A new colour vision arrangement test to detect functional changes in diabetic macular oedema

N Maár, M Tittl, M Stur, B Zajic, A Reitner

Abstract

Aim—A study was undertaken to investigate the correlation between colour discrimination tests and the presence of macular oedema in patients with type I diabetes to find a sensitive diagnostic tool for the detection of early functional changes.

Methods—The study was performed in 39 type I diabetic patients, 10 with and 29 without macular oedema. The examination included biomicroscopy, fundus photography of the macula, videofluorescein angiography, the LogMAR visual acuity chart, Farnsworth-Lanthony desaturated D-15 test, and the new Mollon-Reffin “Minimalist” test for colour vision deficiencies version 6.0.

Results—A highly significant correlation was found between the tritan value of the Mollon test and the presence of clinically significant macular oedema (p<0.0015), with a high sensitivity (88.9%) and specificity (93.3%). The DD-15 test was not significant (p=0.345) and showed low sensitivity for the presence of macular oedema (36%). All variables concerning the grading of macular oedema showed a highly significant association with the tritan values of the Mollon test (p<0.0001).

Conclusion—The results suggest that the Mollon-Reffin “Minimalist” test version 6.0 is the best colour discrimination test for detecting macular oedema, with higher specificity and sensitivity than the other methods used in the study.


Macular oedema is a major cause of visual loss in diabetic patients.1 The Wisconsin epidemiological study found an incidence of 20.1% over a period of 10 years for macular oedema in the younger onset group.2 Photocoagulation treatment decreases the risk of visual impairment,3 and an early diagnosis of diabetic macular oedema is essential for optimal prevention of functional loss. Macular oedema is difficult to detect by routine ophthalmological examination, visual testing, or biomicroscopy.4 Fluorescein angiography carries the risk of an allergic reaction and the required set up is not always available. It is therefore necessary to develop an additional non-invasive diagnostic method to evaluate retinal dysfunction caused by diabetic macular oedema.

Several studies5–12 have reported reduced colour discrimination in patients with diabetic retinopathy of all degrees of severity, but there has been no analysis of the influence of macular oedema. Later studies9–12 have investigated colour discrimination in patients with diabetic macular oedema using the Farnsworth-Munsell 100 hue test or the Farnsworth-Lanthony desaturated D-15 (DD-15) test and a more pronounced tritan defect was found in affected eyes.

We have examined whether a correlation exists between the degree of macular oedema of diabetic patients and the results of several functional tests, including colour discrimination as tested with the Farnsworth-Lanthony DD-15 test and the Mollon-Reffin “Minimalist” test.

Methods

SUBJECTS

The patient group was recruited prospectively from our outpatient department of retinal vascular and macular diseases and consisted of 16 men and 23 women aged 17–47 years. Thirty seven (95%) of the 39 patients had undergone intensive insulin treatment (baseline injection of long acting insulin, several injections of short acting insulin at mealtimes) and two had received conventional insulin treatment (one or two daily injections of long acting insulin), and for all but one patient a recent glycosylated haemoglobin (HbA1C) value could be obtained.

Admission criteria included type I diabetes with or without diabetic retinopathy and with best corrected visual acuity of >0.4 LogMAR (0.4 Snellen value). Because of the influence of nuclear sclerosis on colour vision, only patients younger than 50 years of age and with no lens opacities according to the Lens Opacities Classification System (LOCS) III13 were included. Only one eye of each patient was included in the statistical analysis. The eye to be studied was designated on the basis of the patient’s birth month: in patients born in the first half of the year the right eye was selected, and in those with a birth month in the second half of the year the left eye was used. In patients who had macular oedema in one eye only, this eye was designated for the study. Nineteen right eyes (49%) and 20 left eyes (51%) were included in the analysis, resulting in an equal distribution between left and right eyes.

Exclusion criteria were congenital colour vision deficiencies, cataract, glaucoma, any retinopathy, new vessels, or chorioretinal scars in the macula. We also excluded patients with more than mild proliferative diabetic retinopathy and patients who had a history of intraocular surgery or laser therapy, because patients can have a tritan defect as a side effect of laser therapy.14 15

Department of Ophthalmology, University of Vienna, Austria
N Maár M Tittl M Stur A Reitner

Institute for Statistics, Operations Research and Computer Analysis, University of Vienna, Austria
B Zajic

Correspondence to:
Dr N Maár, Univ Klinik für Augenheilkunde, Allgemeines Krankenhaus der Stadt Wien, Währinger Gürtel 18–20, 1097 Vienna, Austria
Noemi.Maar@univie.ac.at

Accepted 12 June 2000
STUDY DESIGN

Ophthalmological examination included best corrected visual acuity in LogMAR units using Bailey-Lovie charts numbers 4 and 5 (National Vision Research Institute, Australia), and colour discrimination testing using the Farnsworth-Lanthony desaturated D-15 test and the Mollon-Reffin “Minimalist” test version 6.0.16 The chromaticities used are indicated in the original paper17 so anyone interested can construct their own copy of the test. For identification of subjects with congenital colour vision deficit and their exclusion from our study a Nagel anomaloscope (Schmidt and Haensch) was used. The Log-MAR charts were used at a distance of 3 m with direct illumination of 60–120 cd/m². The colour discrimination tests were conducted using an Osram AG lamp at a temperature of 5400 Kelvin and an illumination of 1200–1500 lux, 50 cm above the colour test. Both tests were first performed binocularly for training purposes and then with the right and left eyes separately.

