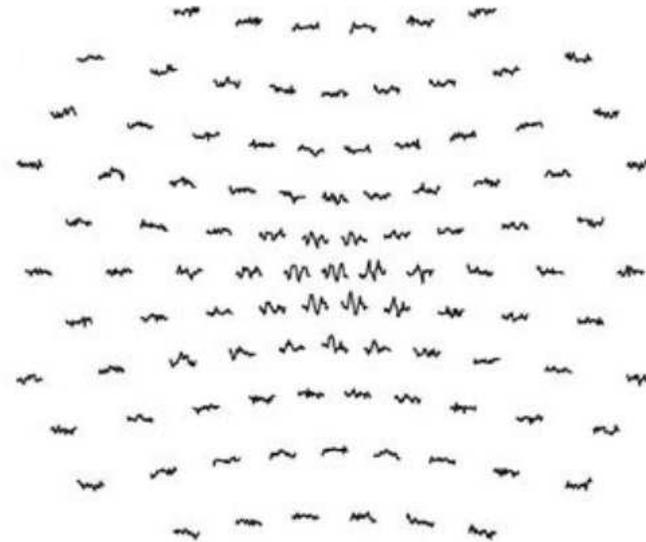


a.



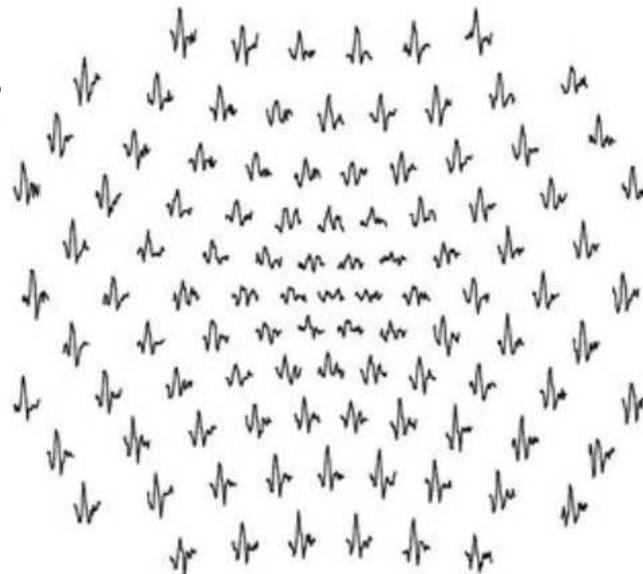
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0 80 ms

b.



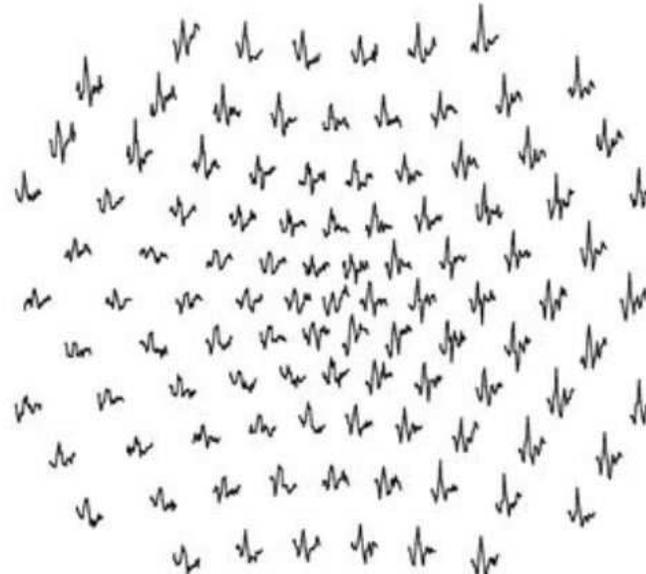
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c.



