Screening for refractive errors in children: accuracy of the hand held refractor Retinomax to screen for astigmatism

Monique Cordonnier, Michèle Dramaix

Abstract

Aims—To assess the reliability of the hand held automated refractor Retinomax in measuring astigmatism in non-cycloplegic conditions. To assess the accuracy of Retinomax in diagnosing abnormal astigmatism in non-cycloplegic refractive screening of children between 9 and 36 months.

Methods—Among 1205 children undergoing a non-cycloplegic refractive screening with Retinomax, 299 (25%) had repeated non-cycloplegic measurements, 302 (25%) were refracted under cycloglea using the same refractor, and 88 (7%) using retinoscopy or an automated on table refractor. The reproducibility of non-cycloplegic cylinder measurement was assessed by comparing the cylindrical power and axis values in the 299 repeated measurements without cycloglea. The influence of the quick mode on cylinder measurement was analysed by comparing the cylinder and axis value in 93 repeated measurements without cycloglea where normal mode was used in one measurement and quick mode in the other. Predictive values of the refractive screening were calculated for three different thresholds of manifest astigmatism (≥1.5, ≥1.75, and ≥2 D) considering as a true positive case an astigmatism ≥2 D under cycloglea condition (measured by retinoscopy, on table, or hand held refractor).

Results—The 95% limits of agreement between two repeated manifest cylinder measurements with Retinomax attained levels slightly less than plus or minus 1 D. The 95% limits of agreement for the axis were plus or minus 46°. The comparison of non-cycloplegic measurements in the quick and normal mode showed no significant difference and 95% limits of agreement plus or minus 0.75 D. The mean difference between non-cycloplegic and cycloplegic cylinder values measured by Retinomax reached 0.17 D and was statistically significant. Manifest thresholds of ≥1.5 D, ≥1.75 D, ≥2 D cylinder value diagnosed 2 D of astigmatism under cycloglea respectively with 71–84%, 59–80%, 51–54% of sensitivity (right eye-left eye) and 90–92%, 95%, 98% of specificity.

Conclusion—Without cycloglea, Retinomax is able to measure cylinder power with the same reproducibility as cycloplegic retinoscopy. No significant difference was found in the cylinder values obtained with the quick and the normal modes. Therefore, the quick mode of measurement is recommended as it is more feasible in children. No difference, which is significant from a screening point of view, exists between the non-cycloplegic and the cycloplegic cylinder value (<0.25 D).

Retinomax diagnoses abnormal astigmatism (≥2 D) in a non-cycloplegic refractive screening at preschool ages with 51–84% sensitivity rates and 98–90% specificity rates, depending on the chosen threshold of manifest astigmatism. If 2 D of manifest astigmatism is chosen as a positive test, the positive predictive value of the screening reaches 81–84% and the negative predictive value 91–90% (right eye-left eye).

Astigmatism is common in the infant population. The incidence of astigmatism of ≥1 D is around 50% in children under 12 months of age.1 During the first few years of life, the extent and incidence of astigmatism decline, mostly during the second year.1

Amblyopia can be the result of a significant uncorrected astigmatism. Gwiazda et al2 showed that meridional amblyopia can result from astigmatism in early childhood (second half of the first year to the end of the second year). Ingram et al3 found that children having 1.5 D or more of astigmatism at 1 year of age have a higher incidence of amblyopia at 3.5 years. Sjöstrand and Abrahamsson4 found that children with constant and increasing astigmatism are “at risk” of developing amblyopia: oblique astigmatism as well as increasing astigmatism associated with high hypermetropia (≥+3.5 D) presented the relatively higher risks. Atkinson et al5 found a strong relation between meridional amblyopia at 4 years of age and the persistence of astigmatism ≥1 D after 2 years of age. This meridional amblyopia partly contributed to poor performance on the acuity tests at 4 years.

In their study, Friedburg and Kloppel6 showed that early (under 4 years) correction of hypermetropic astigmatism ≥1 D resulted in better development of visual acuity of the dominant eye in strabismic children measured at the age of 8 years or later. The proportions of good vision were even higher when the children were corrected under 2.5 years of age. It is therefore important in refractive screening to diagnose astigmatism adequately.
As we are involved in a non-cycloplegic refractive screening with the hand held refractor Retinomax, we wanted to test the reliability and accuracy of this refractor. We have already studied its ability to screen for high hyperopia\(^1\) and decided to further investigate its usefulness in screening for astigmatism.

