Abnormal Visual Adaptation to Flicker in Multiple Sclerosis

J.E. Raymond

ABSTRACT: A visual psychophysical adaptation procedure was used on patients with Multiple Sclerosis (MS) in an attempt to induce a temporary and local exacerbation of subclinical visual impairment. Using a flicker detection task, sensitivity before and after adaptation to a flickering stimulus was measured in 9 MS patients and 9 control subjects. Although only 22% of patient eyes had abnormal flicker sensitivity prior to adaptation, visual deficit was observed in 83% of eyes studied after adaptation. Of the 7 MS eyes studied for which no other sign or symptom of visual involvement was present, 5 were found to have visual deficits after flicker adaptation. In addition, 10 of the 11 eyes affected by MS showed an abnormal response to flicker adaptation. Recovery from the effects of adaptation was complete in all patients within 2 minutes. The results suggest that partial demyelination of visual pathway neurons may exist in patients without signs or symptoms of visual involvement. The prolonged stimulation provided during adaptation may produce a temporary fatiguing or conduction blockade of such neurons which may lead to reductions in sensory sensitivity.

RéSUMÉ: Adaptation visuelle anormale à la stimulation lumineuse intermittente dans la sclérose en plaques

Nous avons utilisé une technique d'adaptation visuelle psychophysique chez des patients atteints de sclérose en plaques (SEP) dans le but d'induire un exacerbation temporaire et locale de l'atteinte visuelle subclinique. A l'aide d'une épreuve de détection du papillotement, nous avons mesuré la sensibilité avant et après adaptation à une stimulation lumineuse intermittente chez 9 patients atteints de SEP et 9 sujets contrôles. Même si seulement 22% des yeux des patients avaient une sensibilité anormale à la stimulation lumineuse intermittente avant adaptation, un déficit visuel a été observé chez 83% des yeux étudiés après adaptation. Parmi les 7 yeux SEP étudiés qui ne manifestaient aucun signe ou symptôme d'atteinte visuelle, 5 présentaient des déficits visuels après adaptation à la stimulation lumineuse intermittente. De plus, 10 des 11 yeux atteints de SEP présentaient une réponse d'adaptation anormale. La récupération des effets de l'adaptation était complète chez tous les patients en dedans de 2 minutes. Ces résultats suggèrent qu'une démyélinisation partielle des neurones des voies de transmission de l'influx visuel peut exister chez des patients ne manifestant aucun signe ou symptôme d'atteinte visuelle. La stimulation prolongée fournie pendant l'adaptation peut produire une fatigue temporaire ou un blocage de la conduction de tels neurones, pouvant induire une diminution de la sensibilité sensorielle.


Extensive demyelination of neurons in the primary visual pathways is almost universally found in patients with advanced multiple sclerosis (MS) who are examined at autopsy.1,2 However, investigations of visual disorders in MS patients have revealed abnormalities in only 30 per cent of patients when conventional testing procedures (e.g., Snellen acuity, static perimetry, fundoscopic examination) are used.3 In view of these findings, it is likely that visual system pathology is present in a considerable proportion of MS patients during life but that it is not always manifested as clinically evident fundoscopic changes or as obvious disturbances in visual function. Recently, psychophysical techniques for measuring visual function have been used experimentally as a means of detecting subtle visual disturbances in MS patients.

One class of such visual tests have employed temporal visual discriminations in an effort to demonstrate abnormally slowed or delayed neural transmission in the visual pathways of MS patients. For example, in double flash4 and multi-flash5 campimetry, subjects view a test spot at various locations in the visual field and are asked to judge whether they perceive a single illumination of a test spot or multiple successive flashes. The duration between successive flashes of light is lengthened until the subject reports the perception of flicker. Using these techniques, it was found that a large proportion of MS patients with a history of optic neuritis could tolerate substantially longer inter-stimulus intervals before flicker perception was reported than healthy control subjects. These results are consistent with neurophysiological data on experimentally demyelinated neurons which suggest that the temporal order of neural signals transmitted by demyelinated axons is under considerable temporal disarray. However, abnormal multi-flash campimetry results may also be found in patients with amblyopia, cataract, macular degeneration and optic neuritis not associated with demyelinating disease.6
Other tests based on flicker perception which have reported abnormal visual function in MS include techniques which assess critical flicker fusion frequency,\textsuperscript{7,8} the Pulfrich illusion,\textsuperscript{9} differences of visual latencies in the two eyes,\textsuperscript{10} DeLange functions,\textsuperscript{11} temporal contrast sensitivity functions\textsuperscript{12} and temporal frequency discrimination functions.\textsuperscript{13} In all of these reports, the MS patient samples consisted of individuals with previously known visual signs or symptoms. Thus, while these psychophysical tests have proved successful at detecting visual system involvement in substantial proportions of carefully selected patients, it is not known whether they are successful at revealing visual system damage in patients with subclinical visual involvement.

