ELECTRO-OCULOGRAPHY AS A FUNCTIONAL TEST IN PATHOLOGICAL CONDITIONS OF THE FUNDUS
I. FIRST RESULTS*

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The modifications of the standing potential in pathological fundus conditions are practically unknown, although electro-oculography (EOG) is a technically simple and clinically very efficient method of obtaining an indirect estimate of its value. Although the EOG is now widely used in order to study ocular motility, it is generally not considered as a functional test: only Riggs (1954) has reported diminished deflections in a case of pigmentary degeneration, and we ourselves have studied the EOG in a few cases of congenital retinal abnormality, such as Oguchi's disease (François and Verriest, 1954) and total colour blindness (François, Verriest, and De Rouck, 1955b). The present paper deals with the technique, physiological values, and results obtained in a few cases of the more typical diseases of the choroid and the retina.

Technique

After a study of the regional differences of conduction on the face of one-eyed subjects (François and De Rouck, 1955), we adopted the following electrode scheme as giving the greatest amplitudes, the most constant regularity, and the most independent records for both eyes (Fig. 1):

1. On the temporal orbital margin, a little below the external canthus of the right eye;
2. On the base of the nose, near the internal canthus of the right eye;
3. On the base of the nose, near the internal canthus of the left eye;
4. On the temporal orbital margin, a little below the external canthus of the left eye;
5. A mass-electrode is placed on the forehead or on the occiput.

We use non-polarizable silver electrodes, attached to the skin with collodion. The skin is cleaned with ether, and contact is aided with commercial electrode jelly. An ink-electroencephalograph is used for the registration.

The subject performs, as rapidly as possible, two ocular movements in an interrupted series:

(i) from the extreme left to the extreme right (LR);  
(ii) from the extreme right to the extreme left (RL).

Fig. 1.—Scheme of arrangement of electrodes.

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The derivation 1/2 gives the registration of the standing potential of the right eye and 4/3 the same for the left eye (in cases of unilateral enucleation with or without implant or prosthesis, the corresponding derivation is nearly zero).

We have adopted for each eye (mean deflection) as the basis of estimating the value:

\[
M = \frac{LR + RL}{2}
\]

These measurements are effected in moderate brightness, without previous dark- or light-adaptation, to prevent a rapid modification of the values (François, Verriest, and De Rouck, 1955a).

**Physiological Values**

We recorded the EOG of both eyes in a series of 34 normal subjects. We also obtained 61 utilizable values of \( M \); these form a normal distribution with a good approximation (Fig. 2). The arithmetical mean \( m = 405 \, \mu V \) and the standard deviation \( \sigma = 60 \, \mu V \). Theoretically 95.4 per cent. of the normal values must be between \( m \pm 2\sigma \), or 285 and 525 \( \mu V \). In one subject we obtained very low values (\( M = 130 \, \mu V \) for the right eye and 215 \( \mu V \) for the left eye). According to Chauvenet’s criterion, we did not include this case in the statistics.

![Figure 2: Physiological values of M.](image)

We compared the values of the two eyes in 28 subjects and established that in normal conditions (Fig. 3, overleaf):

\[
\frac{M_{\text{max}} - M_{\text{min}}}{M_{\text{max}}} < 0.33
\]

**Pathological Eye-Conditions**

(1) **Microphthalmos.**—We examined two cases. In the first the condition was bilateral and the values of \( M \) very low (right eye 115 \( \mu V \); left eye 55 \( \mu V \).
In the second the condition was unilateral and complicated by a coloboma of the optic nerve and the choroid; on the affected side the EOG was markedly reduced (270 μV) in comparison with that of the sound eye (440 μV).

(2) Congenital Functional Abnormalities of the Retina.—We examined three cases of total colour blindness with amblyopia ("rod-monochromatism"). In one we obtained very high normal values (right eye 500 μV, left eye 450 μV); in the second lower, but normal, values (300 μV for both eyes); in the third we also obtained normal values, but these were not comparable, because we were using a different technique.

We also obtained normal values in a case of protanopia (right eye 425 μV; left eye 480 μV), in a case of pronounced protanomaly (400 μV, quotient of anomaly 0·16; marked depression of the spectral sensitivity in the red), and in cases of deuteranopia and deuteranomaly (quotient of anomaly 3·6).

A case of Oguchi’s disease showed a normal EOG (440 and 425 μV).

However, in all these congenital abnormalities, there are no histological features indicating that the functionally absent receptors are missing.

(3) Tapeto-Retinal Degenerations.—Eight cases of primary pigmentary degeneration were tested:

Four showed pathologically low values: 140 μV for both; Right 100 μV, left 170 μV; 200 μV for both; Right 250 μV, left 200 μV.

