Diagnostic value of nystagmus: spontaneous and induced ocular oscillations

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Research over the past 20 years has provided a clearer understanding of the pathogenesis of most forms of nystagmus and other ocular oscillations. To the clinician, these advances translate into greater accuracy of nystagmus as a diagnostic sign. However, to capitalise on these advances, it is important to systematically examine eye movements and interpret the findings with reference to pathophysiology. In this review we describe a scheme for examining the patient with nystagmus and interpreting common ocular oscillations; some examples are provided as video clips.

Enough is now known about nystagmus to make it a valuable clinical finding, provided that it is characterised systematically, and interpreted in the context of the general neurological evaluation. Nystagmus may be defined as repetitive, to and fro, involuntary eye movements that are initiated by slow drifts. Nystagmus may consist mainly of sinusoidal slow phase oscillations (pendular nystagmus) or, more commonly, of an alternation of slow drift and corrective quick phase (jerk nystagmus). Common waveforms are schematised in figure 1. Thus, each slow phase takes the eye away from the preferred direction of gaze and the corrective quick phase (a saccade) brings the eye back towards the visual target. Although nystagmus is often described by the direction of its quick phases (for example, downbeat nystagmus), it is the slow phase that reflects the underlying disorder. Nystagmus should be differentiated from saccadic intrusions and oscillations, in which rapid movements (saccades) take the eye away from the target (fig 1).

Modern classifications of nystagmus are based on current knowledge of the neurobiology of eye movements, and disorders of the three mechanisms that normally hold gaze still (visual fixation, the vestibulo-ocular reflex, and the eccentric gaze holding mechanism). However, at the bedside, the clinician must rely on the observed characteristics of nystagmus and a systematic examination of eye movements to deduce the significance of any ocular oscillations. Here we describe how to elicit nystagmus and saccadic oscillations, and then summarise the observed characteristics of nystagmus and a systematic examination of eye movements to deduce the significance of any ocular oscillations. Here we describe how to elicit nystagmus and saccadic oscillations, and then summarise the general significance of common forms of nystagmus. To realise the full diagnostic value of nystagmus, whether it interferes with vision or causes oscillopsia (illusory motion of the visual world), and whether it is accompanied by other neurological symptoms. In general, oscillopsia is a feature of acquired, not congenital, nystagmus. Ask if associated visual symptoms are worse when viewing far or near objects, during specific directions of gaze (for example, downgaze), or when the patient is in motion (which implies a vestibular disorder). Try to determine if the nystagmus is congenital by asking about “jumping eyes,” strabismus, eye patching, or eye operations since childhood. Check the patient’s current medications for agents with effects on the brain (for example, anticonvulsants). If the patient habitually tilts or turns the head, determine whether these findings are evident on old photographs, which would suggest congenital nystagmus.

EXAMINATION

Before starting an evaluation of nystagmus, note any abnormality of head posture, examine the visual system (acuity, fields, colour, stereopsis), optic nerves, lids, pupils, and look for signs of ocular albinism. Several congenital forms of nystagmus are associated with disorders of the visual system.

Before examining for nystagmus, determine whether there is a full range of the movements of each eye, and note any static deviation (strabismus). Then observe the stability of gaze as the patient attempts to fix upon a stationary target (such as the letter X) at a viewing distance greater than 2 m. With the patient’s eyes close to central position, determine whether there is any nystagmus. For each eye, note the directions in which the nystagmus occurs: horizontal, vertical, torsional (rotational around the line of sight), or mixed. Compare the nystagmus in each eye, and note whether the direction or size of movement differs, or if there is any asynchrony. If the size of the oscillations differs in each eye, it is referred to as dissociated nystagmus. If the direction of the oscillations in each eye differs, it is called conjugate or disjunctive nystagmus. Cover each eye in turn to check for latent nystagmus (see video “Latent_Nystagmus”). Some nystagmus is intermittent and requires sustained observation over several minutes. Low amplitude nystagmus may only be detected during ophthalmoscopy; note that the direction of horizontal or vertical nystagmus is inverted when viewed through the

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Evaluation of nystagmus is incomplete without a systematic examination of each functional class of eye movements—vestibular, saccades, smooth pursuit, and vergence; selective examination of each functional class of eye movements—sustained convergence.

A characteristic sign of congenital nystagmus is that during testing with a horizontal optokinetic stimulus, the quick phases may be directed in the same direction as drum motion—“optokinetic inversion.” Horizontal nystagmus that changes direction when the eyes are alternatively covered (beating away from the covered eye) is usually “latent nystagmus,” which is associated with lack of normal binocular development (absent stereopsis), and childhood strabismus (see video, “Latent Nystagmus”).

