Neuro-ophthalmologic aspects of multiple sclerosis: Using eye movements as a clinical and experimental tool

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Abstract: Ocular motor disorders are a well recognized feature of multiple sclerosis (MS). Clinical abnormalities of eye movements, early in the disease course, are associated with generalized disability, probably because the burden of disease in affected patients falls on the brainstem and cerebellar pathways, which are important for gait and balance. Measurement of eye movements, especially when used to detect internuclear ophthalmoplegia (INO), may aid diagnosis of MS. Measurement of the ocular following response to moving sinusoidal gratings of specified spatial frequency and contrast can be used as an experimental tool to better understand persistent visual complaints in patients who have suffered optic neuritis. Patients with MS who develop acquired pendular nystagmus often benefit from treatment with gabapentin or memantine.

Keywords: multiple sclerosis, eye movements, ocular motor disorders

Introduction

No evaluation of a patient either suspected or diagnosed with MS is complete without a systematic examination of their vision and eye movements (Leigh and Zee 2006). Although disturbances of vision and eye movements are recognized as important for making the diagnosis of MS, the neuro-ophthalmologic examination also provides insights into the nature of the disorder in any patient and even in estimating prognosis (Frohman et al 2005). Here we review four aspects of the role of eye movements in MS: (1) the value of clinical examination of eye movements in making the diagnosis and estimating the prognosis in MS; (2) how measurement of eye movements may contribute to the diagnosis of MS; (3) how eye movements can be used as an experimental tool to investigate the visual disturbance due to optic neuritis; and (4) current concepts of treatment of acquired forms of nystagmus due to MS and their visual consequences.

Clinical examination of eye movements

A systematic examination of eye movements in patients with MS shows a range of disorders (Table 1). The most common eye movement abnormalities are saccadic dysmetria, internuclear ophthalmoplegia (INO), disorders of the vestibulo-ocular reflex (VOR), and gaze-evoked nystagmus (Downey et al 2002; Serra et al 2003; Derwenskus et al 2005). Saccades are the rapid, brief eye movements by which we shift the point of fixation from one feature of interest to the next. Normal subjects show mild undershoots (hypometria), especially for larger saccades; however, with repetitive saccades, the eye lands on the visual target. Patients with MS may show persistent dysmetria, including overshoots (hypermetria) and directional dysmetria (eg, an inappropriate vertical component during a horizontal saccade, requiring...
In normal subjects, saccades of each eye are conjugate vertically, but are mildly disjunctive horizontally, with the abducting eye initially leading (producing transient divergence) and the adducting eye eventually catching up (causing transient convergence) (Collewijn et al 1988). This minor, transient disconjugacy of horizontal saccades in normal subjects cannot be discerned at the bedside. However, in INO, the adducting movement is visibly slowed (adduction lag) compared with the abducting movement. Slowing of adduction occurs because the pulse of innervation that causes rapid contraction of the medial rectus muscle is deficient, since the demyelinated axons of the medial longitudinal fasciculus (MLF) cannot conduct the high-frequency discharge (Leigh and Zee 2006). In classic INO, the findings include restricted adduction and adduction lag in the eye ipsilateral to the MLF lesion and dissociated abducting “nystagmus” of the contralateral eye (which is discussed in the last section). Milder forms of INO cause adduction lag but no restriction of adduction or abducting “nystagmus” (Crane et al 1983; Müri and Meienberg 1985). This adduction lag is most easily visualized with rapid eye movements; thus, INO is best detected by testing saccades, and may be missed if only pursuit movements are examined (“follow my finger”).

Carefully conducted clinical studies have established that patients who have abnormal eye movements are more disabled than those with normal motility, and that this difference is sustained after two years (Downey et al 2002; Serra et al 2003; Derwenskus et al 2005). The reason for the increased disability in MS patients with abnormal eye movements is less due to their impaired vision than to their disturbance of gait. In such patients, MRI scans confirm that both abnormal eye movements and gait disturbance are caused by prominent involvement of the brainstem and cerebellar pathways.

### Measuring eye movements to aid diagnosis of MS

Clinical detection of INO and saccadic abnormalities in MS may be challenging (Frohman TC et al 2003), and measurement of eye movements may help confirm the diagnosis during early stages of the disease. A representative example is shown in Figure 1.

