Electrophysiological evaluation of visual loss in Müller cell sheen dystrophy

Ulrich Kellner, Hannelore Kraus, Heinrich Heimann, Horst Helbig, Norbert Bornfeld, Michael H Foerster

Abstract

Aims—To describe the clinical picture and electrophysiological findings in Müller cell sheen dystrophy, a recently reported retinal dystrophy.

Method—A basic ophthalmological evaluation as well as recording of standard electro-oculography and electoretinography were performed in one patient at the onset of visual loss and after 1 year of follow up.

Results—A 61 year old woman presented with visual loss in the right eye. Multiple folds at the level of the internal limiting membrane were seen at the posterior pole in both eyes. Macular oedema was present in the right eye. The visual acuity of the right eye was 6/30 and of the left 6/9. A paracentral scotoma was found in the right eye. Electro-oculographic examination of both eyes gave normal results. Electoretinography (ERG) revealed reduced b-wave and flicker amplitudes in the right eye; these potentials were normal for the left eye. The ON response in the right eye was reduced and delayed; it was normal in the left eye. A further loss of visual function was noted 1 year later in the right eye, but the ophthalmoscopic findings were unchanged. The ERG of the right eye had a negative waveform when dark adapted. Light adapted responses showed an unusual delayed b-wave, broad and delayed ON and OFF responses and a missing flicker response, suggesting a Müller cell dysfunction. Light adapted responses were slightly reduced in the left eye.

Conclusions—Electrophysiological data indicate Müller cell dysfunction as a background of functional loss in Müller cell sheen dystrophy. This is in agreement with previously reported histological findings in this disorder.

Patient and methods

The female patient was first examined at 61 years of age. She had noted acute loss of visual acuity in the right eye 4 weeks previously. The patient had had normal vision before this; further, her general medical history was unremarkable. A basic ophthalmological evaluation, as well as recording of standard EOG and ERG, was performed. The family history could only be partially obtained. The patient’s father had no vision problems up to his death at 80 years of age. The patient did not know her mother and she had no brothers or sisters. Her daughter had no visual problems up to the age of 38 years. Our patient had lost contact with her daughter for several years and therefore the daughter was not available for examination.

EOG and ERG were performed as described in detail previously. The recording techniques were in accordance with the standard...
ards for clinical EOG and ERG. Briefly, EOG was performed with maximal dilated pupils (2.5% phenylephrine and 0.5% tropicamidine) using a method described by Behrens et al. The response was described by the ratio between the maximum amplitude and the amplitude before the luminance was increased (light peak versus baseline). Normal ranges for baseline and light peak were defined by calculation of the median values and the 95% confidence intervals in one eye of 40 probands.

For ERG recording a Nicolet Spirit (Nicolet, Madison, WI, USA) in combination with a Nicolet Ganzfeld was used. Recordings were performed with a Jet contact lens electrode and with maximal dilated pupils. Averaging was not performed. ON and OFF responses were recorded with a LED stimulator (light emitting diode, Roland Consult, Brandenburg, Germany) using either red or green LEDs (3 cd.s/m²) and flashes of long duration (200 ms) presented on background illumination. Recordings were done after 10 minutes of light adaptation (10 cd/m²): 128 responses were averaged. The normal ranges for ERG and ON and OFF responses were defined by calculation of the median values and the 95% confidence intervals in one eye of 23 age matched probands.

All examinations were performed in agreement with the Declaration of Helsinki after informed consent was obtained from the patient.

Figure 1 (A) The posterior pole of the right eye shows multiple folds and an oedema in the lower part of the macula. (B) The posterior pole of the left eye shows similar folding, but no oedema. (C) The upper mid periphery of the right eye shows smaller folds in the mid periphery and no folds towards the periphery. (D) The upper nasal quadrant of the right eye shows folds nasally to the optic disc, and no folds towards the periphery. Comparison with (C) shows the variation in visibility of the folds depending on the angle of light as seen in the retina above the optic disc.

Figure 2 (A) Fluorescein angiography on the right eye reveals normal filling of retinal vessels. In the late venous phase dye leakage begins in a spot near the fovea. (B) In the late phase, dye leakage has stained the lower part of the macula.
Results

CLINICAL EXAMINATION

At the initial examination (May 1996), the patient complained of hazy vision in the right eye. She had no complaints in the left eye. Visual acuity was right eye 6/30 (+5.0 −0.75 axis 52) and left eye 6/9 (+5.75 −0.75 axis 126). Ocular pressure was within the normal range. The anterior segments were unremarkable. Multiple folds of the internal limiting membrane were seen at the posterior pole extending into the mid periphery in both eyes (Fig 1). The outer periphery appeared normal. The degree of folding was similar in both eyes. An additional macular oedema was present in the right eye and was confirmed on fluorescein angiography (Fig 2). Vascular leakage was not present outside of the macula.

