Visual Defects Associated with Vigabatrin: A Study of Epileptic Argentine Patients

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ABSTRACT: Objective: The aim of the present study was to assess visual alterations in a population of Argentine patients treated with the antiepileptic drug vigabatrin. Methods: Twenty patients receiving vigabatrin and 15 patients receiving carbamazepine were examined with automated perimetry using a Humphrey 120-point full screening strategy. In addition, scotopic flash electrotinograms were performed. Results: Of 20 patients treated with vigabatrin, two were unable to cooperate with testing. Of the remaining 18 patients, all but two showed at least one non-detected point inside the central 40° of the visual field of each eye. Of the 15 carbamazepine-treated patients, three were unable to perform the study. None of the remaining 12 patients showed visual field defects. Both a- and b-wave amplitudes of the scotopic electrotinogram were significantly reduced in 12 patients receiving vigabatrin. Conclusions: Visual field defects among patients on vigabatrin therapy may occur with a higher frequency than previously recognized. The Humphrey 120-points full field screening test and electrotinography are useful tools to assess the visual dysfunction associated with vigabatrin.

RÉSUMÉ: Déficits visuels associés au vigabatrin: étude de patients argentins. Objectif: Le but de cette étude était d’évaluer les changements visuels chez un groupe de patients argentins traités par le vigabatrin, un médicament anti-épileptique. Méthodes: Vingt patients recevant du vigabatrin et 15 patients recevant de la carbamazépine ont subi une périmétrie de dépistage plein écran de 120 points au moyen de l’appareil automatisé Humphrey. De plus, des électroténotogrammes au flash scotopique ont été effectués. Résultats: L’étude n’a pu être réalisée chez deux des 20 patients traités par le vigabatrin à cause de leur manque de coopération. Chez les 18 autres patients, tous sauf deux avaient au moins un point non détecté au champ visuel central de 40° de chaque œil. Trois des 15 patients recevant de la carbamazépine n’ont pu compléter l’étude. Aucun des 12 autres patients n’avait de déficit au niveau des champs visuels. L’amplitude des ondes a et b de l’électroténotogramme scotopique était significativement diminuée chez 12 patients recevant le vigabatrin. Conclusions: Des anomalies du champ visuel sont peut-être plus fréquentes qu’on ne l’estimait chez les patients recevant du vigabatrin. La périmétrie automatisée Humphrey plein écran de 120 points et l’électroténotigraphie sont des méthodes utiles pour évaluer la dysfonction visuelle associée au vigabatrin.


Vigabatrin is a highly effective antiepileptic drug used for the treatment of infantile spasm and partial seizures, particularly those unresponsive to classic antiepileptic drugs. Its efficacy is related to the ability to interfere irreversibly with GABA (γ-aminobutyric acid) transaminase activity, the enzyme involved in the catabolism of GABA, thereby increasing the levels of this neurotransmitter in the central nervous system.

Vigabatrin was thought to be a relatively free of side-effects until 1997, when Eke et al.⁴ described constriction of the visual fields of three patients taking vigabatrin. The loss of field persisted after the drug was stopped. Since then, persistent visual field constriction following vigabatrin treatment has been confirmed in several reports⁵ but the prevalence of visual dysfunction is still controversial. We examined visual functions in a population of Argentine epileptic patients receiving this drug. A control group of 15 epileptic patients taking carbamazepine was also evaluated.
Patients diagnosed with partial epilepsy were recruited from the Epilepsy Center, “Ramos Mejía” Hospital, Buenos Aires, Argentina. Twenty patients treated with vigabatrin and 15 patients treated with carbamazepine (who had never taken vigabatrin) consented to the study. This research adhered to the tenets of the Declaration of Helsinki and was approved by the Institutional Review Board/Ethics Committee.

Ophthalmologic Examination

A full ophthalmologic evaluation was performed to exclude other visual disorders. We recorded visual symptoms, best-corrected visual acuities, colour vision (Ishihara 24-plate test), intraocular pressures, ocular alignment and findings on biomicroscopy and direct and indirect ophthalmoscopy.

Visual Fields

Perimetry was performed with a Humphrey field analyzer (Humphrey Systems INC, San Leandro CA, U.S.A.), using the 120-Points Full Field Screening Test, which tested a field-radius of 60 degrees. The visual field examinations were performed with normal-sized pupils, i.e. without pharmacologically induced mydriasis or miosis.