Three gave a subnormal value for one eye: Right 270 μV, left 340 μV; Right 245 μV, left 390 μV; Right 285 μV, left 200 μV.

One gave high normal values: Right 500 μV, left 510 μV.

This last case was an initial one, with a monophasic dark-adaptation curve (final global threshold: log asb 0·05—2·00) and an extinguished ERG as in the other cases.
In a case of atrophy gyrata, a young girl with typical fundus modifications and myopia, mild concentric contraction of the visual fields, high monophasic dark-adaptation-curve (global threshold after 20 min: log asb 0.7—3.0), and an abolished ERG, the values were 335 μV for the right eye and 360 μV for the left eye.

In a case of abiotrophic juvenile macular degeneration, with normal ERG, objective and subjective CFF, and dark-adaptation curve, the values were again normal: 375 μV right eye and 400 μV left eye.

In a case of unilateral senile macular degeneration (retinitis disciformis) the result was 220 μV on the affected side and 330 μV on the sound side.

(4) Chorio-retinitis.—Among our cases of cicatrical diffuse or disseminated chorio-retinitis, two showed a normal EOG (the first had a normal and the second a diminished ERG), and three others gave pathological values. The first of these three cases had a bilateral, heavily pigmented, diffuse chorio-retinitis with impairment of central vision, central and paracentral scotomata, a slightly subnormal dark-adaptation curve, and a subnormal b-wave in the ERG. The values of M were right eye 50 μV, left eye 260 μV.

In the second case, the lesions were circumscribed at the posterior pole, the values of M were 145 μV in the right eye and 125 μV in the left eye.

The third patient was a woman aged 42 years with very pigmented lesions, macular choroidal sclerosis, abolition of central vision in the right eye (left eye: 1.0), central and paracentral scotomata, a normal red-green colour-sense, a high biphasic retarded dark-adaptation curve with a subnormal final threshold (log asb 0.6—5.0), a diminished ERG (amplitude of the b-wave with the apparatus of Karpe: 50 μV), and a reduced objective CFF. The values of M were 210 μV in the right eye and 250 μV in the left.

Two cases of toxoplasmic cicatrical central necrotizing chorio-retinitis had a normal EOG.

We examined a case of secondary pigmentary retinitis after measles. The EOG was at the inferior limit of the physiological range in the right eye (265 μV) and gave a normal value in the left eye (360 μV). This case presented in the beginning an extinguished ERG; one year later, there was an isolated negative deflection (P III). The visual fields presented an annular scotoma and there was a marked hemeralopia.

A case of retinitis septica, with healed secondary retinal detachment of the left eye, showed values of 330 μV in the right eye and 140 μV in the left eye. The ERG was absent in both eyes.

(5) Myopia Gravis and Myopigenic Retinal Detachment.—In four cases of high myopia with pronounced choroidosis the values of the EOG were normal (470 and 450 μV; 335 μV; 400 and 500 μV; 490 and 420 μV).

Five eyes with actual or old, unhealed total detachment showed very low pathological values (145 μV; 50 μV; 50 μV; 170 μV; 250 μV).
Two cases of healed detachment were examined. The first (Fig. 4) showed a diminished value (200 μV; 500 μV for the sound eye), and the second showed normal values (470 μV) although the ERG was absent.

(6) Vascular Disturbances.—We obtained normal values in cases of hypertensive retinopathy, diabetic retinopathy, embolism of the central artery, periphlebitis, pre-retinal haemorrhage, and diffuse choroidal sclerosis.

(7) Glaucoma.—We also obtained normal values in cases of wide-angle, narrow-angle, and traumatic glaucoma. In a case of congenital glaucoma we found a subnormal value in one eye (210 μV), and in another case a normal value (470 μV), the other eye being enucleated.

A patient with very advanced wide-angle glaucoma, no visual perception in the right eye, and a thrombosis of the central vein with haemorrhagic glaucoma in the left eye gave the following results: right eye 160 μV, left eye 175 μV. The ERG was subnormal for both eyes and the dark-adaptation curve in the left eye highly pathological.

(8) Optic Nerve and Higher Optic Pathways.—We found normal values in cases of papillitis and retrobulbar neuritis, and in a case of cerebral thrombosis with homonymous triquadrantanopsia, acquired total colour blindness, deuteranopic anomalososcopic findings, and normal spectral luminosity.

Summary

With a standard method, we used electro-oculography as an indirect estimation of the standing potential in conjunction with other functional tests in normal and pathological cases. The normal range \((m \pm 2\sigma)\) of the mean deflection \(M\) lies between 285 and 525 μV. Too low values were principally found in cases of microphthalmos, primary pigmentary degeneration, cicatrical chorio-retinitis, and retinal detachment. There is no direct correlation with the ERG, which seems to be more rapidly affected.

REFERENCES