The Mollon-Reffin test contains a set of grey chips of varying lightness which serve as the background chips, a set of coloured chips, the probe chips, and an orange demonstration chip. Five of the grey chips are placed randomly on the black Plexiglass. To these the examiner first adds the orange chip, which does not lie on any confusion line, mixes it with the grey chips, and invites the patient to identify the “coloured chip” by touching with a pointer. If the patient successfully identifies the orange chip, the examiner draws a probe chip from the middle of the proton series. After correct identification of this probe, the examiner then moves forwards along the confusion line and presents the least saturated chip; if, on the other hand, the response to the first proton probe is incorrect, the examiner moves backwards to the most saturated chip. On subsequent trials a simple staircase procedure is used to establish the maximal chroma at which the patient fails. The same process is then repeated for the tritan and deutan axis.

Colour discrimination deficits detected with the DD-15 test were analysed using a software program for calculating the total colour difference score (TCDS).18 For the Mollon-Reffin test the score value was defined as the number of the reliably identified coloured chip for each confusion line.

The intraocular pressure was measured with an applanation tonometer. After the pupil was maximally dilated with 0.5% tropicamide and phenylephrine, the retina was carefully examined by biomicroscopy with a Volk 90D lens. The biomicroscopic finding was documented using the ETDRS final retinopathy grading form.20 The presence of clinically significant macular oedema (CSMO) according to the ETDRS criteria was established by biomicroscopy. Stereoscopic colour fundus photographs of the standard field no 2 were taken with a fundus camera (Canon CF-60UV with Kodak Ektachrome 100HC) to classify the macular oedema. Fluorescin angiography was recorded using a scanning laser ophthalmoscope (Rodenstock, Germany). A bolus injection of 5 ml of 10% fluorescein sodium (Braun Melsungen, Germany) was injected into the antecubital vein and the angiograms were recorded on an S-VHS video recorder (Panasonic 7330).

The fundus photographs and the angiograms were graded by two independent examiners and, where there were differences of opinion, these were resolved by discussion. For the grading of the fundus photographs a transparent overlay with a circle of 1 disc diameter (DD) radius was fixed over the image. By using a Donaldson 5× stereoscopic viewer the degree of macular oedema was determined using the modified Airlie House classification.19 As morphological variables the following lesions were graded: degree of CSMO (1 = no evidence, 2 = questionable involvement, 3 = size of oedema >1 disc area, a part <1 DD from the centre, 4 = thickening/hard exudates <500 µm from the centre); hard exudate at the centre (1 = no evidence, 2 = questionable involvement, 3 = definite involvement by hard exudate); size of macular retinal thickening <1 DD from the centre (1 = no evidence, 2 = questionable involvement, 3 = size of thickening <0.5 disc area, 4 = size of thickening <1 disc area, 5 = size of thickening <2 disc area, 6 = size of thickening = 2 disc area); and maximal retinal thickness at the centre (1 = no evidence, 2 = questionable involvement, 3 = thickness of <1 × reference, 4 = thickness of 1 × reference but <2 × reference, 5 = thickness of 2 × reference but <0.5 DD, 6 = thickness ≥0.5 DD). The videos of the angiograms were studied on a black and white high resolution monitor. The classification of the angiogram was performed in field 2 with a 20° central subfield using the ETDRS fluorescein angiogram grading form.20

STATISTICS

For statistical analysis the software program STATVIEW version 4.5 (Abacus Concepts Inc) was used. Statistical tests including Kendall correlation, analysis of variance (ANOVA), and multiple and logistic regression were used to analyse patient data, particularly for significance of correlation between the results of the functional tests, on the one hand, and age, duration of diabetes, duration of the intensified insulin treatment, HbA₁c stage of retinopathy, and pathological changes detected in the macula; p values of <0.05 were considered to be statistically significant. All tests were used in a two tailed manner.

To determine the sensitivity and specificity the χ² test was used. Sensitivity represents the probability of a test giving a positive result when the disease is present, and specificity is the probability of a negative result when the disease is absent.