Goldmann visual fields showed normal outer isoptres. A small paracentral scotoma was present in the right eye. Colour vision testing with the desaturated Panel D15 test revealed mild (left eye) to severe (right eye) errors without a typical axis of confusion.

At the follow up examination (April 1997), the patient complained of further visual loss in the right eye. Visual acuity in the right eye was reduced to 6/120. It was unchanged in the left eye. The ophthalmoscopic findings with respect to the location and severity of the folds were unchanged in both eyes. A macular oedema was still present in the right eye. The patient did not agree to repeat fluorescein angiography. Goldmann visual fields showed normal outer isoptres. A small paracentral scotoma was present in the right eye. Colour vision testing with the desaturated Panel D15 test revealed mild (left eye) to severe (right eye) errors without a typical axis of confusion.

ELECTROPHYSIOLOGICAL EXAMINATION

An EOG was recorded once at the time of the first examination. Baseline values and light peak were normal in both eyes. Visual fields remained normal in the left eye.

Table 1  ERG amplitudes (µV) (reduced amplitudes are underlined)

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Dark adapted:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rod:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>b</td>
<td>445</td>
<td>327</td>
<td>122</td>
<td>0</td>
<td>387</td>
<td>442</td>
</tr>
<tr>
<td>Rod-cone:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>a</td>
<td>464</td>
<td>376</td>
<td>504</td>
<td>415</td>
<td>505</td>
<td>494</td>
</tr>
<tr>
<td>b</td>
<td>742</td>
<td>600</td>
<td>453</td>
<td>331</td>
<td>744</td>
<td>744</td>
</tr>
<tr>
<td>Light adapted:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Single cone:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>a</td>
<td>103</td>
<td>72</td>
<td>85</td>
<td>100</td>
<td>93</td>
<td>105</td>
</tr>
<tr>
<td>b</td>
<td>246</td>
<td>200</td>
<td>149</td>
<td>127</td>
<td>271</td>
<td>189</td>
</tr>
<tr>
<td>Cone flicker:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>a</td>
<td>241</td>
<td>215</td>
<td>99</td>
<td>0</td>
<td>265</td>
<td>200</td>
</tr>
<tr>
<td>b</td>
<td>744</td>
<td></td>
<td></td>
<td></td>
<td>744</td>
<td></td>
</tr>
<tr>
<td>ON response:</td>
<td>35</td>
<td>28</td>
<td>78</td>
<td>12</td>
<td>49</td>
<td>35</td>
</tr>
<tr>
<td>OFF response:</td>
<td>32</td>
<td>20</td>
<td>20</td>
<td>33</td>
<td>42</td>
<td>38</td>
</tr>
</tbody>
</table>

Table 2  ERG implicit times (ms) (delayed implicit times are underlined)

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Single cone</td>
<td>32.0</td>
<td>34.8</td>
<td>34.0</td>
<td>nd</td>
<td>32.0</td>
<td>30.8</td>
</tr>
<tr>
<td>Cone flicker</td>
<td>29.3</td>
<td>31.5</td>
<td>32.0</td>
<td>nm</td>
<td>28.0</td>
<td>28.0</td>
</tr>
<tr>
<td>ON response</td>
<td>32.0</td>
<td>36.0</td>
<td>38.0</td>
<td>50.0</td>
<td>33.0</td>
<td>34.0</td>
</tr>
<tr>
<td>OFF response</td>
<td>22.0</td>
<td>24.0</td>
<td>26.0</td>
<td>32.0</td>
<td>23.0</td>
<td>24.0</td>
</tr>
</tbody>
</table>

nd=not determinable, because no distinct b-wave peak was present.
nm=not measurable owing to missing response.

ERGs were recorded at the first visit and at the follow up examination. The values for amplitudes and implicit times are given in Tables 1 and 2; the recordings are shown in Figures 3 and 4.