**Material**

**RETINOMAX HAND HELD REFRACTOR**

This hand held refractor has been fully described elsewhere.\(^2\) In the normal mode, a fogging system aims to minimise accommodation. The quick mode disables the fogging system and speeds the measurement process which is helpful in very young children: only 10 seconds are needed to start measuring and obtain 16 data points for both eyes. It also allows easy repetition of measurements.

The width of the target (christmas tree on a green grass and blue sky background) is 2 cm, the subject eye is situated 6 cm away from the target. If the subject does not look at the christmas tree but at either side of the target, this will induce a disparity of approximately 10° between the fixation axis and the axis of autorefraction. According to Banks,\(^3\) this could induce a spurious astigmatism of 0.75 to 1.5 D in children.

**Subjects and methods**

Since November 1995, we have organised free visual screening for children between 9 to 36 months at our university hospital situated in Brussels. Specifications concerning the screening have been described elsewhere.\(^4\) Among other tests, this visual screening required a measurement of manifest refraction with the Retinomax autorefractometer. If the child was reluctant and had to be forcibly steadied, care was taken not to push on his eye when opening the lids during the measurements to avoid artefactual astigmatism. In our tests, the success rate of measuring refraction with the hand held refractor in children of this age reached 98.5%.\(^7\) The 1.5% of children who could not be successfully refracted with the Retinomax were refracted by another method (Viva off axis binocular videorefraction, Tomey Inc) and were not included in this study.

Among the 1205 screened children, 20% were screened positive because they had one or more criteria of abnormal manifest refraction (hyperopia >1.5 D, myopia >3 D, astigmatism >2 D, anisometropia >1.5 D) and/or a squint.

These children were referred to an ophthalmologist outside the hospital (for deontological reasons, given the fact that the screening does not show any defect in the eyes of your child.) If the child was screened positive, the parents agreed to cycloplegia being performed. If the child was screened positive because they had one or more criteria of abnormal manifest refraction (hyperopia >1.5 D, myopia >3 D, astigmatism >2 D, anisometropia >1.5 D) and/or a squint. The other 57% were screened negative.

We performed the analysis of agreement between the 302 non-cycloplegic and cycloplegic cylinder values measured by Retinomax.

Whenever non-cycloplegic measurements had been repeated, we chose the most positive representative value for the sphere and the least negative for the cylinder, in order to minimise accommodation.

Eighty eight (7%) of these 1205 screened children had a cycloplegic refraction performed by an on table autorefractor (78 children) or by retinoscopy (10 children): they
belonged to the 20% of children screened positive and the cycloplegic refraction was performed not more than 6 months later by the ophthalmologist to whom the child was referred. As our feedback ratio was only 40% and as some of the feedback forms were unworkable (the refraction was not specified, and as some of the feedback forms were returned), we could not gather these refractive data for all the positive children.

Predictive values of our refractive screening were calculated for three different thresholds of manifest astigmatism (≥1.5, ≥1.75, and ≥2 D) considering as a true positive case an astigmatism ≥2 D under cycloplegia. These predictive values were estimated in the 302 cases having cycloplegic refraction measured by Retinomax and in the 88 cases having cycloplegic refraction performed by retinoscopy or on table refractor.

Results

Figure 1 shows the age distribution in months of the 1205 screened children, of the 302 children refracted under cycloplegia by the hand held refractor, and of the 88 children refracted under cycloplegia by retinoscopy or on table refractor.

Table 1 shows the mean difference of non-cycloplegic minus cycloplegic value. d = mean difference of non-cycloplegic minus cycloplegic value. S = significant.

Discussion

Twenty per cent of children are screened positive for refractive anomalies and/or squint. This figure is approximately twice that found in the literature. The reason is that our screening is expected by their parents or paediatrician. Among others, we selected many children with a family history of strabismus or amblyopia and also children having a visual anomaly suspected by their parents or paediatrician.
The the relative proportion of positive (43%) and negative (57%) results among the 302 children controlled under cycloplegia is different from the one of the screening (20% versus 80%). In our opinion, the reason is that the parents were more inclined to accept the cycloplegia for their child if it was to confirm an anomaly.