Detection of a saccadic abnormality may be better when targets are presented randomly, so that neither their time of onset nor their location can be predicted. Large saccades (20 degrees or greater) are more likely to show changes in velocity than are small saccades. In making paired comparison of the peak velocity of abducting and adducting saccades, it is important to take account of the fact that normal subjects show somewhat greater peak velocities in the abducting eye, depending on the methodology being used (Collewijn et al 1988). A more reliable approach is to measure the ratio of abducting/adducting movements during horizontal saccades, and compare the patient’s value with that of group of normal subjects recorded in the same laboratory (Flipse et al 1996; Frohman et al 2002; Frohman EM et al 2003). Patients with INO have abduction/adduction ratios for peak velocity or

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### Table 1 Summary of common disorders of ocular motility in MS (Downey et al 2002)

| Abnormal finding                      | Occurrence in MS patients who show abnormal eye movements |
|---------------------------------------|----------------------------------------------------------|
| Misalignment of visual axes           | 32% (usually exotropia)                                  |
| Gaze-evoked nystagmus                 | 36%                                                      |
| Nystagmus in central position         | 18% (pendular; downbeat)                                 |
| Saccadic dysmetria                    | 91% (59% in both planes)                                 |
| Internuclear ophthalmoplegia          | 68% (27% bilateral)                                     |
| Smooth pursuit impaired               | 32%                                                      |
| Vestibulo-ocular reflex impaired      | 36%                                                      |
| Vergence impaired                     | 23%                                                      |

Abbreviation: VOR, vestibulo-ocular reflex.
Bilateral internuclear ophthalmoplegia impairs the vertical vestibulo-ocular reflex, gaze holding, smooth pursuit, and eye-head tracking— all functions that also depend on the medial longitudinal fasciculus (Ranalli and Sharpe 1988). Patients with head tremor and an impaired vestibulo-ocular reflex may report oscillopsia—illusory motion of the visual world (Proudflock et al 2002).

Less commonly, MS may cause horizontal and vertical gaze palsies (Wall and Wray 1983; Martyn and Kean 1988; Tan and Kansu 1990; Milea et al 2001), or individual oculomotor, trochlear or abducens palsies due to brainstem plaques (Leigh and Zee 2006), upbeat and downbeat nystagmus (Fisher et al 1983; Masucci and Kurtzke 1988; Baloh and Yee 1989), and a range of vestibular and optokinetic abnormalities (Katsarkas 1982; Huygen et al 1990; Todd et al 2001). MRI often aids identification of brainstem or cerebellar lesions responsible for such abnormalities; thin cuts and proton density images may be required.

Diagnosis of multiple sclerosis depends on demonstration of lesions disseminated throughout the nervous system (Frohman et al 2005). Early diagnosis has become more important because prompt initiation of immunotherapy has the best chance of arresting the disease. Although detection of subtle deficits of ocular motility, such as INO, may be useful in making a diagnosis, some caution is required since these tests are not specific for multiple sclerosis. Thus, the clinician must weigh the results of ocular motor studies with other clinical or laboratory findings before making a diagnosis in the context of current criteria (Dalton et al 2002). Routine examination of eye movements during follow-up visits to the MS clinic (along with careful evaluation of vision) may provide a useful method of evaluating progress and prognosis. As noted above, patients who are more generally disabled are more likely to develop abnormalities of eye movements, probably because the brunt of the disease is affecting brainstem and cerebellar connections (Downey et al 2002; Serra et al 2003; Derwenskus et al 2005).

**Application of eye movements to investigate visual disorders in MS**

The signature disturbance of vision in MS is optic neuritis, which typically presents as monocular loss of central vision, accompanied by discomfort with eye movements (Frohman et al 2005). Thanks to the longitudinal Optic Neuritis Treatment study, we now have a more detailed knowledge of the clinical characteristics and course of the syndrome, in which over 95% of patients recover 20/40 or better vision.
by 12 months after the onset of symptoms (Beck and Cleary 1993; Atkins et al 2006).

Some patients who suffer optic neuritis make a good recovery of vision as tested by high-contrast Snellen optotypes but show residual visual impairment, particularly with contrast sensitivity and with low-contrast optotypes such as Sloan letters (Balcer et al 2003). Color vision may also be asymmetrically impaired in such patients.

We recently investigated whether motion vision could remain compromised after recovery from optic neuritis in a 42-year-old man with MS (Rucker et al 2006). Three years after left optic neuritis had reduced his vision to 20/40 OS (20/20 OD) with Sloan letters but remained at 20/200 OS (20/32 OD) with Sloan letters at 10% contrast. Color vision was normal OD but essentially absent OS. Despite this improvement, he reported subjective visual difficulties OS, which he noted while he was in motion. Rather than perform a subjective test of motion vision, we measured his ocular following response (OFR). The OFR is a measure of first-order, luminance driven motion vision which consists of a sudden conjugate eye movement elicited at ultra-short latencies by rapid shift of a large visual stimulus (Sheliga et al 2005). One advantage of studying motion vision by measuring the short-latency OFR is that this eye-tracking response is rather automatic and not substantially influenced by the level of attentiveness of the subject. The stimuli we applied consisted of ¼-cycle shifts of a sine-wave grating with a spatial frequency of either 0.153 or 0.458 cycles/degree. We found that his responses to the higher spatial frequency stimulus were impaired in the affected left eye, suggesting predominant post-optic neuritis involvement of parvocellular visual function (Figure 2). Responses to the low spatial frequency stimulus were similar between the two eyes and to controls, suggesting preservation of the magnocellular visual function, which may function as a visual field-holding reflex (Miles 1998). Although OFR has been applied to patients with cortical disturbances of vision, this preliminary study suggests that OFR may be a valuable, objective means of further study of visual dysfunction following optic neuritis.