Results

Most of the eyes were classified as having low levels of retinopathy; 12 (31%) had no diabetic retinopathy, nine (23%) had early, nine (23%) had mild, four (10%) had moderate, three (8%) had severe non-proliferative diabetic retinopathy and two (5%) had mild prolifera-
Table 1 Mean (SD) data of patients with and without clinical significant macular oedema (CSMO)

<table>
<thead>
<tr>
<th></th>
<th>With CSMO (n=10)</th>
<th>Without CSMO (n=29)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>33.7 (7.75)</td>
<td>28.07 (5.67)</td>
</tr>
<tr>
<td>Duration of diabetes (years)</td>
<td>22.8 (7.00)</td>
<td>12.31 (7.22)</td>
</tr>
<tr>
<td>HbA1c (%)</td>
<td>6.22 (2.33)</td>
<td>6.94 (0.68)</td>
</tr>
<tr>
<td>Functional tests</td>
<td>0.07 (2.01)</td>
<td>0.6 (0.17)</td>
</tr>
<tr>
<td>TCDs</td>
<td>2.1 (0.74)</td>
<td>1.03 (0.19)</td>
</tr>
<tr>
<td>FAZ outline</td>
<td>144.8 (23.34)</td>
<td>132.23 (28.44)</td>
</tr>
<tr>
<td>Cystoid spaces</td>
<td>3.8 (0.42)</td>
<td>1.0 (0.00)</td>
</tr>
<tr>
<td>Leakage</td>
<td>1.0 (0.00)</td>
<td>1.0 (0.00)</td>
</tr>
<tr>
<td>Capillary dilatation</td>
<td>3.3 (1.42)</td>
<td>0.07 (0.37)</td>
</tr>
</tbody>
</table>

Fluorescence angiography

- FAZ size
- FAZ outline
- Cystoid spaces
- Leakage
- Capillary dilatation

Dependence on age and duration of diabetes

Significant correlation was found between age and the DD-15 colour discrimination tests (p<0.0001; Kendall), as well as between age and capillary loss (p=0.0236), and age and capillary dilatation (p=0.0124). The degree of macular oedema was not significantly correlated with age.

None of the tests showed significant correlation with duration of diabetes. Generally, a higher significance was established between duration of diabetes and the morphological variables, but this trend was also age dependent (capillary loss, p=0.0001; capillary dilatation, p=0.0019; leakage, p=0.0181; but CSMO, p=0.189; Kendall). Patients with CSMO or more severe retinopathy had a significantly longer duration of diabetes (p=0.0003 and p<0.0001; ANOVA).

Dependence on retinal morphology

Patients with macular oedema had poorer visual acuity (p=0.692) and a higher TCDs (Fig 1, p=0.345) but only the Mollon T score differed significantly (Fig 2, p=0.0015; logistic regression r² = 0.565). Table 2 shows the p values and adjusted r² of the correlation between the T value of the Mollon-Reffin test and the morphological variables of the retina. Thus, it can be asserted that the T score of the Mollon-Reffin test showed a significant corre-
with the FM-100 test, macular oedema, capillary loss and leakage are correlated with the error score. The highest significance level was associated with leakage, which is similar to our results. These authors and an ETDRS report\textsuperscript{12} have reported that the tritan axis is more severely affected than the protan and deutan axes in patients with diabetic maculopathy and that this tritan-like defect increases in magnitude with increasing severity of macular oedema. Hudson et al\textsuperscript{25} found increased sensitivity of short wavelength pathways in detecting CSMO. The position of the localised field loss corresponded to the clinical mapping of the extent of the oedema. A correlation between the selective loss of the short wavelength pathway sensitivity and the severity of diabetic macular oedema has already been demonstrated using an incremental threshold technique.\textsuperscript{26}

Macular oedema reduces the transmission of light to the photoreceptors.\textsuperscript{27} This might affect the blue rather than the red-green mechanism as a result of the lower density and number of blue cones in the human fovea.\textsuperscript{28} Another possible explanation for the tritan colour defect—similar to retinal detachment\textsuperscript{29–31}—might be the oblique orientation of the photoelements that occurs after retinal detachment\textsuperscript{32} and other retinal pathologies.\textsuperscript{33}

The best correlation was found between the presence of macular oedema and the Mollon-Re\textsuperscript{“Minimalist”} test version 6.0. The DD-15 test only identifies the confusion axis for blue-yellow defects while the Mollon-Re\textsuperscript{“Minimalist”} test examines the tritan axis separately, directly along the confusion line. This is probably the reason for the higher sensitivity and specificity of the Mollon-Re\textsuperscript{“Minimalist”} test for detecting acquired colour discrimination defects in the tritan axis. With regard to the sensitivity and specificity, we found that the Mollon-Re\textsuperscript{“Minimalist”} test was the only colour discrimination test that might be appropriate for clinical use. In our study we included only patients without any lenticular opacities. Since patients with diabetes are more likely to develop cataracts than those without,\textsuperscript{34} a blue-yellow colour defect indicated from the lenticular opacities may occur earlier in this group than in the normal population. The use of blue-yellow colour vision tests without examination of the lens in diabetic patients older than 30 is therefore inadvisable.

In summary, our data indicate the usefulness of a simple colour discrimination method, the Mollon-Re\textsuperscript{“Minimalist”} test version 6.0, as part of the screening and follow up examination for macular oedema in young patients with juvenile onset diabetes.

Proprietary interest: none.


