Gyrate atrophy of the choroid and retina: ERG of the neural retina and the pigment epithelium

Christina Raitta, Synnöve Carlson, Kaarina Vannas-Sulonen

Abstract

The function of the neural retina and the pigment epithelium in 10 patients with gyrate atrophy was examined by the conventional alternating current electroretinogram and by direct current electroretinography to study the c-wave. The a- and b-wave responses were subnormal in all patients and the ERG was undetectable in patients with an advanced stage of the disease. The c-wave was recordable by the DC ERG from three patients with a less advanced stage of the disease. In one patient the c-wave could be recorded only from the other eye but the a- and b-wave responses were recordable in both his eyes.

TABLE I Age and sex distribution, visual acuity, and stage of disease and cataract of the GA patients

Case	Age	Sex	GA stage	Vision		Cataract	
				OD	OS	OD	OS
1	7	F	I	0.6	0.7	—	—
2	12	F	I	0.8	0.7	Central opacity	—
3	14	F	I-II	0.7	0.4	Small opacity	—
4	16	M	I-II	0.5	0.2	—	—
5	21	M	I	0.8	0.6	—	—
6	36	M	III	0.2	0.2	Aphakia	—
7	37	M	IV	0.25	0.3	Aphakia	—
8	38	F	II-III	0.2	0.3	Aphakia	—
9	41	F	IV	0.2	0.2	Aphakia	—
10	46	M	II	0.6	0.4	Posterior opacity	—

Grading¹⁶: Stage I: Round, separate, or fused atrophic areas in the midperiphery of the fundus. Stage II: Fused atrophic areas approaching the posterior pole. Stage III: A large degenerative area round the disc. Stage IV: all the posterior pole except the macula are atrophic.

Gyrate atrophy of the choroid and retina (GA) is a rare form of hereditary retinal dystrophy accompanied by a defect in ornithine metabolism.¹ The clinical ophthalmological findings in GA include myopia, cataract, and chorioretinal degeneration. The metabolic defect presents as hyperornithinaemia, ornithuria, and deficiency of ornithine aminotransferase activity.² The human gene responsible for GA has been localised to chromosome 10.³

The full-field ERG has been shown to be reduced in amplitude at an early stage in GA patients,⁴ and no normal ERG recordings have been reported in them.⁵ Ophthalmoscopic and fluorescein angiographic findings of young patients suggest that the retinal pigment epithelium is affected early in the course of the disease.^{4,6} One histological study on animals indicates that the primary changes in GA are located in the pigment epithelium.⁷

We studied 10 GA patients with a full-field flash ERG to establish the degree and cell level of the disturbance in the function of the neural retina. The c-wave of the ERG, which originates in the pigment epithelium,^{9,10} was recorded by a direct current (DC) method.⁸ The purpose of this study is to evaluate the function of the pigment epithelium as revealed by the ERG c-wave and to correlate these findings with the a- and b-waves of the conventional alternating current (AC) ERG in GA patients.

Material and methods

Ten GA patients, five females and five males, aged 7-46 years (mean 27 years) were studied. The results were compared with the recordings of eight normal subjects (aged 20-46 years, mean 31 years). The GA diagnosis was based on ophthalmoscopic findings, hyperornithinaemia, ornithuria, and established deficiency of the ornithine aminotransferase enzyme.¹¹ The latest plasma ornithine level of the patients was at 205-

1300 $\mu\text{mol/l}$ (mean 733 $\mu\text{mol/l}$, normal <70 $\mu\text{mol/l}$). The visual functions of the patients are summarised in Table I.

The subjects were dark adapted for 30 minutes before the ERG recording. The pupils were dilated with 0.5% tropicamide (Oftan Tropicamide) which produced a dilatation of 7-8 mm. The other eye was covered. The recordings were performed with the subjects supine. Both eyes were examined in the same session, the right eye first. There was an additional 15 min dark adaptation between the recordings of the two eyes.