Peripheral vestibular nystagmus (fig 1A) commonly beats horizontally, is commonly congenital. Congenital nystagmus usually remains horizontal in all gaze angles, may be suppressed during convergence (see video, “Congenital_Nystagmus”), and is sometimes associated with anomalous head positions, head oscillations and strabismus. A characteristic sign of congenital nystagmus is that during testing with a horizontal optokinetic stimulus, the quick phases may be directed in the same direction as drum motion—“optokinetic inversion.” Horizontal nystagmus that changes direction when the eyes are alternatively covered (beating away from the covered eye) is usually “latent nystagmus”, which is associated with lack of normal binocular development (absent stereopsis), and childhood strabismus (see video, “Latent Nystagmus”).

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examination and history of lifelong eye problems. However, sometimes it is necessary to record eye movements, because most waveforms of congenital nystagmus are diagnostic, especially increasing velocity waveforms (fig 1C) with superimposed “fvection periods,” when the eyes are momentarily still. The diagnostic significance of these congenital forms of nystagmus to the neurologist are that they usually require no further evaluation. Horizontal nystagmus that regularly reverses direction about every two minutes is periodic alternating nystagmus (PAN); it is associated with disease of the midline cerebellum (nodulus and uvula). To identify this oscillation, which is usually unaffected by visual fixation, an observation of a few minutes is required, and the examiner should also look for periodic alternating head turns that may be associated. A congenital variant of PAN, which is not related with cerebellar disease, usually does not reverse direction at regular intervals.

Vertical jerk nystagmus (downbeat or upbeat), as well as purely torsional jerk nystagmus, with the eye near to central position is usually attributable to a central involvement of the vestibular pathways. Both downbeat and upbeat nystagmus are poorly suppressed by visual fixation and may be exacerbated by simply placing the patient in a head hanging position; they may also be increased, suppressed, or inverted by convergence. Typically downbeat nystagmus is best evoked on looking down and laterally (see video “Downbeat_Nystagmus”) whereas upbeat nystagmus generally increases on looking up. Downbeat nystagmus is usually encountered with lesions of the vestibulocerebellum, craniovascular anomalies, and drug intoxications; it is also a feature of the calcium channelopathy, episodic ataxia type II. Upbeat nystagmus is less well localised, being mainly reported with lesions in the medulla, or close to the superior cerebellar peduncle (see video “Upbeat_Nystagmus”).

Pure torsional nystagmus with the eye close to central position is uncommon and is usually associated with medullary lesions, such as syringobulbia and lateral medullary infarction (Wallenberg’s syndrome). It is often accompanied by ocular tilt reaction or unilateral internuclear ophthalmoplegia.

Acquired pendular nystagmus has a quasi-sinusoidal waveform (fig 1D), and often shows trajectories that are oblique, elliptical, or circular, depending on the size and temporal relation between horizontal and vertical components. Congenital pendular nystagmus is usually horizontal in direction. Acquired pendular nystagmus may be conjugate, disconjugate (dissimilar trajectories of each eye), or dissociated (different size of oscillations in each eye). It occurs most commonly in association with disorders of central myelin, especially multiple sclerosis (see video, “Pendular_Nystagmus”), and after brain stem stroke as part of the syndrome of oculopatalol tremor (“myoclonus”).

Pendular oscillations that are about 180° out of phase in the horizontal plane causes a type of convergent-divergent nystagmus, which occurs in some patients with cerebral Whipple's disease, sometimes with associated oscillatory movements of the jaw, face, or limbs (oculomasticatory myorhythmia) and vertical gaze palsy. Pendular convergent-divergence nystagmus has also been reported in patients with multiple sclerosis and brain stem stroke. Convergence-divergence nystagmus should be differentiated from conjugate nystagmus that is affected (suppressed, increased, or inverted) by convergence, a feature of a variety of acquired forms of nystagmus (for example, downbeat, upbeat nystagmus). Congenital nystagmus is often suppressed during convergence.

Seesaw nystagmus (pendular or jerk) consists of elevation and intorsion of one eye and synchronous depression and extorsion of the other eye in the first half cycle, followed by change in direction during the next half cycle. To appreciate the seesaw nature of the oscillation, it often helps to look at the bridge of the patient's nose. Pendular seesaw nystagmus occurs with disorders that interfere with crossing axons at the optic chiasm and cause bitemporal hemianopia, such as pituitary tumours, as well as a variant of congenital nystagmus (see video “Seesaw_Nystagmus”). Jerk seesaw nystagmus is reported in patients with lesions in the region of the interstitial nucleus of Cajal.