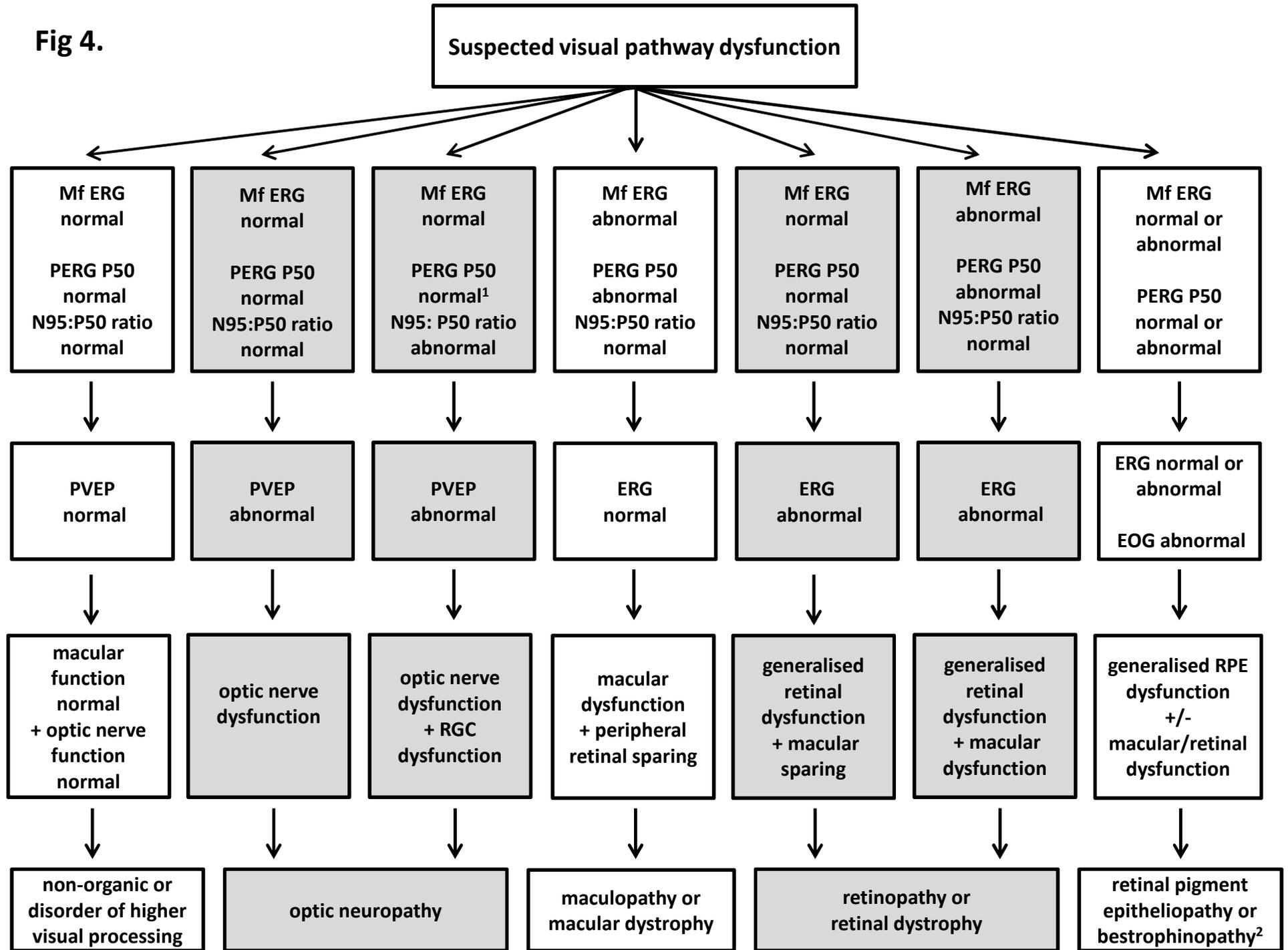
200 nV
0 80 ms

d.



200 nV
0 80 ms

Fig 4.



