Non-ophthalmologist screening for retinopathy of prematurity

Richard A Saunders, Margaret L Donahue, Jerry E Berland, Eric L Roberts, Billy Von Powers, Philip F Rust

Abstract

**Aim**—To determine if a non-ophthalmologist can accurately screen for retinopathy of prematurity (ROP) by evaluating the posterior pole blood vessels of the retina. ROP is a common ocular disorder of premature infants and may require multiple screening examinations by an ophthalmologist to allow for timely intervention. Since there is a strong correlation between posterior pole vascular abnormalities and vision threatening ROP, screening examinations performed by non-ophthalmologist may yield useful clinical information in high risk infants.

**Methods**—Infants born at the Medical University of South Carolina who met screening criteria (n = 142) were examined by a single non-ophthalmologist using a direct ophthalmoscope to evaluate the posterior pole blood vessels for abnormalities of the venules and/or arterioles. To determine the accuracy of the non-ophthalmologist’s clinical observations, infants were also examined by an ophthalmologist, using an indirect ophthalmoscope, who graded the posterior pole vessels as normal, dilated venules, or dilated and tortuous venules and arterioles (including “plus disease”).

**Results**—There was significant correlation (p < 0.001) between the non-ophthalmologist’s and ophthalmologist’s diagnoses of posterior pole vascular abnormalities. 47 infants had normal posterior pole blood vessels by the non-ophthalmologist examination. Of these, 31 (66%) were considered to have normal vessels and 16 (34%) to have dilated venules by the ophthalmologist. The non-ophthalmologist correctly identified abnormal posterior pole vessels in all 21 infants diagnosed with abnormal arterioles and venules by the ophthalmologist. No infants with clinically important ROP (“prethreshold” or worse) would have failed detection by this screening method.

**Conclusion**—Using a direct ophthalmoscope, a non-ophthalmologist can screen premature infants at risk for ROP by evaluating the posterior pole blood vessels of the retina. While not necessarily recommended for routine clinical practice, this technique may nevertheless be of value to those situations where ophthalmological consultation is unavailable or difficult to obtain.

(Br J Ophthalmol 2000;84:130–134)
little is known about the potential effectiveness of alternative screening programmes.

A recent study in premature infants has shown a significant correlation between the appearance of the posterior pole blood vessels and the severity of peripheral retinal disease in ROP.14 Infants with normal appearing posterior pole vessels had mild or no ROP, lessening the need for concurrent indirect ophthalmoscopic examination of the peripheral retina. Examination of the posterior pole vessels can be accomplished using a standard hand held (direct) ophthalmoscope and, more importantly, potentially be performed by non-ophthalmologists with reasonable accuracy after only minimal training.15 Our current study was undertaken as part of the training of a staff neonatologist (MLD) to evaluate the status of the posterior pole blood vessels in premature infant eyes. If deemed sufficiently reliable, preliminary screening by non-ophthalmologists could be used to detect infants with evidence of progressive ROP who might then receive expedited referral to a specialist for evaluation and appropriate intervention.

Subjects and methods
Premature infants admitted to the intensive care nursery at the Medical University of South Carolina Children’s Hospital between October 1993 and December 1994 (excluding 15 October to 7 December 1993) with birth weights less than 1600 g, were enrolled in the study. Approval of the protocol was obtained by the university’s institutional review board for human research. Routine screening examinations for ROP were performed before discharge or by 33 weeks post-conceptional age. Follow up examinations were performed as necessary, depending on clinical findings. Before each examination, the pupils were dilated with either Cyclomydril or sequential instillation of cyclopentolate 0.5% and phenylephrine HCl 2.5%. Using a Cook-style paediatric eyelid speculum,16 funduscopic examination was performed 30–60 minutes later at the bedside.

A non-ophthalmologist (MLD) examined both retinas of each infant using a halogen bulb direct ophthalmoscope and recorded whether the blood vessels were normal or abnormal, based on reference photographs (Figs 1–4).

Other than explanation of the fundus photographs and viewing several examples of vascular abnormalities through the teaching mirror of the indirect ophthalmoscope, the non-ophthalmologist examiner received no in-depth training in ophthalmoscopy or the evalu-
ation of posterior pole blood vessels before beginning this study. A vessel pair was considered abnormal if there was venous dilatation with or without accompanying dilatation or tortuosity of the retinal arterioles. Venules were considered dilated if they were greater than twice the calibre of normal appearing arterioles in the same eye, whereas the normal diameter ratio of venules to arterioles is approximately 3:2. Twelve eyes could not be evaluated using direct ophthalmoscopy because of vitreous haze, poorly dilated pupils, or inability to obtain adequate focus on the posterior pole structures.

After the non-ophthalmologist’s findings had been recorded, a paediatric ophthalmologist (RAS) or paediatric ophthalmology fellow (JEB or ELR) examined both eyes using the indirect ophthalmoscope and graded the posterior pole blood vessels as 1, normal, 2, dilated venules, or 3, dilated and tortuous arterioles and venules using the same photographic guidelines. Dilated and tortuous vessels did not necessarily imply that “plus disease” was present, but plus disease is included as a subset in this third group. Finally, an examination of the peripheral retina was performed using scleral depression and the findings recorded using the International Classification of ROP. The ophthalmologist examiner was not aware of the findings of the non-ophthalmologist examiner until each posterior pole examination was complete and the data recorded for both eyes. Infants with previously identified retino-vascular abnormalities who were familiar to the non-ophthalmologist were excluded from the data analysis.