A photostimulator (Grass PS22) with a xenon flash lamp was used. The duration of the flash was 10 μs and the maximum intensity 3.38 $\mu\text{J}/\text{cm}^2$ at the eye level (measured with Model 350 Linear/Log Optometer with a photometric filter 21043, United Detector Technology). The stimulus light was attenuated with neutral density filters (Kodak). To obtain a near uniform illumination of the retina a full-field stimulator was used (LKC Systems Inc), which reduces the problems related to uncertain fixation.^{12,13}

The recordings were performed with a disposable corneal wick electrode described by Sieving *et al*¹⁴ and slightly modified by us. The recording method is described in detail elsewhere.⁸ As no lid speculum was used, the subject was able to blink. The electrode was placed on the surface of the topically anaesthetised eye (0.4% oxybuprocaine hydrochloride, Oftan Obucain), where it settled above the margin of the lower lid. The electrode is well tolerated, and occasional blinking keeps it moist.

Beckmann's Ag-AgCl disk electrodes were used as ground (on the forehead) and reference (on the ipsilateral temple) electrodes. The preparations for the recording were performed in a darkened room with the help of a dim red slit-lamp (luminance 2 cd/m^2). The corneal potentials were amplified (amplification 1000) with an

University of Helsinki,
Finland
Department of
Ophthalmology
C Raitta
K Vannas-Sulonen

Department of
Physiology
S Carlson

Correspondence to:
Christina Raitta, MD,
Department of
Ophthalmology, University of
Helsinki, Haartmaninkatu
4 C, SF-00290 Helsinki,
Finland.

Accepted for publication
18 January 1990

TABLE II The a- and b-wave implicit times and amplitudes of the GA patients and normal subjects

	Patients			Controls			Student's <i>t</i> test	
	<i>n</i>	Mean (ms)	SD	<i>n</i>	Mean (ms)	SD	<i>t</i>	<i>p</i>
IMPLICIT TIME								
<i>a-Wave</i>								
KW 47B+ND 0.6	11	29.3	4.1	11	27.7	0.72	1.27	—
KW 25	11	17.9	2.8	11	16.3	0.47	1.86	—
ND 1	11	19.9	4.4	11	17.0	0.44	2.18	<0.05
Bgr, KW 25	10	15.9	2.6	11	14.0	1.13	2.23	<0.05
Bgr, ND 1	10	15.9	2.0	11	14.6	0.98	1.90	—
<i>b-Wave</i>								
KW 47B+ND 2	10	85.5	9.9	11	81.9	4.1	1.10	—
KW 47B+ND 0.6	11	54.6	6.6	11	51.1	2.1	1.48	—
KW 25	11	51.6	3.9	11	52.7	1.5	0.87	—
ND 1	11	45.5	7.3	11	43.0	3.1	1.05	—
Bgr, KW 25	11	31.8	3.6	11	28.2	0.45	3.29	<0.01
Bgr, ND 1	11	29.7	2.9	11	27.3	0.72	2.64	<0.05
Flicker	11	32.8	1.6	11	29.9	1.27	4.71	<0.001
							Wilcoxon test	
	<i>n</i>	Med. (μV)	Range min/max	<i>n</i>	Med. (μV)	Range min/max	<i>u</i>	<i>p</i>
AMPLITUDE								
<i>a-Wave</i>								
KW 47B+ND 0.6	11	3.5	3.5/42	11	63	46/101	3.94	<0.001
KW 25	11	17.5	3.5/49	11	45.5	32/81	3.84	<0.001
ND 1	11	38.5	3.5/119	11	136	88/220	3.74	<0.001
Bgr, KW 25	10	7.9	3.5/13	11	17.5	14/32	3.83	<0.001
Bgr, ND 1	10	9.4	1.8/28	11	21	15/35	3.17	<0.001
<i>b-Wave</i>								
KW 47B+ND 2	10	40.2	21/74	11	157.0	95/220	3.83	<0.001
KW 47B+ND 0.6	11	49.0	28/154	11	213.0	147/329	3.87	<0.001
KW 25	11	45.5	18/175	11	248.0	147/413	3.84	<0.001
ND 1	11	38.5	18/108	11	157.0	84/325	3.81	<0.001
Bgr, KW 25	11	24.5	14/30	11	71.7	44/129	3.94	<0.001
Bgr, ND 1	11	32.5	6/49	11	96.2	51/140	3.94	<0.001
Flicker	11	24.5	7/61	11	59.5	25/110	2.99	<0.01

KW=Kodak Wratten. ND=neutral density. Bgr=background. n=Number of eyes.