	Typical or common fundus/ocular abnormalities	Acquired disorder or gene/s implicated ¹	Macular function	Rod system function		Cone system function		Comments including VEP, EOG and other electrophysiological findings where relevant.
			PERG P50 or MfERG	DA 0.01	DA10.0	LA 3.0 30Hz	LA3.0 2Hz	
Adult vitelliform macular dystrophy	Small vitelliform foveal lesion due to a subretinal cyst with or without paracentral drusen and mild RPE changes.	PRPH2, BEST1, IMPG1, IMPG2	N/A	N	N	N	N	The EOG is normal or mildly subnormal and distinguishes most cases from Best vitelliform macular dystrophy.
Albinism	Blonde fundus and foveal hypoplasia are common. There may be iris transillumination and nystagmus.	TYR, OCA2, TYRP1, SLC45A2, GPR143	A	N	N	N	N	Multichannel VEPs show bilateral contralateral predominance to pattern onset-offset (adults) or flash stimulation (children). Assessment of macular function may be precluded by the effects of nystagmus; in the absence of nystagmus there may be evidence of mild macular dysfunction.
Autosomal dominant vitreoretinopathy (ADVIRC)	Liquefied vitreous. Preretinal white dots and neovascularization often present. Peripapillary atrophy can occur. Abnormal pigment often extends to an equatorial demarcation line at the posterior border.	BEST1	A	A	A	A	A	The EOG light peak to dark trough ratio is severely abnormal.
Autosomal recessive bestrophinopathy (ARB)	Diffuse RPE irregularity extending to the vascular arcades associated with patchy RPE atrophy and punctate white dots.	BEST1	A/N	A/N	A/N	A/N	A/N	The EOG light peak to dark trough ratio is severely abnormal. ERG is initially normal but mild abnormalities usually develop in late childhood or adolescence and then worsen progressively.
Batten disease (juvenile onset neuronal ceroid lipofuscinosis)	Normal or Bull's eye lesion	CLN3	A	A	A (-ve)	A	A (-ve)	Electronegative ERG (-ve) may be detected before fundus changes.
Birdshot retinochoroidopathy	Multiple pale sub-retinal lesions. Inflammatory signs such as vitritis, vasculitis and CMO are common.	Acquired	A/N	A/N	A(-ve)/N	A/N	A/N	Variable. Mf ERG and PERG often reveal macular dysfunction, especially if there is CME. ERG is normal to abnormal depending on severity and efficacy of treatment. LA 30Hz ERG commonly delayed and DA strong flash ERGs in some cases have a low b:a ratio before treatment.
Best vitelliform macular dystrophy ("Best disease")	Variable but often characterized by a vitelliform yellow macular lesion due sub-retinal cyst, which may evolve in some to become vitelliruptive with eventual atrophy. Fundus can be normal.	BEST1	N/A	N	N	N	N	The EOG light peak to dark trough ratio is abnormal. Macular dysfunction occurs as macular lesions become vitelliruptive.
Bulls-eye maculopathy	Concentric paracentral changes with foveal sparing.	Acquired or Genetic	A A A	N N A	N N A	N A A+	N A A+	Maculopathy or macular dystrophy Cone dystrophy Cone-rod dystrophy PERG/mfERG evidence of macular dysfunction; mfERG may reveal paracentral dysfunction with localized or relative foveal sparing.
Carcinoma Associated Retinopathy	Fundus initially normal. RPE atrophy, mottling and vessel attenuation may develop.	Acquired	A	A+	A+	A+	A+	Often severe photoreceptor dysfunction causing an undetectable ERG or severe a-wave reduction. Cone system is most affected in some. In rare cases there is an electronegative ERG.