**Acquired nystagmus in MS and treatment of its visual consequences**

Several forms of nystagmus occur in MS. So-called dissociated nystagmus with INO is more correctly classified as a disorder of saccadic metrics: the adducting eye fails to get on target, and the brain then programs a series of corrective saccades that are not evident in the paretic, adducting eye, but are observed in the abducting eye (Leigh and Zee 2006). Patching the affected eye in unilateral INO for several days reduces “dissociated nystagmus” while patching the normal eye increases it, indicating that plastic-adaptive mechanisms contribute to the nystagmus (Zee et al 1987).

Although upbeat, downbeat, and seesaw nystagmus all may occur in MS, it is acquired pendular nystagmus (APN) that is most visually disabling. APN is also a feature of other disorders of central myelin, including Pelizaeus-Merzbacher disease (Trobe et al 1991), peroxisomal assembly disorders (Kori et al 1998), and in toluene abuse (Maas et al 1991). Since patients with MS who have pendular nystagmus often also have evidence of optic nerve demyelination, it follows that prolonged response time of the visual processing might be responsible for the ocular oscillations. Support for this hypothesis comes from the observation that oscillations are usually larger in the eye with evidence of more severe optic nerve demyelination (Barton and Cox 1993). However, the nystagmus often remains unchanged in darkness (when visual inputs have no influence on eye movements). Furthermore, spontaneous ocular oscillations can be induced in normal subjects by experimentally delaying the latency of visual feedback during fixation; however, the frequency of these induced oscillations is less than 2 Hz, which is lower than in...
most patients with pendular nystagmus due to multiple sclerosis (Averbuch-Heller et al. 1995). Furthermore, when this experimental technique was applied to patients with acquired pendular nystagmus, it did not change the characteristics of the nystagmus, but instead superimposed lower-frequency oscillations similar to those induced in normal subjects (Averbuch-Heller et al. 1995). Thus, disturbance of visual fixation due to visual delays cannot wholly account for the high-frequency oscillations that often characterize acquired pendular nystagmus.

An important observation concerning acquired pendular nystagmus in multiple sclerosis is that some patients show transient suppression of their oscillations following a saccade (Figure 3). Systematic comparison of the oscillations prior to, and following, a saccade has demonstrated that the oscillations are phase-shifted (reset) leading to the hypothesis that the oscillations of APN in patients with multiple sclerosis arise in the neural integrator for eye movements, and that large saccades affect the timing of the oscillations by “resetting” the integrator with the large pulse of neural activity (Das et al. 2000). The nucleus prepositus hypoglossi and medial vestibular nucleus are important components of the neural integrator, which is probably affected by feedback of signals via cell groups of the paramedian tracts (PMT) to the cerebellar flocculus (Leigh and Zee 2006). Patients with APN show a preponderance of multiple sclerosis plaques in their paramedian pons in the region of the PMT cell groups (Lopez et al. 1995). Thus, it is postulated that the neural integrator loses normal feedback, becomes unstable, and begins to oscillate, causing acquired pendular nystagmus. The neurotransmitters used by neurons that contribute to normal neural integration of ocular motor signals are known to include glutamate and gabapentin (Straube et al. 1991; Arnold et al. 1999). This has led to clinical trials that have demonstrated gabapentin (Averbuch-Heller et al. 1997; Bandini et al. 2001), and memantine (Starck et al. 1997) may ameliorate the visually disabling acquired pendular nystagmus that occurs in multiple sclerosis. Our personal experience is that most patients with APN due to MS visually benefit from either gabapentin or memantine. Both drugs are usually well tolerated, although gabapentin sometimes increases ataxia.

**Summary**

In the past twenty years, eye movements have moved beyond the bounds of neuro-ophthalmology and are now widely used by neuroscientists, visual scientists, and psychologists, as well as clinicians in several specialties. The popularity of eye movements as an experimental tool arises, in part, because they can be conveniently and accurately measured and analyzed, but also because much is known concerning their neural substrate. It is therefore, no surprise, that eye movements have been commonly applied to better understand the visual and motor disorders in patients with MS. In this review we draw attention to how eye movements can aid diagnosis of MS at the bedside and in the laboratory. In addition, eye movements can be used as an experimental tool to better understand the disorders of vision that occur in MS. Finally, better knowledge concerning neuropharmacology has led to effective drug treatments for abnormal eye movements that degrade vision in patients with demyelinating disease.

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