AC/DC ERG amplifier (Biotronics, PO Box 29, SF-02421 Jorvas, Finland) and filtered (AC recording; low pass frequency 400 Hz, DC recording; low pass frequency 30 Hz). The recorded potentials were displayed on a digital oscilloscope (Gould 1425), which also provided the possibility for signal averaging. The implicit times (measured from the onset of the stimulus to the peak of the waves) and the amplitudes (measured from the baseline to the peak of the waves) were measured with a cursor. For the a-wave the earliest negative peak was measured. The waveforms were drawn from the oscilloscope with a pen recorder (Gould LS270) with a slow drawing speed (20 s/trace).

In the first recording session both eyes were studied for rod and cone mediated responses. Dark-adapted responses were elicited with the following test stimuli in the absence of a background illumination:

(1) Blue flash (KW47B, intensity $0.247 \times 10^{-1} \mu\text{J}/\text{cm}^2$) attenuated with neutral density filters to determine the b-wave threshold. (2) Blue flash (KW47B with 2.0 log units attenuation, intensity $0.195 \times 10^{-3} \mu\text{J}/\text{cm}^2$) to elicit a rod mediated response. (3) Blue flash (KW47B with 0.6 log unit attenuation, intensity $0.140 \times 10^{-1} \mu\text{J}/\text{cm}^2$) to elicit a rod mediated response. (4) Red flash (KW25, intensity $0.459 \mu\text{J}/\text{cm}^2$) to elicit a mixed cone and rod mediated response; the red stimulus and the previous blue stimulus (KW47B with 0.6 log unit attenuation) were matched subjectively to be equally strong stimuli for the rod system.¹⁵ (5) White flash with 1.0 log unit attenuation (intensity $0.394 \mu\text{J}/\text{cm}^2$) to elicit a mixed cone and rod mediated response.

With a white background illumination in the

globe (luminance 28 cd/m²) the following test stimuli were presented to elicit cone mediated responses:

(1) Blue flash (KW47B with 0.6 log unit attenuation). With this stimulus no response was recordable in normal subjects, so that the same stimulus in the dark-adapted conditions elicited the response mainly from the rod system. This control test was performed only in normal subjects. (2) Red flash (KW25). (3) White flash with 1.0 log unit attenuation. (4) White 30 Hz flicker (intensity of the white flash $3.38 \mu\text{J}/\text{cm}^2$). The c-wave was recorded in a separate session.

The patient was prepared for the recording as described above. A DC recording method was used which enabled the recording of the slow c-wave. A small red fixation light was lit inside the globe to help the patient maintain fixation for several seconds. The subjects were instructed to avoid movement and blinking during several seconds after the stimulus. The stimulus was a white flash with 1.0 log unit attenuation (intensity $0.394 \mu\text{J}/\text{cm}^2$). A prestimulus time of 0.96 second was provided for the recordings by the recording system, and the response to each stimulus was recorded for 10 seconds. No signal averaging was used to avoid contamination of the recordings by occasional traces containing eye blinks or eye movements. Several single responses were recorded. All recordings were performed in the morning.

Results

AC ERG was performed in both eyes of eight patients (cases 1–5, 7, 9, and 10). In two patients (cases 6 and 8) only one eye was studied because their other eye had recently been operated on for cataract. When a- and b-waves were undetectable, no c-wave recording was attempted (cases 6–9). Thus the DC ERG was performed on six patients (cases 1–5 and 10).

The AC ERG was isoelectric in both eyes of two patients (cases 7 and 9), in the recorded eye of cases 6 and 8, and in one eye of case 3. The b-wave threshold was raised in all patients with measurable responses.