Central retinal artery occlusion	Inner retinal oedema and a cherry red spot at the macula in the early stages. Eventual arteriolar attenuation and disc pallor.	Acquired	A (variable)	A	A (-ve)	A	A	Decreased oscillatory potentials. Relative sparing of visual acuity and of the PERG/central mfERGs if there is a cilioretinal artery.
Central retinal vein occlusion	Dilatation and tortuosity of retinal veins, dot and flame hemorrhages, cotton wool spots, optic disc and macular edema, hyperemia. Ischemic form may result in severe vascular leakage and rubeosis.	Acquired	A (variable)	A	A(-ve)	A	A	Reduced oscillatory potentials. Ischemic CRVO associated with more severe ERG changes and a more reduced DA ERG b:a ratio than non-ischemic disease. ERG a-wave involvement in severe cases.
Choroideremia	Loss of RPE and choriocapillaris. Inner retina and optic disc normal. Late involvement of macula	REP 1	A	A+	A+	A	A	Severe (+) rod>cone or undetectable ERGs. Late macular involvement.
	Female heterozygotes may show mild pigmentary changes or patchy RPE degeneration.		N	N	N	N	N	ERG is usually normal in female heterozygotes but worsening can occur in middle age.
Cone dystrophy	Fundus may be normal. Disc pallor, granular RPE, bull's eye lesion, central atrophy.	see Ret Net (many)	A	N	N	A	A	See text. PERG and MfERG usually show severe and early macular involvement.
Cone-rod dystrophy	Fundus may be normal. Disc pallor, granular RPE, bull's eye lesion, central atrophy.		A	A	A	A+	A+	
CSNB 1.Schubert-Bornschein a)complete a) incomplete 2. Riggs-type	Fundus normal (+/- myopic changes).	NYX, GRM6, TRPM1, LRIT3, GPR179	A	U	A (-ve)	A	A	Fundus are normal; see text for details. See also fundus albipunctatus and Oguchi disease (forms of CSNB with abnormal fundus).
	Fundus normal (+/- myopic changes).	CACNA1F, CABP4	A	A	A (-ve)	A+	A+	
	Fundus normal	PDE6B, RHO, GNAT1, SLC24A1	N	A	A	N	N	
Dominant optic atrophy	Disc pallor typically wedge shaped and temporal but may be diffuse.	OPA1, OPA3, OPA4, OPA5, OPA8	PERG P50 N or mildly subnormal and of short peak time MfERG N	N	N	N	N	PERG N95 may be abnormal in early stages. In severe cases PERG P50 may be reduced with shortening of P50 peak time. Pattern VEP often shows delay and reduction but abnormalities can be mild in the early stages.
Enhanced S-cone syndrome	Normal to nummular pigment clumping in RPE in vicinity of vascular arcades. Macular schisis can occur.	NR2E3	A	U	A	A+	A	Pathognomonic ERG abnormalities. DA3, DA10 and LA3 ERGs are severely delayed with a simplified waveform. LA3 ERG a-wave is larger than the severely abnormal LA 30Hz ERG. S-cone ERG is enlarged.
Fundus albipunctatus	Multiple small white/yellow spots with sparing of the macula.	RDH5	A/N	A	A	A/N	A/N	See text. DA ERGs improve or normalize after prolonged dark adaptation. Approximately 50% have mild LA ERG abnormalities.
Ischemic optic neuropathy	In arteritic form optic disc swelling, disc pallor +/- flame hemorrhages	Acquired	PERG P50 N or mildly subnormal	N	N	N	N	Pattern VEPs show reduction without significant delay. More severe in arteritic than non-arteritic cases. There may be eventual PERG

			and of short peak time MfERG N					N95 reduction in keeping with retinal ganglion cell dysfunction and with reduction/shortening of P50 peak time in some. Usually unilateral.
KCNV2-retinopathy ("cone dystrophy with supernormal rod ERG")	Normal in young but BEM and macular RPE atrophy may develop. Disc pallor in some. Peripheral retina normal.	KCNV2	U	A	A	A	A	Pathognomonic ERG abnormalities. Generalized cone system dysfunction with unusual rod system involvement; DA ERGs to dim flashes are small and delayed and ERG b-waves to strong flashes large. DA 10 ERG a-wave has a distinctive broad trough with a late negative component.
Leber congenital amaurosis	Pigmentary & atrophic changes with age. Hypoplastic/swollen discs common.	see Ret Net (many)	A	A+	A+	A+	A+	ERG typically undetectable or severely reduced from early infancy.
Leber hereditary optic neuropathy	Nerve fibre layer swelling in acute stages. Enlarged or telangiectatic and tortuous peripapillary vessels. Optic atrophy.	G11778A, T14484C, G3460A	PERG P50 N or mildly subnormal and of short peak time MfERG usually N	N	N	N	N	PERG N95 may be abnormal in acute stage. Pattern VEPs are undetectable or severely abnormal. Absence of fluorescein leakage from the swollen disc, distinguishing LHON from other forms of disc swelling. Both eyes
Melanoma Associated Retinopathy	Fundus usually normal. Vitreous cells, vessel attenuation and disc pallor may develop in some.	Acquired	A	U	A(-ve)	A	A	Long duration On-Off ERG shows On response b-wave reduction with sparing of the Off-response. Full-field ERGs identical to those in complete CSNB.
Oguchi disease	Golden fundus sheen which resolves following prolonged dark adaptation (Mizuo-Nakamura phenomenon).	SAG, rhodopsin kinase	N	U	A	N	N	DA ERGs show severe rod dysfunction after 20 minutes in the dark. LA ERGs are normal. After prolonged DA a single strong flash elicits a normal ERG; subsequent flashes elicit subnormal responses and further prolonged DA is needed to recover.
Optic neuritis	Disc pallor and thinning of retinal nerve fibre layer may be evident.	Acquired	PERG P50 N or mildly subnormal and of short peak time MfERG N	N	N	N	N	Pattern VEP is usually delayed with or without amplitude reduction. PERG P50 is usually normal but in 35% cases there is PERG N95 reduction in keeping with retinal ganglion cell dysfunction and with reduction/shortening of P50 peak time in some. May be subclinical involvement of other eye.
Pattern dystrophy	Various patterns of pigment deposition within the macula including adult-onset vitelliform macular dystrophy, butterfly-shaped, reticular, multifocal pattern dystrophies and fundus pulverentulus.	PRPH2	A/N	N	N	N	N	The EOG is normal or mildly subnormal. The ERG is usually normal although there can be marked variability in fundus appearance and ERG phenotype within families with PRPH2 mutation.
Retinitis Pigmentosa (rod-cone dystrophy)	Classically bone-spicule formation, RPE atrophy, attenuated vessels, disc pallor. Normal or near-normal in some.	see Ret Net (many).	A/N	A+	A+	A	A	See text. Rod-cone dystrophy of variable severity. Variable macular involvement. In X-linked pedigrees, female heterozygotes usually have ERG abnormalities with inter-ocular ERG asymmetry. .
Rod monochromacy ("achromatopsia")	Usually normal, macular granularity may develop.	CNGA3, CNGB3, GNAT2, PDE6C, PDE6H, ATF6	A	N	N/ sl. A	U	U	See text. There may be mild reduction in DA strong flash ERGs due to loss of the normal cone system contribution to the a- and b-waves.