In the earlier stages of MS, when diagnosis is a critical issue, lesions may be too small to result in measurable deficits of visual function. Techniques which could temporarily exacerbate the effect of such lesions so as to render symptoms psychophysiologically observable could be potentially useful as diagnostic procedures. For example small elevations in body temperature\textsuperscript{14} or prolonged physical exercise\textsuperscript{15} has been reported to produce temporary deficits in vision in MS patients.

Research on demyelination in animals has shown that a temporary increase in the probability of complete axonal conduction blockade can also be produced by stimulation of a partially demyelinated neuron with high frequency trains of electrical impulses.\textsuperscript{16,17} This suggests that excessive stimulation of visual neural units through prolonged exposure to specific suprathreshold visual stimuli (i.e., adaptation) might increase conduction blockade in MS patients, thus leading to a temporary worsening of visual symptoms. Unlike elevation in body temperature, the temporary exacerbation of symptoms would be local to the visual system and not cause a generalized discomfort to the patient.

Visual adaptation (e.g., to high contrast grating patterns) has been shown to produce small transient changes in visual (e.g., contrast) thresholds obtained from healthy subjects (e.g., 18). These well-studied psychophysical adaptation effects demonstrate the impact prolonged exposure to specific stimuli has on the healthy visual system and have been useful for gaining an understanding of visual mechanisms. If the visual pathways of MS patients are only partially demyelinated, then visual adaptation procedures may be particularly successful at revealing the visual pathology that is undetectable using conventional techniques. Given the physiological data on the effects of demyelination on neural transmission and the psychophysical data on flicker perception abnormalities in MS patients, adaptation techniques involving temporal discriminations may be particularly successful at revealing subtle visual disturbance in MS.

Using healthy observers, Smith\textsuperscript{19} reported that after prolonged viewing of a highly modulated flickering disk, the minimum depth of modulation needed to detect flicker of the same frequency as the adapting stimulus was elevated relative to that required before adaptation. The present experiment investigated this adaptation effect in MS patients. The results indicate that relative to healthy controls, MS patients undergo abnormally rapid and large losses in flicker sensitivity during adaptation.

**METHODS**

**Patients and Control Subjects**

The control group consisted of 5 females and 4 males chosen from university students and staff whose ages ranged from 20 to 28 years (mean = 23.1 years). All subjects had 20/30 acuity or better in both eyes and no history of neurological disorder. Control subjects were paid an hourly fee. The MS group was composed of 9 patient volunteers (8 females and 1 male) ranging in age from 21 to 53 years (mean = 36.3). All patients were diagnosed "definite" MS according to the criteria set out by Schumacher et al.\textsuperscript{20} and were rated on the disability status scale of Kurtzke.\textsuperscript{21} Clinical data for the patients are listed in Table I. All patients had good manual dexterity and experienced no difficulty manipulating the knob used to control flicker in the display.

Patients were assessed for visual acuity, optic disk colour, pupillary defects and, in most cases, contrast sensitivity and double flash sensitivity. Two of the nine patients had no sign or symptom of visual involvement in either eye. Three patients had evidence of monocular involvement and the remaining four showed bilateral involvement. None of the patients were on medication at the time of the experiment.

**Apparatus and Stimuli**

Flickering stimuli were generated by light emitting diodes (LEDs) driven by custom electronics. A single Wein bridge electronic oscillator sinusoidally modulated the luminance of both the adapting and test stimuli. LEDs were used because they have a rapid response to input current. A tightly packed matrix of 13 LEDs attached to the end of a 105 cm long mirror box and viewed through a 2 mm thick milk glass diffuser was used to produce the adapting field display. When viewed by the subject, a spatially uniform red (approximately 650 nm) circular field 2.5 deg in diameter was seen. During adaptation, the luminance of the field was increased to just under threshold levels.

