Clinical Ophthalmology

Methodology

Visual field examination method using virtual reality glasses compared with the Humphrey perimeter

This article was published in the following Dove Press journal:
Clinical Ophthalmology
7 August 2017
Number of times this article has been viewed

Stylianos Tsapakis
Dimitrios Papaconstantinou
Andreas Diagourtas
Konstantinos Droutsas
Konstantinos Andreanos
Marilīta M Moschos
Dimitrios Brouzas

1st Department of Ophthalmology,
National and Kapodistrian University of Athens, Athens, Greece

Purpose: To present a visual field examination method using virtual reality glasses and evaluate the reliability of the method by comparing the results with those of the Humphrey perimeter.

Materials and methods: Virtual reality glasses, a smartphone with a 6 inch display, and software that implements a fast-threshold 3 dB step staircase algorithm for the central 24° of visual field (52 points) were used to test 20 eyes of 10 patients, who were tested in a random and consecutive order as they appeared in our glaucoma department. The results were compared with those obtained from the same patients using the Humphrey perimeter.

Results: High correlation coefficient ($r=0.808, P<0.0001$) was found between the virtual reality visual field test and the Humphrey perimeter visual field.

Conclusion: Visual field examination results using virtual reality glasses have a high correlation with the Humphrey perimeter allowing the method to be suitable for probable clinical use.

Keywords: visual fields, virtual reality glasses, perimetry, visual fields software, smartphone

Introduction

Automated perimetry is a useful method to assess visual fields in many ophthalmic and neurological diseases. Current perimeters are accurate, but they have a number of disadvantages. Visual field testing is a time-consuming process. It is inconvenient and stressful for debilitated, claustrophobic, ill, or elderly patients to keep their heads still in the perimeter bowl throughout the test. To overcome these problems, visual field testing using a video projector has been proposed. The majority of computerized perimeters are specialized pieces of hardware/software. They typically consist of a projection area, an embedded microcontroller, an input device for the operator, and a button for the patient. These devices, built for physicians’ offices or hospitals, are bulky, heavy, and expensive. They are not portable, and they cannot be used at bedside. However, smartphones are found everywhere, and they are inexpensive. Virtual reality (VR) glasses have some advantages in visual field testing. They are lightweight, portable, comfortable, and affordable, and there is no need for an eye patch.

The possibility of using VR glasses for visual field testing has been described since 1998, patent no: US5737060A. However, at that time, hardware and software was an issue. Smartphones and similar portable devices were not as improved as they are today. VR glasses for smartphones did not exist. Win98 was actually just a shell over DOS.2–4

Correspondence: Dimitrios Brouzas
10 G Papandreou Street, Byron,
Athens 16231, Greece
Tel/fax +30 21 0765 2909
Email brouzas@yahoo.com

DOI: http://dx.doi.org/10.2147/OPTH.S131160

Clinical Ophthalmology 2017:11 1431–1443

© 2017 Tsapakis et al. This work is published and licensed by Dove Medical Press Limited. The full terms of this license are available at https://www.dovepress.com/terms.php
and incorporate the Creative Commons Attribution – Non Commercial (unported, v3.0) License (http://creativecommons.org/licenses/by-nc/3.0/). By accessing the work you hereby accept the Terms. Non-commercial use of this work is permitted without any further permission from Dove Medical Press Limited, provided the work is properly attributed. For permission for commercial use of this work, please see paragraphs 4.2 and 5 of our Terms (https://www.dovepress.com/permissions).
Commercially available visual reality glasses with built-in displays do not perform well. These VR glasses are usually built for gaming, and the display is usually small with low resolution. This requires moving the fixation point, which confuses older patients, whereas custom-built VR glasses with bigger displays are more expensive and lack standardization. For these reasons, widespread testing of visual field using VR glasses has been limited.

Today’s smartphones are much more powerful, affordable, have bigger displays, and standardization can be achieved by selecting proper hardware/software.