The studies were performed according to the manufacturer’s recommendations, using a white, size III (4 mm²) stimulus against a background illumination of 31.5 asb. Other test parameters were at their default values. Patient data, including age, corrective lens used (based on distance prescription with age-appropriate convex spherical add), pupil diameter, and visual acuity, were recorded. Fixation was monitored using a magnified infrared camera image every one to two minutes. Virtual cursors were manually adjusted before each test to set a fixation “window” which, if transgressed by the pupil margins, would stop the test until fixation was restored. The test time for the procedure was approximately seven minutes per eye, but rest breaks were allowed when requested. The same instructions were given to each participant.

Humphrey perimetry was considered unreliable if the blind spot could not be plotted. In order to assess the reproducibility of the visual field results, almost 30% of the patients from both groups were asked to repeat the study at least 24 hours after the first examination. In these cases, no differences between the first and the second examination were found. Two of 20 patients treated with vigabatrin, and three of 15 treated with carbamazepine, were unable to cooperate with testing. All visual field results were assessed by one ophthalmologist (JB), without knowledge of the clinical details of each patient. Visual field was considered defective when at least one non-detected point inside 40º for both eyes was found.

Electrophysiology

Electroretinography (ERG) was performed with a flash stimulator; twenty minutes of dark adaptation occurred before the stimulus was presented in a scotopic ambient. The active electrode was a gold foil surface electrode placed on the skin over the infraorbital ridge near the lower eyelid. The reference electrode was placed 2 cm temporal to the lateral orbital rim and the ground electrode was placed slightly above the supraorbital rim. ERGs were recorded in response to 1 Hz flash stimulation (flash strength 3 cd/m²). ERGs were filtered at 1.5 to 800 Hz, and the signal was averaged. Each eye was tested separately while the second eye was occluded. The amplitudes and latencies of the a- and b- wave were measured and averaged for each patient.

RESULTS

Demographic and clinical data

The clinical details related to the 20 vigabatrin-treated and 15 carbamazepine-treated patients are presented in Table 1. The age range was 19 to 49 years old for the vigabatrin group and 15 to 52 years old for the carbamazepine group. The vigabatrin patients were taking the drug on a daily basis and the range of the duration of treatment was one to eight years. No patients had noted vision problems. All patients had corrected visual acuities of 20/20 in both eyes except patient 20 who had monocular congenital glaucoma. This eye was excluded from the study. No other abnormalities of visual acuity or colour vision were detected in vigabatrin-treated or carbamazepine-treated patients. Fundus inspection revealed no pathology.

The visual fields shown in Figure 1 exemplify the normal results typically observed in the carbamazepine group (patient 23) and the patterns of field loss seen in the vigabatrin-treated patients: nasal constriction (patient 9), more extensive nasal constriction with mild temporal constriction (patient 18), and marked generalized constriction with patchy central involvement (patient 10).

Table 2 summarizes the Humphrey perimetry results of both eyes of the patients receiving vigabatrin. Patients 8 and 12 were unable to cooperate with visual field assessment, leaving 18 assessable patients. All but two patients (patient 1 and 17) failed to detect at least one point inside the 40º radius with each eye. None of the 12 assessable carbamazepine patients showed this number of non-detected points within the central 40º. Of the affected vigabatrin patients, 45% had marked field constrictions with central involvement and the rest had more mild constriction. The nasal field was involved in all affected eyes, and the temporal field was involved in 58% of the eyes. A right -left eye pairing (assessed as the total number of non-detected points for each eye) resulted in a highly significant correlation for all vigabatrin patients (p< 0.0001), indicating that the dysfunction was bilateral.

No correlation between the extent of the visual field defects and the duration, daily dosage, or cumulative dosage of vigabatrin was observed. The visual field loss of one patient who received vigabatrin for two years (patient 10; Figure 1) was similar to that of patient taking vigabatrin for six years (patient 11). The two patients who had normal visual fields had taken vigabatrin for three and five years. We also found no relationship between the extent of visual field defects and the type or duration of the epileptic syndrome.

The scotopic ERG of one patient receiving carbamazepine and three patients receiving vigabatrin are shown in Figure 2. Of the 20 patients treated with vigabatrin, patient 2 failed to attend, leaving 19 assessable patients. Figure 3 shows the means (± SD) of the scotopic a- and b- wave amplitudes of the 19 vigabatrin-treated and 15 carbamazepine-treated patients. The amplitudes were significantly reduced in patients receiving vigabatrin (for a-wave: control = 7.89 ± 2.71 µV, vigabatrin = 5.31 ± 2.22 µV,
p<0.01; for b-wave: control = 14.16 ± 4.76 μV, vigabatrin = 9.19 ± 3.74 μV, p< 0.01). A right-left eye pairing of scotopic a- and b-wave amplitudes resulted in a significant correlation for all vigabatrin patients (p<0.05 for a-wave, p<0.001 for b-wave).