The age distribution in the 1205 screened children and in the 302 children also refracted under cycloplegia by the hand held refractor is similar, showing a good breakdown of the population by age groups. Less than 15% of the children did not have the intended screening age and we deemed it unnecessary to discard them for the statistical analysis.

The distribution of age of the 88 children screened positive and refracted under cycloplegia by the ophthalmologist to whom the child was referred shows a definite trend towards higher ages. As we decided in this study to discard the refractive feedback that took place more than 6 months after the screening, the delay between the screening and the consultation cannot explain this difference. We think that the main reason is the following: the ophthalmologist was reluctant to send us a dated card the refractive feedback that took place more than 6 months after the screening. As we decided in this study to discard the feedback, we received concerned children of 2 years and over who were refracted by an on table refractor (78 children) rather than by retinoscopy (10 children).

There is no significant difference between non-cycloplegic repeated measurements of cylinder and axis with Retinomax. The 95% limits of agreement between two repeated cylinder measurements reach levels slightly less than ±1 D. This is comparable with the agreement between repeated measurements along one meridian by non-cycloplegic autorefraction (−0.72 to 0.71 D) and by cycloplegic retinoscopy (−0.87 to 1.02 D).13

The 95% limits of agreement for the axis are ±46°, which conforms with the literature: Bermann et al.13 found it to be ±33° in repeated non-cycloplegic retinoscopy and ±65° in repeated manifest autorefraction (Canon R-1). McBrien and Millodot14 found d ±20° with the same Canon autorefractor in manifest conditions. Wood et al.15 found d ±40° with the Nidek autorefractor in manifest conditions.

The comparison of non-cycloplegic measurements in the quick and normal mode gives similar results: no significant difference and 95% limits of agreement around ±0.75 D. This confirms our previous study17 concerning the amount of sphere in hyperopic children: no difference is seen between the quick and the normal modes in minimising accommodation. We therefore suggest choosing the quick mode for screening children of this age, since measurements in this mode are much easier to obtain and nearly always successful in our test.

The mean difference between non-cycloplegic and cycloplegic cylinder values measured by Retinomax reaches 0.17 D and is statistically significant. This means that the value of the cycloplegic cylinder is generally higher than the non-cycloplegic one (all our cylinders are negative). As already suggested by Rubin and Harris,16 this could imply that the accommodative change in a human eye is not entirely spherical in nature but may include an additional cylindrical component. This difference of 0.17 D is, however, not clinically important from a screening point of view.

Several concerned ophthalmologists do not prescribe glasses in children of 12 months of age if an astigmatism with coincident axes below 2 D.15 We chose therefore a value of 2 D or more of astigmatism under cycloplegic condition as true positive case to establish the sensitivity, specificity, and predictive values of the screening.

In our opinion, Table 4 gives a more accurate idea of the overall sensitivity and specificity rates of our screening than Table 5: the sample is larger (302 children versus 88), the age distribution is comparable with the one of the screened population (Fig 1), and the refraction is better controlled (all measurements were made the same day, by the same operators, with the same refractor, and using the same cycloplegic protocol). The results of both tables are however rather similar. The threshold of manifest astigmatism ≥ 2 D gives the best figures regarding specificity and positive predictive value. The thresholds of ≥ 1.75 D and ≥ 1.5 D give a better combination of sensitivity and specificity.

Although a high threshold of abnormal astigmatism diminishes the sensitivity of the screening, it improves its specificity. To validate our visual screening and as astigmatism is not a severely disabling eye condition, we preferred a highly specific test to avoid overreferrals and to have a good positive predictive value.18

### Table 4: Performances of the non-cycloplegic screening for three different thresholds of manifest astigmatism (true positive case if cycloplegic cylinder value ≥ 2 D)