Materials and methods

To test the reliability of visual fields using visual reality glasses, 20 eyes of 10 patients, who were chosen randomly and consecutively at our glaucoma department, were tested successively using a Humphrey perimeter and the VR glasses method within hours for comparison. Approval was obtained from the Ethical Committee of the General Hospital of Athens “G Gennimata”. Written informed consent was obtained from all patients in the study.

Trust EXOS 3D VR glasses and Alcatel One Touch Pixi 4 (6) 8050D smartphone with 6 inch display were used. The patients were allowed to wear his/her glasses during testing if they felt it was necessary (Figure 1A–C).

Virtual display focus distance is adjusted with the 2 rotating knobs on the sides. Trial glasses were not used as the patient could wear his/her glasses during testing if necessary (Figure 1C).

Proprietary software implementing a fast-threshold 3 dB step staircase algorithm at central 24°/52 points of visual field was used for the purpose of testing (Figure 2). The projected stimuli intensity was distributed on a logarithmic scale.

The typical luminosity of a LCD screen is 250 cd/m<sup>2</sup>. The results of a visual field test depend on the luminosity of the examination display. As different smartphone models may be used for visual field testing, the luminosity of a display must be adjusted in order to make sure that the data are consistent from one visit to another and between successive tests. This allows for the data to be analyzed over time and between different installations.

Contrast ratio is the ratio of luminance between the brightest white and the darkest black that can be produced. Brightness sets the black point and determines the low light output level (black level) of the display.

Gamma describes the relationship between the pixel level and the luminance of the monitor (the light energy it emits).

LCDs are considered linear devices; therefore, technically they do not need gamma correction. Gamma correction, however, corrects for the deficiencies (non-linearity) of cathode ray tube (CRT) monitors.

The software uses gamma 1.0 because LCDs are linear but gamma is adjustable to match the viewing system’s gamma for optimum performance (Figure 3).

The VR glasses gamma is set separately (Figure 4).

The display’s gamma/brightness/contrast can be visually calibrated. Visual calibration is sufficiently reliable to be used as an alternative to calibration using an expensive photometer. The software uses a gray scale step wedge for display adjustment. The settings should be set to a point that makes the shades of gray distinct and clearly visible (Figure 5A–D).

In our case, for better accuracy and comparability, a photometer was used and the luminosity of white color was set
at 130 Lux (approximately 410 asb). This was about 50% of the maximum available brightness for the smartphone used (Alcatel Pixi 4(6) 8050D).

Software features
1. Fast-threshold, 3 dB step staircase strategy, 52 points, central 24° of visual field.
2. The software uses the Heijl–Krakau blind spot method to monitor fixation. The software detects the blind spot by projecting stimuli at maximum luminosity at expected blind spot locations until finding the correct response.
3. The software pauses the test in case of fixation loss.
4. Supra threshold stimuli are used to check for false negative results. The software also checks for false positive responses.
5. Variable stimuli presentation rate, adjusted to patient’s response time.
6. Stimuli presentation time 250 ms.
7. Initial patient’s response waiting time 500 ms, adjusted to patient’s response time.

The software includes eye tracking capability using AForge. NET computer vision and artificial intelligence language. The source code and binaries of the project are available under the terms of the Lesser GPL and the GPL (GNU General Public License). Pupil diameter and eye movements were not recorded during examination because they were not supported by the VR glasses used. The points are projected using proper trigonometry adjustment to compensate for the classical perimeter bowl of VR glasses so that stimuli appear on the retina as if they were projected from a classical bowl perimeter (Figure 6).

Examination procedure
During testing, the patient should sit comfortably, put on the VR glasses, and adjust the head straps. The VR glasses should not be tilted, off-center, too high, or too low. Pupil distance should be adjusted with the rotating knob on top. To optimize image quality, focus distance should be adjusted with the 2 rotating knobs on both sides of the VR headset until the picture is sharp.

