The objective of the test, instrumentation, procedure, and what information obtained from a visual field test depends on the ability of the patient to reliably perform the test. Standardised reliability criteria have been adopted and consist of fixation loss rate less than 20%, false positive response rate (FP) less than 33%, and false negative response rate (FN) response rate less than 33% of test catch trials.

A number of studies have shown that 29–45% of full threshold SAP test results are unreliable using these standardised reliability indices, with most of the unreliable fields attributable to fixation losses. Katz et al found that 19% of normals, 28% of ocular hypertensives, and 37% of glaucoma patients were unreliable on their first C30–2 full threshold field. Studies of continuous patient monitoring during testing have shown either no significant difference or a positive group effect, with no effect on individual reliability indices in a more recent study.

It is also possible that test duration may influence reliability and, in particular, fatigue has been shown to influence reliability in glaucomatous subjects.

The importance of adequate and careful patient instruction both directly and indirectly by training of technicians has frequently been emphasised as a factor playing a major part in obtaining a reliable result. Moreover, perimetrists’ instructions have been shown to significantly affect obtained automated perimetry thresholds.

The aim of this study was to evaluate the effect of a patient information video on visual field test result reliability. The video was designed for patients who had not previously performed a visual field test and provided information on the objective of the test, instrumentation, procedure, and what would be expected of them.

MATERIALS AND METHODS
The study was performed in a hospital eye service “new patient” glaucoma clinic. The inclusion criteria were new referral, no previous threshold visual field tests, absence of hearing or cognitive impairment, understanding of English language, and best corrected visual acuity of 6/36 or better in both eyes.

After informed consent consecutive, eligible patients attending the clinic were randomised in to either a control or “video” group. The control group proceeded normally through the clinic, which involved a routine visual field test and a subsequent consultation with the clinician. The video group were individually shown the standard information video and then preceded in the same manner as the control group. All patients having field tests would receive instruction from the technician monitoring the test. The level of monitoring was at the discretion of the technician. Technicians performing the visual field tests were masked to each patient’s randomisation status and patients were instructed not to disclose whether they had been shown the video.

The audiovisual patient information video was produced in house by the audiovisual unit and was 4.5 minutes in duration. It explained the purpose, rationale, and events surrounding a standard automated visual field test, with emphasis on the sources of unreliable visual field results and a visual representation of the perimeter bowl as perceived by the patient, including the fixation target and test stimuli.

Visual fields were performed by one of seven technicians, who supervised tests on both eyes of the patient. The following data were recorded for all the patients: age, sex, whether or not a visual field test were performed by a referring optometrist, technician ID, best corrected visual acuity (BCVA), diagnosis, duration of visual field test, fixation loss rate (FL), false positive response rate (FP), false negative response rate (FN), mean deviation (MD), pattern standard deviation (PSD), and glaucoma hemifield test (GHT) result. Standard reliability criteria were employed: fixation loss rate less than 20%, false positive response rate less than 33%, and false negative response rate less than 33%.

The study was published in the journal "British Journal of Ophthalmology."
Patient diagnosis was recorded under five categories: normal ocular examination; glaucoma suspect; glaucoma, including primary open angle and normal tension glaucomas; ocular hypertension. The fifth category labelled “other” included all miscellaneous diagnoses such as cataract, age related maculopathy, congenital disc anomalies, amblyopia, and other unconfirmed diagnosis.

Ethical approval for the study was obtained from the United Bristol Healthcare research and ethics committee.

Statistical analysis
The demographics of the video and control groups were compared using the unpaired two sample t test for continuous variables and the Pearson χ² test for nominal variables where proportions were compared. The paired t test was used to compare field parameters of the patients’ right and left eyes.

RESULTS
Of the 306 consecutive patients attending the new patient clinic, 244 patients were eligible for inclusion. One hundred and thirty two patients were randomised to the control group and 112 to the video group. Summary data for the video and control groups are provided in Table 1. No significant differences were found between these groups for age, sex and BCVA. Furthermore, there was no significant difference (p = 0.356) in the proportion of diagnoses in either group (Fig 1).

There was a considerable spread of visual field defect magnitudes in both groups (Table 1). The MD or PSD for the right and left eye were not significantly different between the groups.