Right eye

At the first examination, recording at dark adaptation revealed normal a-wave amplitudes and reduced b-wave amplitudes (Fig 3). At maximum stimulus intensity, the b-wave amplitude was reduced to 61% of normal median. The ERG response had the configuration of a negative ERG (b-wave amplitude smaller than a-wave amplitude). At light adaptation, the a-wave amplitude, was in the lower normal...
Electrophysiological evaluation of visual loss in Müller cell sheen dystrophy

653

The functional disturbance in the left eye of our patient was mild despite the marked folds at the posterior pole. Indeed, she had not noted any vision problems in the left eye at all. Slowly progressive functional loss was, however, obvious in the light adapted ERG recordings after 1 year of follow up. Functional loss was, in contrast, more severe in the right eye. The most interesting findings were seen in the ERG; the negative waveform at dark adaptation, the delayed implicit times in light adapted responses, and the loss of the 30 Hz flicker response. Based on functional and ophthalmoscopic findings other retinal disorders associated with a negative waveform in the ERG (for example, congenital stationary night blindness, X linked retinoschisis, subtypes of retinitis pigmentosa or cone dystrophies, melanoma associated retinopathy, and severe retinal vascular disorders) could be excluded in our patient. Reduced ERG amplitudes and a negative waveform have been described in other patients with Müller cell sheen dystrophy.13

The normal a-wave amplitudes as well as the normal EOG in both eyes of our patient indicate normal function of the photoreceptor-pigment epithelium complex. Retinal bipolar cells and Müller cells are predominantly involved in the generation of the b-wave of the clinical ERG.14-20 The reduced b-wave and flicker amplitudes at both examinations in the right eye and at the last examination in the left eye suggest a functional defect either in bipolar cells or Müller cells. The ON response reflects activity of the depolarising bipolar cells, and the OFF response activity of the hyperpolarising bipolar cells.20 At the last follow up examination in the right eye of our patient, the ON and OFF responses were normal. Both responses showed a broader waveform with markedly delayed implicit times. This broader waveform of both responses could be explained by a delayed uptake of K⁺ ions in the Müller cells. A similar delay of implicit time of ON and OFF responses is in accordance with normal bipolar cell function in combination with a Müller cell defect. Therefore, the negative ERG at dark adaptation and the delayed implicit times at light adaptation in the right eye of our patient most likely indicate Müller cell dysfunction. The delayed implicit times of ON and OFF responses have two interesting consequences with regard to the waveform of the light adapted ERG. The b-wave of the single cone response is broader than usual and the return to baseline is delayed. The complete loss of the 30 Hz flicker response within 1 year can
be explained by the increased delay of both ON and OFF responses. The potentials of both dopolasing and hyperpolarizing bipolar cells and, to a lesser degree, the cone photoreceptor potentials have been recently demonstrated to constitute the 30 Hz flicker response. The delayed timing of both bipolar cell potentials apparently alters the summation for the flicker response, resulting in the elimination of a recordable flicker response.

The basement membranes of the Müller cells form the main part of the internal limiting membrane. Histologically, a diffuse thickening and an undulation of the internal limiting membrane with small inner retinal schisis cavities have been demonstrated in Müller cell sheen dystrophy. We can speculate that the primary defect in this disorder is within the Müller cells. During the early stage of the disease, Müller cells may produce a material which has yet to be characterised in nature and which accumulates in the internal limiting membrane. This material deposition induces thickening of the membrane and its subsequent folding. At this stage, the membrane properties of the Müller cells show normal function as far as they contribute to the generation of the b-wave of the ERG. This is demonstrated in the left eye of our patient at the first examination. At a later stage, the increased accumulation of material leads to disruption of the contact between the internal limiting membrane and the Müller cells and, consequently, inner retinal schisis cavities develop. The functional result includes the deterioration of the b-wave amplitude and the delayed implicit times, as seen in the right eye of our patient. The widespread increased retinal vascular permeability in the late stage of the disease, as observed in some patients, may be due to secondary damage to retinal capillaries, which are surrounded by a Müller cell basement membrane. It is reasonable to assume that the most severe folding occurs at the posterior pole since the internal limiting membrane is thicker at the posterior pole than in the periphery. The onset of clinical signs at 50 years of age or later may be accounted for by the normal thickening of the internal limiting membrane with age.

Treatment has not proved to be of benefit for patients with Müller cell sheen dystrophy. Vitrectomy performed in three eyes did not improve their visual function. We initiated a treatment with systemic acetazolamide and local flurbiprofen because this has been proved to be helpful in some cases of macular oedema from other causes. A positive effect of this treatment, however, could not be observed in our patient.

This study was partially supported by the Deutsche Forschungsgemeinschaft, Germany (U Kellner, grant DFG-Ke442/7-1).

18 Gurevich L, Slaughte MM. Comparison of the waveforms of the ON bipolar neuron and the b-wave of the electroretinogram. Vis Res 1993;33:2431-5.
19 Turn N, Slaughte MM. Correlation of dynamic responses in the ON bipolar neuron and the b-wave of the electroretinogram. Vis Res 1995;35:1359-64.