### Table I: Patients’ Clinical Data

<table>
<thead>
<tr>
<th>Age</th>
<th>Sex</th>
<th>Kurtzke Scale</th>
<th>Visual Indications</th>
<th>Years Since Onset</th>
<th>Snellen Acuities</th>
<th>Acuity R</th>
</tr>
</thead>
<tbody>
<tr>
<td>23</td>
<td>f</td>
<td>2</td>
<td>right optic neuritis and pale discs bilaterally</td>
<td>2</td>
<td>20/30</td>
<td>20/20</td>
</tr>
<tr>
<td>21</td>
<td>f</td>
<td>2</td>
<td>left optic neuritis</td>
<td>1</td>
<td>20/15</td>
<td>20/20</td>
</tr>
<tr>
<td>47</td>
<td>f</td>
<td>3</td>
<td>pale discs bilaterally</td>
<td>11</td>
<td>20/30</td>
<td>20/30</td>
</tr>
<tr>
<td>40</td>
<td>f</td>
<td>1</td>
<td>left disc pallor</td>
<td>2</td>
<td>20/20</td>
<td>20/20</td>
</tr>
<tr>
<td>45</td>
<td>m</td>
<td>1</td>
<td>episodes of bilateral vision loss and bilateral pale discs</td>
<td>1</td>
<td>20/30</td>
<td>20/20</td>
</tr>
<tr>
<td>37</td>
<td>f</td>
<td>1</td>
<td>episodes of reduced vision left eye</td>
<td>3</td>
<td>20/30</td>
<td>20/30</td>
</tr>
<tr>
<td>32</td>
<td>f</td>
<td>2</td>
<td>episodes of bilateral vision loss</td>
<td>1</td>
<td>20/40</td>
<td>20/60</td>
</tr>
<tr>
<td>53</td>
<td>f</td>
<td>3</td>
<td>none</td>
<td>7</td>
<td>20/20</td>
<td>20/20</td>
</tr>
<tr>
<td>29</td>
<td>f</td>
<td>2</td>
<td>none</td>
<td>5</td>
<td>20/20</td>
<td>20/30</td>
</tr>
</tbody>
</table>
The adapting field was sinusoidally temporally modulated at a frequency of 7 Hz with a luminance modulation depth of 92%. Modulation depth is defined as \( \frac{(L_{\text{max}} - L_{\text{min}})}{(L_{\text{max}} + L_{\text{min}})} \times 100\% \) where \( L_{\text{max}} \) is the maximal luminance of the display and \( L_{\text{min}} \) is the minimal luminance. Modulation depth was measured with a photomultiplier. The time average luminance of this display (in the presence or absence of modulation) was 14 cd/m² as measured with an SEI photometer.

A single red LED, positioned at a right angle to the mirror box, was used to generate a 0.5 deg circular test stimulus. With the aid of a small beam splitter, the test spot appeared to the subject to be centered on the adapting field. A diagram of the apparatus is shown in Figure 1. A small test spot was used so that flicker measurements were not contaminated by edge effects. Apertures limiting the size of the test and adapting stimuli were placed at an equal distance from the observer so that no shift in accommodation was required upon presentation of the test stimulus. The mean luminance of the test spot was 27 cd/m². For all modulations, the test spot was always visible against the adapting field. The subject’s task in the experiment was to adjust the modulation depth in the flicker of the test spot until flicker was just barely detectable and then signalled the experimenter. (Note that unlike the measurement of critical flicker fusion, the frequency of flicker in the test spot was never varied.) Subjects experienced no difficulty in determining thresholds in the interval provided. At the beginning of each session, subjects were given verbal instructions regarding the task and two practice trials were conducted.

**General Procedure**

In each experiment, four trials were conducted for each eye of each subject. The eye used was alternated after every second trial. A single experimental trial consisted of a 4.5 min period of viewing the adapting field. This period was interrupted every 15 seconds with a 5 sec long presentation of the test stimulus. Whenever the test stimulus was present, the adapting field remained unmodulated to minimize edge contrast. At the onset of a test interval, the modulation depth in the test stimulus was set to a subthreshold value. The subject then adjusted the depth of modulation in the test spot until flicker was just barely detectable and then signalled the experimenter. (Note that unlike the measurement of critical flicker fusion, the frequency of flicker in the test spot was never varied.) Subjects experienced no difficulty in determining thresholds in the interval provided. At the beginning of each session, subjects were given verbal instructions regarding the task and two practice trials were conducted.