The results of the a- and b-wave recordings of those GA patients whose ERG was not isoelectric are listed in Table II together with the results of the normal recordings. The implicit times of the a- and b-waves were slightly delayed in GA patients in comparison with normal subjects, but the difference was statistically significant only for the cone mediated responses (statistical analysis was by Student's *t* test for independent samples) (Table II). The implicit time of the a-wave was significantly delayed in GA patients in comparison with normal subjects also in one test for mixed rod and cone mediated response (white flash without background illumination). In all recordings of the GA patients the a- and b-wave amplitudes were reduced. The difference in results between the GA patients and normal subjects was also statistically significant (non-parametric Wilcoxon test for independent samples) (Table II). Figure 1 shows an example of the normal recording. The responses of three patients (cases 2, 4, and 8) are illustrated in Figure 2A, B, C.

Figure 1: Normal AC ERG recording. From top to bottom: b-wave threshold (blue stimulus, KW 47B, ND 3.3, intensity $0.118 \times 10^{-4} \mu\text{J}/\text{cm}^2$); rod mediated response (KW 47B, ND 2); rod mediated response (KW 47B, ND 0.6); mixed rod and cone mediated response (red stimulus, KW 25); mixed rod and cone mediated response (white stimulus, ND 1); no response (see text) (white background illumination, $28 \text{ cd}/\text{m}^2$, and blue stimulus, KW 47B, ND 0.6); cone mediated response (white background illumination and red stimulus, KW 25); cone mediated response (white background illumination and white stimulus, ND 1); cone mediated response (white background illumination, 30 Hz white flicker, flash with no attenuation). Dashed line indicates the occurrence of the stimulus (duration 10 μs). Each recording is the average of four single responses. B=blue, R=red, W=white, bgr=background illumination.

Figure 1

Figure 2: Flash ERG of three patients. A: A 16-year-old male (case 4), right eye. B: A 12-year-old girl (case 2), right eye. C: A 38-year-old woman (case 8). The stimuli from top to bottom: (1) blue (KW 47B, ND 2); (2) blue (KW 47B, ND 0.6); (3) red (KW 25); (4) white (ND 1); (5) white background illumination (28 cd/m^2), red flash (KW 25); (6) white background illumination, white flash (ND 1); (7) white background illumination, 30 Hz white flicker, flash with no attenuation. Each recording is the average of four single responses. B=blue, R=red, W=white, bgr=background illumination.

Figure 2A

Figure 2B

The c-wave was recorded in six patients (cases 1–5 and 10). In one patient (case 1) the DC recording failed. This 7-year-old girl could not avoid moving the eyes during the recording. In one patient (case 2) the c-wave was recordable in both eyes (Fig 3). She was the second youngest (12 years) of the patients in this study. The clinical ophthalmological findings of her eyes corresponded to stage I in the grading described by Takki¹⁶ (Fig 4). In the 16-year-old male (case 4) the c-wave was recordable from the right eye but not from the left (Fig 3). This finding was verified by repeating the recordings the following day. The clinical ophthalmological findings

Figure 2C

Figure 3: Examples of the c-wave recordings. Top: Normal c-wave recording. Second: Case 2. C-waves were recordable in both eyes. Third: Case 4. C-wave was recordable in the right eye (left) but not in the left eye (right). Bottom: Case 10. A small positive deflection at a latency characteristic of the c-wave in the recordings from both eyes. The stimulus, a white flash with 1.0 log unit attenuation (duration 10 μ s), was presented just before the occurrence of the a- and b-waves with a latency of 0.96 s from the beginning of the recording. Two to four responses are superimposed.

TABLE III Implicit time and amplitude of the c-wave in GA patients and in normal persons

		Implicit time (s)		Amplitude (μ V)	
Case 2	OD	0.79		127	
	OS	0.85		124	
Case 4	OD	0.79		98	
	OS	—		—	
Case 10	OD	0.76		10	
	OS	0.80		13	
Normal	n=9	0.90	0.08	147	80
		(Mean)	(SD)	(Mean)	(SD)

of his eyes corresponded to stage I–II (Fig 5). A small positive deflection with an implicit time characteristic of the c-wave, was recorded from case 10 (stage II). In the other two patients (cases 3 and 5) no c-wave was recordable from either eye. The results of the c-wave recordings of the GA patients are listed in Table III together with the normal values.