The VR glasses should be positioned appropriately to avoid lens rim artifact (LRA), which can sometimes be confused as nasal step scotomas. According to a study in central static threshold visual fields (Humphrey 30-2 Program) performed with a corrective lens, LRA was present in 10.4% of 704 fields examined retrospectively and 6.2% of 276 fields evaluated prospectively. If LRA occurs in one of our patients. If it occurs, then the test should be repeated with better placement of the VR glasses (Figure 7).

Figure 2 Computer – Virtual reality glasses – computer setup.

Figure 3 Gamma correction adjustment for PC.
To avoid LRA, the software allows the doctor to project all stimuli (at maximum intensity so that all points are clearly visible, provided there is no absolute scotoma) and make appropriate adjustments. In most cases, this is enough (Figure 8).

The software locates the blind spot automatically and adjusts the location and size of the test points. Furthermore, the location and size of test points can be set manually.

Each eye was tested separately, and no eye patch was used. During testing, the patient should stare at the central fixation point and click a mouse whenever he/she sees a visual stimulus on the display (Figure 9).

The patient is free to change position or move his/her head while testing. VR glasses are lightweight; they weigh ~385 g while the smartphone weighs ~179 g. The patient may use his/her hand to hold the VR glasses, making testing more comfortable.

Twenty eyes of 10 patients appearing randomly and consecutively at the visual fields lab were tested successively using a Humphrey perimeter and the VR glasses method within hours for comparison.

The results were statistically analyzed and compared. The patients tolerated the VR test very well. All the patients reported that it was much more comfortable compared to the standard bowl perimeter (Humphrey).

**Figure 4** Gamma correction adjustment for mobile device. **Abbreviations:** VR, visual reality; APK, Android package kit.

**Figure 5** Mobile device display adjustments and points to be tested. (A) gamma correction, (B) brightness adjustment, (C) left eye points, and (D) right eye points.
Figure 6 Trigonometrical projection to compensate bowl perimetry.

Statistical analysis

Point-to-point correlation coefficient (r) between the VR glasses and the Humphrey perimeter was computed for each eye and for all eyes together using the InStat version 3.05 of GraphPad Software, Inc. When the distribution of values was not normal, nonparametric Spearman correlation coefficient (r) was used.

VR glasses tests are 24° (52 points), whereas Humphrey tests are 30° (76 points). Only the corresponding (common 52 points) between these are taken into consideration (Figures 10–16).

Results

Table 1 Point to point Spearman coefficient (r) between the two methods for each eye

<table>
<thead>
<tr>
<th>Eye</th>
<th>Spearman correlation coefficient (r)</th>
<th>Standard deviation</th>
<th>P-value (one-tailed)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.736955</td>
<td>6.594795</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>2</td>
<td>0.765154</td>
<td>4.90298</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>3</td>
<td>0.875855</td>
<td>5.1637</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>4</td>
<td>0.792082</td>
<td>2.449182</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>5</td>
<td>0.773847</td>
<td>3.754133</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>6</td>
<td>0.75502</td>
<td>5.163674</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>7</td>
<td>0.865649</td>
<td>2.717742</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>8</td>
<td>0.83976</td>
<td>6.698726</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>9</td>
<td>0.838132</td>
<td>2.870508</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>10</td>
<td>0.766863</td>
<td>5.146533</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>11</td>
<td>0.870688</td>
<td>2.422245</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>12</td>
<td>0.848471</td>
<td>2.828427</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>13</td>
<td>0.850762</td>
<td>2.313561</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>14</td>
<td>0.889794</td>
<td>2.154654</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>15</td>
<td>0.745111</td>
<td>9.614359</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>16</td>
<td>0.829142</td>
<td>3.223862</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>17</td>
<td>0.725046</td>
<td>5.796804</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>18</td>
<td>0.806027</td>
<td>3.376511</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>19</td>
<td>0.879466</td>
<td>3.225733</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>20</td>
<td>0.722703</td>
<td>4.385763</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

Total results

<table>
<thead>
<tr>
<th>Mean Spearman correlation coefficient (r)</th>
<th>Mean standard deviation</th>
<th>P-value (one-tailed)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.808537</td>
<td>4.19494</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>
In each eye and in all eyes together, the mean difference value between the two methods was statistically significant at $P<0.0001$.