**Experiment I: Flicker Perception In The Absence Of Flicker Adaptation**

The first experiment was conducted to assess flicker perception abnormalities during prolonged viewing of an unmodulated field. Wright et al.\(^{11} \) have reported that some MS patients have higher unadapted flicker modulation depth thresholds than healthy controls. Therefore, before the effects of flicker adaptation could be assessed, deficits in unadapted flicker thresholds were determined. Using the procedure described above, flicker thresholds were assessed every 15 seconds while subjects viewed an unmodulated, spatially uniform adapting field. Six patients participated in the experiments as did five of the healthy control subjects. Of the 12 patient eyes in this subset, 3 had no sign of symptom of visual involvement.

![Figure 1 — A schematic of the apparatus. See text for details. AF, adapting field; T, test stimulus; LED, light emitting diode; D, diffuser; MB, mirror box; BS, beam splitter.](https://www.cambridge.org/core/core/1017/50317167100027761)

![Figure 2 — Minimum modulation depth needed to just detect flicker in a small circular test stimulus as a function of time viewing an unmodulated adapting field for MS and control groups. Open symbols represent group means for the MS patients and closed symbols represent group means for the control subjects. Vertical line represent ± 1 s.e. of the mean.](https://www.cambridge.org/core/core/1017/50317167100027761)
Results

The group mean modulation depth thresholds obtained during Experiment I are shown in Figure 2. Modulation depth thresholds for the healthy control group did not vary significantly as a function of time of testing and averaged 7.2%. Since the visual system typically performs a logarithmic transformation of sensory stimuli, statistical analyses for this and Experiment II were performed on log flicker thresholds. An analysis of variance on the mean log thresholds obtained from each eye showed that MS patients had slightly but significantly higher modulations on the mean log thresholds obtained from each eye showed testing [F (17,323) = 1.73, p < 0.05]. MS thresholds became significantly different from control thresholds after 75 sees of viewing the unmodulated adapting field. The average threshold of the first four values was used as the base-line flicker threshold against which adapted and decay thresholds were compared. Threshold elevations due to adaptation are expressed as a difference in the log baseline threshold and the log adapted threshold. (Threshold elevation = log adapted threshold - log baseline threshold).

EXPERIMENT II: ADAPTATION TO FLICKER

Since the previous experiment indicated that MS patients’ initial modulation depth threshold may be higher than those of controls, a within-subject design was chosen for flicker adaptation measurement. A within-subject design in which subjects’ responses in a pre-adaptation condition were compared to their own responses in a post adaptation condition was also used to eliminate effects due to differences in pupil size (the level of ambient illumination was constant for both conditions) that may have existed between controls and MS patients. MS patients 1 to 9 participated in the experiment as did all the control subjects. The same basic procedure as above was used. As before, the first four threshold estimates were determined after viewing an unmodulated flickering adapting field. The next eight thresholds (i.e., adapted thresholds) were obtained after viewing a flickering adapting field. The last six thresholds (i.e. decay thresholds) were measured after viewing an unmodulated adapting field. The mean of the first four values was used as the baseline flicker threshold against which adapted and decay thresholds were compared. Threshold elevations due to adaptation are expressed as a difference in the log baseline threshold and the log adapted threshold. (Threshold elevation = log adapted threshold - log baseline threshold).

Results

Figure 3 illustrates the group mean changes in log flicker thresholds during and after adaptation to highly modulated flicker. An analysis of variance on the adapted threshold elevations showed a significant difference in the magnitude of the adaptation effects between the two subject groups [F (1,245) = 29.8, p < 0.001]. Both groups showed a significant increase in
threshold elevation with prolonged viewing of the adapting field \[F(7,245) = 10.1, p < 0.001\]. The interaction effect of group and viewing time was not significant indicating that the rate of development of adaptation effects did not differ after the initial 15 sec exposure to the adapting stimulus. However, the data clearly show that during the first 15 sec of exposure, the build-up of adaptation effects was dramatically faster in MS patients than in controls.