The a- and b-wave amplitudes of 12 patients receiving vigabatrin fell below the lower limit of the 95% confidence interval of control patients. Of the two patients that had normal visual fields, only the a-wave amplitude from patient 17 was abnormal. The scotopic a-wave and b-wave latencies showed no significant differences for control and vigabatrin-treated patients.

**DISCUSSION**

In this study, 89% of the patients taking vigabatrin failed to detect at least one point in the central 40° of the visual fields of both eyes on the Humphrey 120-point full field screening test. No carbamazepine-treated patients had abnormal fields by this criterion. The vigabatrin-treated patients showed a similar number of non-detected points in both eyes. As in previous studies, the nasal field was predominantly affected and no patients complained of visual symptoms.

The prevalence of visual field constriction associated with vigabatrin has varied widely in previous reports, ranging from less than 0.1%, as initially reported by the manufacturer, to more than 50%. The results of the present study showed a higher prevalence than that in most previous studies. The higher prevalence may be related to our use of a full field screening technique rather than the more detailed, but slower and more spatially constricted technique of full threshold estimations. The screening technique is less quantitative than the threshold technique but may be more sensitive in detecting vigabatrin-
related field loss. The Humphrey screening technique assesses a radius of 60 degrees and monitors false-positive and false-negative responses and fixation behavior. The stimuli are above threshold and reliably detect major changes, whereas threshold automated perimetry may be less reliable for testing points outside a radius of 30 degrees. Although we tested a 60° radius, our criterion to define an abnormal field was based on points missed within a 40° radius. Thus, methodological differences may account for the higher prevalence of visual field constriction. This is supported by the observation that, if only the central 30 degrees is considered, the prevalence of field loss in our population is close to that of previous reports. Using similar methodology to ours, Midelfart et al. recently reported a similar frequency of visual field defects (86%) in a group of eighteen epileptic children treated with vigabatrin.

In this study, scotopic ERGs showed a significant reduction of both a- and b-wave amplitudes in vigabatrin-treated patients; over 60% had amplitudes that fell below the lower confidence limit of carbamazepine-treated patients. In previous reports, the prevalence of ERG dysfunction in vigabatrin-treated patients has varied from 0% to 100%. Most ERG abnormalities have been observed under photopic conditions, implicating cone dysfunction. Studies by Daneshvar et al. and Miller et al. have also shown reduction of b-wave amplitudes under scotopic conditions. In the present study, no correlation was observed between the scotopic ERG abnormalities and the visual field results. Miller et al. have observed a strong correlation between cone ERG amplitudes and field constriction as measured by kinetic perimetry. The results of this study confirm that scotopic ERG abnormalities are common in vigabatrin-treated patients and may be a useful indicator of retinal dysfunction, especially in patients for whom reliable perimetry cannot be performed.

The retinal locus and mechanism that account for the visual disturbance associated with vigabatrin therapy are not well understood. Dysfunction of both the cone and the rod systems...
has been postulated. Whether the dysfunction results from nonspecific toxicity or a chronic increase in retinal GABA levels remains an open question. GABA is a major inhibitory neurotransmitter of the vertebrate retina. In several species, the horizontal and amacrine cells are the principal GABAergic neurons in the retina. Perfusion with GABA agonists in the cat alters the responses of horizontal, amacrine, and ganglion cells. In albino rats (but not pigmented rats), vigabatrin has a dose-dependent effect on the retina, characterized by disruption of the outer nuclear layer, which contains the photoreceptor nuclei. GABA selectively affects components of the ERG. In rabbits, low concentrations of extracellular GABA enhance a- and b-wave amplitudes, whereas high concentrations reduce the b-wave amplitude and have no effect on the a-wave amplitude. In albino rats (but not pigmented rats), vigabatrin has a dose-dependent effect on the retina, characterized by disruption of the outer nuclear layer, which contains the photoreceptor nuclei. GABA selectively affects components of the ERG. In rabbits, low concentrations of extracellular GABA enhance a- and b-wave amplitudes, whereas high concentrations reduce the b-wave amplitude and have no effect on the a-wave amplitude. In rats, the inactivation of GABA transaminase is greater in the retina than in the brain; and the retina does not develop tolerance to vigabatrin, unlike the brain and the spinal cord. Reduction of b-wave amplitude under scotopic and photopic conditions could be the result of GABA-induced Müller cell dysfunction.

The results of this study provided further support for a casual relationship between vigabatrin treatment and a specific pattern of visual field loss and electroretinographic dysfunction. Our results suggest that patients receiving vigabatrin should have their visual fields monitored regularly and that the Humphrey 120-point screening technique and electroretinography are useful monitoring tools.

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