The reliability results are shown in Table 2. Because the results of reliability of either eye has a direct impact upon the management of the patient, reliability was expressed by eye and whether a patient had reliable visual fields in both eyes. In the control group 81 (61.4%) patients had reliable fields in both eyes. In the video group 85 (75.9%) patients had reliable fields in both eyes. The difference in reliability in the two groups was significant (p = 0.015).

When only the right (first tested) eye was considered, 93 (83.0%) of the eyes in the video group were reliable compared to 106 (80.3%) in the control group. The difference was statistically significant (p = 0.011).

![Figure 1 Patient diagnosis within the glaucoma and video group.](image)

Table 1  The demographic data on the video and control group. Note that no significant differences existed between groups for any of the variables

<table>
<thead>
<tr>
<th>Control</th>
<th>Video</th>
<th>Significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>132</td>
<td>112</td>
</tr>
<tr>
<td>Age (years)</td>
<td>62.6 (14.4)</td>
<td>62.7 (13.2)</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>70 (53%)</td>
<td>57 (50.9%)</td>
</tr>
<tr>
<td>Female</td>
<td>62 (47%)</td>
<td>55 (49.1%)</td>
</tr>
<tr>
<td>Optician fields</td>
<td>106 (80.3%)</td>
<td>96 (86.5%)</td>
</tr>
<tr>
<td>Right eye</td>
<td></td>
<td></td>
</tr>
<tr>
<td>BCVA &gt;20/80</td>
<td>104 (92.0%)</td>
<td>125 (94.7%)</td>
</tr>
<tr>
<td>MD (SD) dB</td>
<td>−2.98 (5.24)</td>
<td>−2.52 (4.06)</td>
</tr>
<tr>
<td>PSD (SD) dB</td>
<td>3.22 (2.71)</td>
<td>2.89 (2.30)</td>
</tr>
<tr>
<td>Left eye</td>
<td></td>
<td></td>
</tr>
<tr>
<td>BCVA &gt;20/80</td>
<td>103 (92%)</td>
<td>122 (92.4%)</td>
</tr>
<tr>
<td>MD (SD) dB</td>
<td>−2.77 (4.58)</td>
<td>−2.57 (3.80)</td>
</tr>
<tr>
<td>PSD (SD) dB</td>
<td>3.26 (2.92)</td>
<td>3.12 (2.61)</td>
</tr>
</tbody>
</table>

BCVA = best corrected visual acuity, MD = mean deviation, PSD = pattern standard deviation, NA = non-applicable.

*Unpaired t test.

†Two tailed Pearson χ² test.

Table 2  The number and percentage of patients in the video and control groups who had a reliable field test in both eyes and in each eye separately

<table>
<thead>
<tr>
<th></th>
<th>Reliable</th>
<th>Unreliable</th>
<th>Significance*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Both eyes</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Video</td>
<td>85 (75.9%)</td>
<td>27 (24.2%)</td>
<td>0.015</td>
</tr>
<tr>
<td>Control</td>
<td>81 (61.4%)</td>
<td>51 (38.6%)</td>
<td></td>
</tr>
<tr>
<td>Right eye</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Video</td>
<td>93 (83.0%)</td>
<td>19 (17.0%)</td>
<td>0.583</td>
</tr>
<tr>
<td>Control</td>
<td>106 (80.3%)</td>
<td>26 (19.7%)</td>
<td></td>
</tr>
<tr>
<td>Left eye</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Video</td>
<td>97 (86.6%)</td>
<td>15 (13.4%)</td>
<td>0.011</td>
</tr>
<tr>
<td>Control</td>
<td>97 (73.5%)</td>
<td>35 (26.5%)</td>
<td></td>
</tr>
</tbody>
</table>

*Two tailed Pearson χ² test for significance of difference in reliability between the control and video group.
have artefactually improved their testing standard, albeit up bias, whereby technicians’ awareness of the project may affect the first tested eye. It is also possible that the video may reduce other means. It may be postulated that information provided by performing a field test cannot be adequately substituted by

The majority of the visual field tests, 214 (87.7%), were supervised by four of the technicians (Fig 2). There was no significant difference between the proportion of visual fields performed by each technician within the video and control group (p=0.254).

DISCUSSION
This study has demonstrated that 75.9% of patients watching an educational video had reliable visual fields in both eyes on their first attempt, which represented a significant improvement in reliability compared to the control group, of whom 61.4% had reliable fields in both eyes. Assessment of reliability of visual fields by patient, rather than by eye has practical implications for patient management, as this is likely to be affected by the either eye’s reliability.