One notable characteristic of the data was that the standard deviations of individual threshold estimates measured in MS patients were significantly \((p < 0.01)\) greater than those for control subjects. A similar result was reported by Patterson, Foster and Heron\(^\text{23}\) who found that contrast detection threshold variability was abnormally high in MS patients relative to healthy controls during exposure to backgrounds of high luminance levels. This variability may be related to the “fading” of stimuli during adaptation intervals reported by some patients. In all cases patients reported that fading disappeared with the onset of the test stimulus and that it did not interfere with threshold adjustments. Fading was typically reported after at least 2 minutes into a testing trial.

The most consistent and dramatic effects of adaptation in the MS patients were seen after only 15 seconds of adaptation. At this early stage of adaptation, fading was never reported by patients and adaptation effects are relatively uncontaminated by nonspecific light adaptation shown in the previous experiment to occur in some MS patients after longer periods of viewing. The average threshold elevation found in the control group at this interval was 0.066 log units (s.d. = 0.036). Using s.d. in the calculation of confidence intervals, it was found that 5 of the 7 unaffected eyes in the MS group showed an average threshold elevation greater than the upper 99% confidence limit of the control group mean. All 9 MS patients could be distinguished from normal on this basis. Figure 4 shows the distribution of threshold elevations obtained from both groups.

All of the control eyes appeared normal.

The average threshold elevation measured in the control group during the entire adaptation period was 0.122 log units (s.d. = 0.035). Using s.d. in the calculation of confidence intervals, it was found that 3 of the 7 unaffected eyes in the MS group showed an average threshold elevation greater than the 99% confidence limits of the control group mean. Six of the 11 affected eyes in the MS group revealed an average threshold elevation greater than the upper 99% confidence limit of the control group mean. All 9 MS patients could be distinguished from normal on this basis. Figure 4 shows the distribution of threshold elevations obtained from both groups.

Prior to flicker or non-specific light adaptation, only 22 percent of the MS patients’ eyes exhibited abnormal flicker sensitivity. In all cases, signs and symptoms of visual involvement were already present in these eyes. An initial control experiment in which subjects viewed an unmodulated adapting field, revealed that some patients and no controls experience a small loss in flicker sensitivity after viewing an unmodulated field for 2 to 3 minutes. Enoch et al.\(^\text{25}\) observed that when patients viewed an intense, uniformly luminous field for a similar period of time, large losses in grating acuity could be temporarily produced. Both these data and those from the present study suggest that for MS patients, non-specific adaptation to light may cause reductions in sensitivity for a wide variety of visual functions. However, in the present study this effect was found for only 39% of the eyes tested and revealed a deficit in only 1 of the 7 eyes previously found to be uninvolved.

An important clinical contribution of a psychophysical analysis of visual function in MS patients could be to supplement information obtained from fundoscopic examination, visual acuity tests and patient reports of blurring and reduced vision episodes. Procedures which can identify clinically “normal” eyes as abnormal have potential use in the diagnostic process. Although tests of flicker sensitivity without adaptation or with nonspecific adaptation seem unable to detect subtle visual deficits, the present study shows that specific, i.e., flicker, adaptation procedures can produce temporary symptoms in otherwise symptomatic eyes, thus revealing subclinical visual involvement of the disease. After only 15 seconds of adaptation, 15 of the 18 eyes studied (83%) demonstrated an abnormal sensitivity to flicker. Particularly noteworthy is that this procedure was able to identify 5 of the 7 unaffected eyes as abnormal. Contrast sensitivity (CS) measurements were available for the 5 eyes so identified and, in all cases, CS was found to be normal. Double flash data was available for 3 of the 5 eyes and, again, in all cases was found to be normal. Of the twelve eyes in the study for which double flash thresholds were available, 6 of the 6 eyes found to be normal with this test were observed to be abnormal with the flicker adaptation test. While the present pro-
procedure may have significant diagnostic value, additional research is needed to determine how specific the observed effects are for MS as opposed to other neurological and ophthalmological disorders.

The results of the present experiment suggest that partial demyelination of visual pathway neurons may exist in MS patients without signs or symptoms of visual involvement. The prolonged stimulation provided during visual adaptation procedures may produce sufficient exacerbation of neural transmission abnormalities to render reductions in sensory sensitivity. Unlike other methods used clinically to produce temporary exacerbation of symptoms in MS for the purpose of diagnosis (e.g., hot bath or exercise), visual adaptation is local to the visual system, painless, and recovery is complete within a few minutes.

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REFERENCES