<table>
<thead>
<tr>
<th>Manifest cylinder</th>
<th>Positive test n (%)</th>
<th>Negative test n (%)</th>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
<th>PPV (%)</th>
<th>NPV (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>≥ 2 D Right eye</td>
<td>32 (11)</td>
<td>270 (89)</td>
<td>51</td>
<td>98</td>
<td>81</td>
<td>91</td>
</tr>
<tr>
<td>≥ 1.75 D Right eye</td>
<td>42 (14)</td>
<td>260 (86)</td>
<td>59</td>
<td>95</td>
<td>71</td>
<td>92</td>
</tr>
<tr>
<td>≥ 1.5 D Right eye</td>
<td>60 (20)</td>
<td>242 (80)</td>
<td>80</td>
<td>95</td>
<td>78</td>
<td>95</td>
</tr>
<tr>
<td>≥ 2 D Left eye</td>
<td>38 (13)</td>
<td>264 (87)</td>
<td>54</td>
<td>98</td>
<td>84</td>
<td>90</td>
</tr>
<tr>
<td>≥ 1.75 D Left eye</td>
<td>41 (14)</td>
<td>260 (86)</td>
<td>59</td>
<td>95</td>
<td>71</td>
<td>92</td>
</tr>
<tr>
<td>≥ 1.5 D Left eye</td>
<td>60 (20)</td>
<td>242 (80)</td>
<td>80</td>
<td>95</td>
<td>78</td>
<td>95</td>
</tr>
</tbody>
</table>

**PPV** = positive predictive value.
**NPV** = negative predictive value.

### Table 5: Performances of the non-cycloplegic screening for three different thresholds of manifest astigmatism (true positive case if cycloplegic cylinder value by retinoscopy or on table refractor ≥ 2 D)

<table>
<thead>
<tr>
<th>Manifest cylinder</th>
<th>Positive test n (%)</th>
<th>Negative test n (%)</th>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
<th>PPV (%)</th>
<th>NPV (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>≥ 2 D Right eye</td>
<td>20 (23)</td>
<td>68 (77)</td>
<td>52</td>
<td>95</td>
<td>85</td>
<td>76</td>
</tr>
<tr>
<td>≥ 1.75 D Right eye</td>
<td>25 (29)</td>
<td>62 (71)</td>
<td>65</td>
<td>90</td>
<td>76</td>
<td>87</td>
</tr>
<tr>
<td>≥ 1.5 D Right eye</td>
<td>23 (26)</td>
<td>65 (74)</td>
<td>58</td>
<td>93</td>
<td>83</td>
<td>78</td>
</tr>
<tr>
<td>≥ 2 D Left eye</td>
<td>35 (40)</td>
<td>52 (60)</td>
<td>83</td>
<td>81</td>
<td>69</td>
<td>90</td>
</tr>
<tr>
<td>≥ 1.75 D Left eye</td>
<td>31 (35)</td>
<td>57 (65)</td>
<td>82</td>
<td>93</td>
<td>87</td>
<td>89</td>
</tr>
<tr>
<td>≥ 1.5 D Left eye</td>
<td>40 (46)</td>
<td>47 (54)</td>
<td>89</td>
<td>76</td>
<td>65</td>
<td>94</td>
</tr>
</tbody>
</table>

**PPV** = positive predictive value.
**NPV** = negative predictive value.
Therefore, we decided to keep the threshold of 2 D for the ongoing screening.

To conclude, manifest refractive screening with Retinomax diagnoses abnormal astigmatism (>2 D) with 51% to 84% sensitivity rates and 98% to 90% specificity rates, depending on the chosen threshold of manifest astigmatism. If 2 D of manifest astigmatism is considered for positive test, the positive predictive value of the screening reaches 81–84% and the negative predictive value 91–90% (right eye-left eye). As the quick mode of measurement is more feasible in children and as we did not establish any significant difference between the quick mode and the normal mode in measuring manifest refraction, we also suggest choosing this mode of measurement for screening very young children.

This study has been funded by “Les Amis des Aveugles” ASBL, rue de la barrière, 37–39, 7011 Ghlin. The authors would like to thank Moktar Essarhdaoui, the main operator for the screening.

Screening for refractive errors in children: accuracy of the hand held refractor Retinomax to screen for astigmatism
Monique Cordonnier and Michèle Dramaix

Br J Ophthalmol 1999 83: 157-161
doi: 10.1136/bjo.83.2.157

Updated information and services can be found at:
http://bjo.bmj.com/content/83/2/157

These include:

References
This article cites 18 articles, 3 of which you can access for free at:
http://bjo.bmj.com/content/83/2/157#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections
Optic nerve (713)
Optics and refraction (508)
Epidemiology (1077)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/