Retinal toxicity (selected examples)	Chloroquine/hydroxychloroquine	Acquired	A	N/A	N/A	N/A	N/A	MfERG shows annular macular dysfunction in early stages with later central involvement. ERG abnormal in severe cases.
	Desferrioxamine		A	N/A	N/A	N/A	N/A	Macular dysfunction most common; PERG/mfERG +/- ERG abnormality. ERG may be normal but ranges from showing mild rod dysfunction to severe cone-rod dysfunction.
	Quinine		A	A	A(-ve)	A	A	On response electronegative; Off d-wave has an abnormal shape.
Retinitis punctata albescens (Bothnia dystrophy)	Multiple small white/yellow spots with sparing of the macula. Diffuse RPE degeneration, scalloped peripheral atrophy and pigment deposition in late stages.	RLBP1	A/	A	A	A	A	Resembles fundus albipunctatus in early stages and DA ERGs may show partial improvement after prolonged dark adaptation. Eventual progressive rod-cone dystrophy.
S-cone monochromacy ("X-linked incomplete achromatopsia")	Usually normal, macular granularity may develop.	OPN1LW, OPN1MW	A	N	N/ sl. A	U	A	A preserved S-cone ERG distinguishes the disorder from rod monochromacy. There may be relatively mild reduction in DA 0.01 and DA10 ERG a-waves due to high myopia and loss of cone system contribution to the strong flash ERG a- and b-waves.
Stargardt disease / fundus flavimaculatus (ABCA4-retinopathy)	Central atrophy with flecks or widespread flecks across posterior pole with peri-papillary sparing. Extensive RPE atrophy in severe cases.	ABCA4	A A A	N N A	N N A	N A A	N A A	Macular dystrophy Cone dystrophy Cone-rod dystrophy In all 3 phenotypes there is PERG/mfERG evidence of macular dysfunction
Vitamin A deficiency	Normal or white spots across the fundus	Acquired	N	A	A	N	N	See text.
X-linked retinoschisis	Macular cysts common; may progress to macular atrophy in older men. Peripheral schisis occurs in about 50% of cases.	RS1	A	A	A(-ve)	A	A	On b-wave +/- OFF d-wave subnormal. Inner retinal dysfunction of variable severity. PERG and mfERG usually abnormal.

Table 1. Typical electrophysiological abnormalities in selected retinal and visual pathway disorders. N; normal, A; abnormal, sl. A; slightly abnormal; A+ severely abnormal. U; undetectable. "-ve" signifies an electronegative ERG; b-wave that is smaller than a normal or near-normal a-wave (b:a <1). ¹ See RetNet, the Retinal Information Network for details of disease genes and updates on disease genes; RetNet, <http://www.sph.uth.tmc.edu/RetNet/>