Figure 4: Composite photograph of the right eye at stage 1 of a patient aged 12 years (case 2). Mainly separate peripheral chorioretinal atrophic patches. Normal optic disc, macula, and retinal vessels.

Discussion

The first detailed ophthalmological description of the characteristics of GA appeared in 1974.¹⁶ Since then several reports on the electrophysiological findings in GA have recorded that the ERG is severely affected and may become undetectable at an early stage of the disease.^{4 6 17–20} Typical for the disease is that its progress is variable and does not correlate with the age of the patients.^{11 21}

In the present study the ERG was undetectable in patients whose clinical ophthalmological findings corresponded to GA stages II–IV (cases 3 and 6–9). The stage of the disease was less advanced in those patients who still had measurable responses in their ERG (cases 1–5 and 10). Both rod and cone mediated a- and b-wave amplitudes were decreased in the ERG of these patients, indicating that the rod and cone systems are affected jointly in GA. Of the two young patients described by Berson *et al*¹⁸ one had a markedly depressed ERG with rod responses more reduced than cone responses. Kaiser-Kupfer *et al*⁴ on the other hand did not report a difference in the degree of impairment of the two systems.

The c-wave was recordable in both eyes of the 12-year-old patient (case 2, stage I) of this study. In her AC ERG there was already a marked reduction in both cone and rod mediated responses. Of particular interest is case 4 (the 16-year-old male), in whose right eye the c-wave was still recordable but not in the left eye. The rod and cone mediated responses, however, were recordable in both eyes. In the DC ERG of case 10 (a 46-year-old male) a small positive deflection occurred in both eyes at a latency characteristic of the c-wave. The clinical picture of case 10 (stage II) indicates that his disease has progressed slowly and is an example of the heterogeneity of GA. The DC ERG findings of this study indicate that the c-wave can be recorded only at an early stage of the disease.

Since Noell's observation that the c-wave disappears after the pigment epithelium is selectively destroyed⁹ several studies have supported the idea that the c-wave is in large part elicited by the pigment epithelium.^{10 22} When retina is illuminated, there is a decrease in the extracellular levels of K⁺ around the photoreceptors which gives rise to a hyperpolarisation of the apical membrane and to a less extent of the basal membrane of the pigment epithelium.²³

The retinal pigment epithelium maintains the photoreceptor cell homeostasis. The light sensitive disc membranes in the outer segments of the photoreceptor cells are renewed normally throughout life. This process involves multiple steps,²⁴ and a defect in one step may disturb the renewal mechanism and lead to cell abnormalities or cell death. The disappearance of the c-wave at an early stage of the disease is an indication of the disturbed function of the pigment epithelium.

To our knowledge this is the first time that any electrophysiological evidence of the function of the pigment epithelium in GA has been presented. The c-wave was detectable only in the early stages of the disease. In all patients with recordable c-waves the photoreceptor responses were subnormal. With clinical ERG it is not

Figure 5: Composite photograph of the left eye at stages 1-2 of a patient aged 16 years (case 4). Sharply demarcated confluent atrophic patches of garland shaped configuration. Normal optic disc and retinal vessels. Small pigmentary defect parafoveally in the left eye.

possible to determine whether the pigment epithelium is affected earlier in GA than the neural retina. However, in case 4 the absence of the c-wave in the other eye, in which a- and b-waves were still recordable, may be an indication that the functional disturbance originates in the pigment epithelium.

We thank Mrs Pirjo Cleve for technical assistance. This study was supported by a grant from the Academy of Finland and from The Eye Foundation.

- 1 Simell O, Takki K. Raised plasma-ornithine and gyrate atrophy of the choroid and retina. *Lancet* 1973; i: 1031-3.
- 2 Sengers RCA, Trijbels JMF, Brussaart JH, Deutman AF. Gyrate atrophy of the choroid and retina and ornithine-ketoacid aminotransferase deficiency. *Pediatr Res* 1976; 10: 894.