Nystagmus induced by moving the eyes to an eccentric position—gaze evoked nystagmus

The commonest form of nystagmus encountered in clinical practice occurs only when the eyes are moved into eccentric gaze, especially in lateral and up gaze (fig 1B). This gaze evoked nystagmus has quick phases that are directed away from central position (see video, “Gaze&Rebound_Nystagmus”). Holding the eyes in such eccentric positions requires a tonic contraction of the extracocular muscles, which is achieved by a sustained discharge of motoneurons. This ability is impaired by cerebellar and brainstem disorders as well as a number of intoxications. Thus, the finding of nystagmus on lateral or upgaze may be clinically important. Firstly, however, it is important to differentiate between pathological gaze evoked nystagmus and its variant, called end point nystagmus, frequently reported in normal subjects. The latter is usually unsustained, of low frequency and amplitude, and not accompanied by other ocular motor abnormalities. Pathological gaze evoked nystagmus may be attributable to a peripheral process, being a sign of extraocular muscle weakness, for example, in patients with myasthenia gravis (“fatigue nystagmus”). It also occurs as a result of central disorders that involve the gaze holding neural network, which includes the nucleus prepositus hypoglossi and medial vestibular nucleus for horizontal gaze, the interstitial nucleus of Cajal for vertical gaze, and the vestibulocerebellum, which optimises gaze holding. Structural lesions affecting this gaze holding network may cause gaze evoked nystagmus, because the eyes tend to drift back to central position (fig 1B). Most commonly, gaze evoked nystagmus is encountered as a side effect of medications, including sedatives and anticonvulsants, or with drug intoxications (for example, alcohol). When patients with gaze evoked nystagmus attempt to hold eccentric gaze for a number of seconds, and then return their eyes to the central position, a transient rebound nystagmus, with quick phases opposite to the direction of the prior eccentric gaze, may occur (see video “Gaze&Rebound_Nystagmus”). Such rebound nystagmus is prominent in patients with disease affecting the vestibular cerebellum.

When nystagmus on lateral gaze is greater in the abducting eye, the cause may be internuclear ophthalmoplegia, and saccades should be tested, looking for slowing of the adducting eye. In most cases, dissociated nystagmus is a manifestation of the brain's attempt, with a series of saccades, to put the fovea of the weak, abducting eye on target. Tumours of the cerebellopontine angle may cause a combination of low frequency, large amplitude horizontal nystagmus looking ipsilaterally, because of defective gaze holding, and a high frequency, small amplitude nystagmus looking contralaterally because of vestibular imbalance (Bruns' nystagmus).

One form of nystagmus induced by upgaze is convergence-retraction nystagmus. It occurs with dorsal midbrain lesions (for example, pineal tumours). Convergence-retraction nystagmus may be elicited either by asking the patient to make an upward saccade, or by using a handheld optokinetic drum or tape, moving the stripes down. In some cases the convergent movements appear to be opposed vergence movements, but in others it consists of opposed, asynchronous saccades.

Nystagmus (either jerk or pendular, binocular or monocular) occurs commonly in patients with visual impairment or loss, who cannot accurately direct their gaze. In such patients, both horizontal and vertical components are present, and the
direction of nystagmus fluctuates during a period of observation. Their nystagmus probably reflects an uncalibrated ocular motor system.22

**Saccadic intrusions and oscillations**

Several types of inappropriate saccadic movements may intrude upon steady fixation; more common variants are schematized in figure 1E–G. Because saccadic intrusions are rapid and brief, it is usually necessary to measure eye and target position and eye velocity to identify accurately the saccadic abnormality.

Square wave jerks (fig 1E) occur in healthy subjects,23 but are a prominent finding in progressive supranuclear palsy and some spinocerebellar atrophies (especially Friedreich’s ataxia).24 They are small, conjugate saccades, ranging from 0.5 to 5.0 degrees in size, which take the eye away from the fixation position and then return it there after a period of about 200 ms. They may be most evident during smooth pursuit, and are easily detected during ophthalmoscopy.

Macrosaccadic oscillations (fig 1F) usually consist of horizontal saccades that occur in bursts, building up and then decreasing in amplitude, with intersaccadic intervals of about 200 ms.25 These oscillations are usually induced by a gaze shift. They are a sign of midline cerebellar disease (affecting the fastigial nucleus), including spinocerebellar degenerations (see video, “Macrosaccadic oscillations”),26 but have also been reported with pontine lesions.27

Saccadic pulses are brief intrusions that take the eye movement away from the fixation position, with a rapid drift back. They are reported in patients with internuclear ophthalmoplegia.28 There is a continuum between saccadic pulses and saccadic oscillations without an intersaccadic interval (fig 1G). The latter may occur in the horizontal plane, ocular flutter,29 or may consist of saccadic oscillations with horizontal, vertical, and torsional components, opsoclonus (see video, “Opsoclonus”). The frequency of oscillations is usually typically 10 to 15 cycles per second, being higher with smaller size movements. Flutter and opsoclonus are often brought out by eye lid closure or during gaze shifts between far and distant targets.30 Most cases of flutter and opsoclonus conform to four main diagnostic groups: parainfectious brainstem encephalitis (including multiple sclerosis),31 paraneoplastic syndromes, metabolic-toxic states, or idiopathic. Differentiation between these disorders depends on the general neurological findings and laboratory testing.32

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