The correlation coefficient ($r$) in all tests between the two methods was statistically extremely significant at $P<0.0001$.

**Discussion**

VR glasses perimetry has many similarities to classical bowl perimetry. There are some differences due to the hardware used. In all bowl perimeters, the results are comparable to a significant degree, but they are not identical because each perimeter is different from others.

For example, in the Octopus perimeter, a 5 dB attenuation is equal to 316 asb, whereas in the Humphrey perimeter,
Evaluation of visual field examination method using virtual reality glasses

Figure 10 Results, eye 1–3.

Figure 11 (Continued)
a 5 dB attenuation is equal to 3160 asb. In Humphrey, 0 dB correspond to 10,000 asb, whereas in Octopus, 0 dB correspond to 1,000 asb stimulus. Such differences make comparisons more difficult between different devices. This justifies the statistical difference between the mean values of the VR glasses perimetry method and the Humphrey perimeter, yet the correlation coefficient (r) between the two methods was statistically extremely significant (r=0.808, P<0.0001; Table 1). For this reason, if we want the results to be comparable, then the same device should be used for consecutive tests.
Visual field testing is a subjective examination. The variability is significant, and the more visual field damage there is, the greater is the variability of the results. Testing the same eye/patient twice in the same day using the same machine does not produce identical results. It should be noted that the differences between devices are mainly due to the differences in the hardware used and the luminosity of the devices. As the available luminosity and luminosity steps of one device approaches the other, the results become more comparable, if both perimeters are
running the same algorithm. The results between different perimeters are similar but not identical. Other studies have found corresponding results.10–15

The most important advantages of VR glasses method are the ease of use and the comfortable patient position; in fact, it has been found that the patients tolerated the test well and fixation losses occurred rarely.16 The patients moved their heads freely. Furthermore, VR glasses method has low cost, and this makes it suitable for use when cost is an important factor.

High correlation coefficient between VR glasses and the Humphrey perimeter shows that the method is reliable at least when compared to the Humphrey perimeter and probably suitable for clinical use.

![Figure 13 Results, eye 10–12.](image1)

![Figure 14 (Continued)](image2)
An additional application for smartphones is Visual Fields Easy designed to use the iPod screen to perform a fast screening test of the visual fields developed at the University of Iowa (Iowa City, IA, USA).

Virtual Eye perimeter is another device operated through a portable Windows computer (laptop or desktop). A simple, single-screen graphical user interface was designed to emulate the performance of standard instruments such as the
Humphrey field analyzer (HFA II), from Carl Zeiss Meditec (Dublin, CA, USA), or Easyfield from Oculus (Wetzlar, Germany). This device requires VR goggles with proprietary interface electronics and a trial lens holder; when the stimulus is detected, the fixation point moves to the position of the detected stimulus.\textsuperscript{14–16}

The Kasha visual field is a system that uses two full color 0.7 inch × 0.7 inch LCD systems. Early trials comparing this
head-mounted perimetry device with the Humphrey field analyzer have found comparable results in terms of field classification. The authors stated that further trials were necessary in order to fully evaluate this device relative to the standard perimetry tools such as the Humphrey or Goldmann field analyzers.4

The advantages of our system are that it does not require proprietary hardware; the screen is large enough, which eliminates the requirement of moving the fixation point, and the patient uses his/her own glasses.

The software is freely available to non-profit institutions by contacting the corresponding author or by sending an email at info@visual-field.com.

Disclosure
The authors report no conflicts of interest in this work.

References