- 3 O'Donnell JJ, Vannas-Sulonen K, Shows T, Cox D. Ornithine aminotransferase maps to human chromosome 10 and mouse chromosome 7. *Cytogenet Cell Genet* 1985; 40: 716A.
- 4 Kaiser-Kupfer MI, Ludvig IH, de Monasterio FM, Valle D, Krieger I. Gyrate atrophy of the choroid and retina. Early findings. *Ophthalmology* 1985; 92: 395-401.
- 5 Kaiser-Kupfer MI, Valle DL. Clinical, biochemical, and therapeutic aspects of gyrate atrophy. *Prog Retinal Res* 1987; 6: 179-206.
- 6 Takki K, Simell O. Gyrate atrophy of the choroid and retina with hyperornithinemia (HOGA). *Birth Defects* 1976; 12: 373-84.
- 7 Kuwabara T, Ishikawa Y, Kaiser-Kupfer MI. Experimental model of gyrate atrophy in animals. *Ophthalmology* 1981; 88: 331-4.
- 8 Carlson S, Raitta C, Kommonen B. A d.c. electroretinography method for the recording of a-, b-, and c-waves. *J Neurosci Methods*. (In press).
- 9 Noell W. The origin of the electroretinogram. *Am J Ophthalmol* 1954; 38: 78-9.
- 10 Steinberg R, Schmidt R, Brown K. Intracellular responses to light from cat pigment epithelium: origin of the electroretinogram c-wave. *Nature* 1970; 227: 728-30.
- 11 Vannas-Sulonen K. Gyrate atrophy of the choroid and retina. MD thesis, University of Helsinki. ISBN 952-90013-4-7, Helsinki, 1988.
- 12 Ikeda H. Retinal mechanisms and the clinical electroretinogram. In: Haliday AM, Butler SR, Paul R, eds. *A textbook of clinical neurophysiology*. New York: Wiley, 1987: 569-94.
- 13 Niemeyer G. Information von Der Netzhaut durch Electroretinografie. *Graefes Arch Clin Exp Ophthalmol* 1979; 211: 129-37.
- 14 Sieving P, Fishman G, Maggiano J. Corneal wick electrode for recording bright flash electroretinograms and early receptor potentials. *Arch Ophthalmol* 1978; 96: 899-900.
- 15 Gouras P. Electroretinography: some basic principles. *Invest Ophthalmol Vis Sci* 1970; 9: 557-69.
- 16 Takki K. Gyrate atrophy of the choroid and retina associated with hyperornithinaemia. *Br J Ophthalmol* 1974; 58: 3-23.
- 17 Berson EL, Schmidt SY, Rabin AR. Plasma amino-acids in hereditary retinal disease; ornithine lysine and taurine. *Br J Ophthalmol* 1976; 60: 142-7.
- 18 Berson EL, Schmidt SY, Shih VE. Ocular and biochemical abnormalities in gyrate atrophy of the choroid and retina. *Ophthalmology* 1978; 85: 1018-27.
- 19 Rinaldi E, Stoppoloni GP, Savastano S, Russo S, Corticelli L. Gyrate atrophy of choroid associated with hyperornithinemia: report of the first case in Italy. *J Pediatr Ophthalmol Strabismus* 1979; 16: 133-5.
- 20 Hayasaka S, Saito T, Nakajima H, et al. Gyrate atrophy with hyperornithinemia: different types of responsiveness to vitamin B6. *Br J Ophthalmol* 1981; 65: 478-83.
- 21 Kaiser-Kupfer MI, Kuwabara T, Askanas V, et al. Systemic manifestations of gyrate atrophy of the choroid and retina. *Ophthalmology* 1981; 88: 302-6.
- 22 Brown K, Wiesel T. Localization of origins of electroretinogram components by intraretinal recording in the intact cat eye. *J Physiol (Lond)* 1961; 158: 257-80.
- 23 Oakley B, Green D. Correlation of light induced changes in retinal extracellular potassium concentration with the c-wave of the electroretinogram. *J Neurophysiol* 1976; 39: 1117-33.
- 24 Boulton M, Marshall J. A mechanistic approach to the inherited retinal dystrophies and the role of tissue culture as an investigative probe. In: Sheffield JB, Hilfer SR, eds. *Cell and developmental biology of the eye. Heredity and visual development*. New York: Springer, 1985: 115-70.