The introduction of a standardised information video ensures that key points are brought to the attention of every patient and serves to reinforce technician instructions. The video aimed to clearly explain to the patient how to correctly perform the visual field test. This would entail emphasising the importance of maintaining fixation, not guessing a response, and resisting the tendency to be “trigger happy” with responses. Another aim was to clarify some of the ambiguities arising during the first visual field test, such as reminding the patient that although they should maintain fixation, they are allowed to blink during the test, that a pupil is acceptable and should be independent of the brightness of the stimulus and only based on whether a given stimulus is seen or unseen. The final aim of the video was to reassure the patient of some of the sources of stress and anxiety associated with the unfamiliarity of the patient with SAP such as the experience of transient darkening and subjective loss of sensitivity to the perimeter bowl.

It is of interest that the impact of the information video on the reliability of the visual field of the right (first tested) eye was not significant. This would suggest that the video achieves its effect by reducing the rate of unreliable fields in the second tested (left) eye. It is likely that familiarity with SAP achieved by performing a field test cannot be adequately substituted by other means. It may be postulated that information provided by the video reinforces the learning experience gained with the first tested eye. It is also possible that the video may reduce any fatigue effect.

The design of this study may have produced some “work up” bias, whereby technicians’ awareness of the project may have artefactually improved their testing standard, albeit unintentionally. A historical cohort of fields performed by the same technicians may have enabled a quantification of this effect.

During the design of the study it was recognised that the lack of standardisation of the technician instructions may introduce differences between the control and video group and arguably weaken the strength of conclusions. In anticipation of this, the randomised, control study design was employed to minimise such effects and also the effects of any unanticipated confounding variables. A rigid standardisation of technicians’ instructions, although possible, was not considered representative of typical hospital eye service clinics. It is acknowledged that some variability in the quality of instructions and supervision provided by technicians would be inevitable, but the absence of any significant difference in the proportion of fields performed by each technician within the two groups (p = 0.254) makes this unlikely.

The level of the perimetric experience before recruitment into the study may also have produced an element of learning effect for future field tests. Eighty two per cent of referred patients were reported to have had a field test by the referring optometrist. For the purpose of this study it was acceptable for patients to have been exposed to a single field test not performed in an ophthalmic clinic, as most UK optometrists use suprathreshold screening strategies, rather than thresholding algorithms.8 The video and control group did not significantly differ in the proportion of patients in each group who had performed a visual field test at the referring optometrist’s practice.

The benefits of careful patient instruction by technicians performing visual field tests is not a novel idea and has been repeatedly and frequently advocated. The constraints of time and resources, however, limit the extent and quality of information delivered to patients during routine visual field testing. The incorporation of a video guiding and reassuring the patient on taking the visual field test is an effective way of using available clinic time. A reduction in the number of patients requiring attendance for a “repeat visual field” can reduce demand on this frequently used service.

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ECHO

Hyperactive neurones conjure up hallucinations

A

n Australian professor has put forward a hypothesis for the origin of hallucinations which, if correct, would help their diagnosis and possible treatment. His thinking stems from personal experience of simple Charles Bonnet hallucinations.

These were brought about by the development of a macular hole <400 µm in the left eye four years after one >400 µm in the right. Hallucinations started 11 weeks afterwards, when acuity in the left eye was 6/12–6/18. They were simple black and white, non-evoked, geometric arrays. The first looked like “brickwork” (fig Aa-c) within an area <1°. Later came arrays of arches angled at 45°. Then after seven days groups of dark spots appeared (fig B; a, b) and then lozenges angled at 45° (fig C). Finally, all but the arches and previously seen “flashes” faded about 10–12 days after the hallucinations first began. Faint hallucinations returned about 38 days later and lasted 10 days or so. Brickwork reappeared briefly during an episode of macular cystoid oedema in the left eye.

Professor Burke extends current thinking that links particular hallucinations with particular areas of the brain, deducing that they result from “deafferentation” of visual structures in the brain or silencing of the principal afferents to them. This induces changes leading to increased excitability of affected neurones and spontaneous activity—perceived as hallucinations. As the neurones gradually become reactivated the hallucinations fade and vanish.

Charles Bonnet hallucinations occur after injury to the brain or other parts of the visual system, most commonly after age related macular degeneration.

Please visit the British Journal of Ophthalmology website [www.bjophthalmol.com] for link to this full article.