Results

Our results are summarised in Tables 1–4. A total of 142 infants were evaluated by both an ophthalmologist and non-ophthalmologist examiner. Twelve infants had incomplete examinations by the non-ophthalmologist because of inability to assess the posterior pole with the direct ophthalmoscope. These 12 infants are included in the “abnormal blood vessel” category in Tables 1 and 2. On peripheral retinal examination, eight had immature vessels without ROP, three had zone II, stage 2 ROP, and one had zone II, stage 3 ROP.

Table 1 shows the correlation between the non-ophthalmologist’s interpretation of the posterior pole vessels and the ophthalmologist’s findings for the same infants. Of the 142 infants examined by the non-ophthalmologist, 95 (67%) were felt to have retinovascular abnormalities.
Table 4 Posterior pole vascular findings. Three dimensional frequency analysis of interreader interaction, non-ophthalmologist/retinopathy of prematurity interaction, and ophthalmologist/retinopathy of prematurity interaction

<table>
<thead>
<tr>
<th>Non-ophthalmologist’s examination</th>
<th>Peripheral retinal findings</th>
<th>Normal</th>
<th>Abnormal</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal blood vessels</td>
<td>Mature</td>
<td>5</td>
<td>7</td>
<td>12</td>
</tr>
<tr>
<td></td>
<td>Immature or less than prethreshold</td>
<td>26</td>
<td>9</td>
<td>35</td>
</tr>
<tr>
<td></td>
<td>Prethreshold or threshold</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>31</td>
<td>16</td>
<td>47</td>
</tr>
<tr>
<td>Abnormal blood vessels</td>
<td>Mature</td>
<td>0</td>
<td>11</td>
<td>11</td>
</tr>
<tr>
<td></td>
<td>Immature or less than prethreshold</td>
<td>16</td>
<td>51</td>
<td>67</td>
</tr>
<tr>
<td></td>
<td>Prethreshold or threshold</td>
<td>0</td>
<td>17</td>
<td>17</td>
</tr>
<tr>
<td></td>
<td>Total</td>
<td>16</td>
<td>79</td>
<td>95</td>
</tr>
</tbody>
</table>

Discussion

Our results suggest that a non-ophthalmologist can be trained to screen premature infants for retinovascular abnormalities associated with severe ROP. While the standard of practice in the United States currently calls for ophthalmological screening for high risk premature infants, staffing considerations may make this impractical in other parts of the world. A screening protocol relying on non-ophthalmologists could potentially be more comprehensive and cost effective by obviating the need for specialist consultation for every infant, yet allowing appropriate referral for the majority of high risk cases potentially requiring surgical intervention. In our experience, preventable blindness from ROP has almost always been associated with failure to perform appropriate screening or follow up examinations, not failure to diagnose correctly. Therefore, even in nursery environments where ophthalmic consultation is more readily available, examination of posterior pole retinal blood vessels by non-ophthalmologists may still have diagnostic value.

Although a non-ophthalmologist screening protocol may be a useful clinical tool, there are several points that must be addressed. Firstly, consistent agreement on the diagnosis of dilated venules, even among ophthalmologists, was difficult to achieve. The comparison of the diameter of venules to arterioles was often borderline at a 2:1 ratio and might vary among vessel pairs within the same eye. In an emmetropic eye, image magnification with the direct ophthalmoscope is approximately five times greater than the indirect ophthalmoscope using a 20 dioptre condensing lens, and seven times greater than a 30 dioptre condensing lens. A certain amount of disagreement among examiners using different ophthalmoscopes would therefore be expected. Furthermore, during examination with the direct ophthalmoscope, often only one vessel pair can be adequately visualised. It is possible that the specific pair examined may or may not be abnormal, although vessel pairs that were not examined may have dilated venules or even arteriolar tortuosity. This problem can be overcome by examining more than one vessel pair in each fundus. Examiner persistence and perhaps prolonged or sequential examinations may sometimes be required.

While our high sensitivity for detecting posterior pole vascular abnormalities indicates that a non-ophthalmologist would not be likely to miss clinically important ROP on routine screening, specificity was poor. In this study, “abnormal” posterior pole blood vessels were identified in two thirds of infants undergoing screening examination by the non-ophthalmologist examiner using a direct ophthalmoscope (Table 1). This would lead to many unnecessary referrals of low risk infants. Our arbitrary definition of dilated venules (greater than 2:1 ratio of the diameter of venules to arterioles) may therefore need to be revised to improve specificity in the correlating of posterior pole findings with important peripheral retinal disease. More precise grading of posterior pole vascular abnormalities, as we have proposed using the indirect ophthalmoscope, might also be possible.

Secondly, dilatation of the pupils and use of an eyelid speculum is generally required regardless of whether a non-ophthalmologist or ophthalmologist performs the screening examination. Dilating eye drops need to be available in appropriate concentrations for premature infant to minimise potential medication side effect. The non-ophthalmologist will require appropriate training to place an eyelid speculum in the eye without scratching the cornea or damaging adnexal structures.

Thirdly, training of our non-ophthalmologist examiner using reference photographs and funduscopic examination through the teaching mirror of an indirect ophthalmoscope probably improved our results. Other non-ophthalmologists interested in using this screening protocol would presumably require
similar instruction to achieve an acceptable level of accuracy. Currently, there are no known programmes teaching these techniques to non-ophthalmologists. However, the experience needed to become competent in posterior pole vessel evaluation with the supervision of an ophthalmologist may be as few as 20–25 examinations, assuming that these would include infants with varying degrees of abnormal posterior pole vascular findings.

It must also be stressed that the finding of normal posterior pole vessels at one point in time does not rule out the potential for developing blinding ROP later on. This screening protocol requires sequential examinations, documenting repeatedly normal posterior pole vessels or, alternatively, referral to an ophthalmologist for further evaluation. Previous studies on the natural progression of ROP have documented that the majority of infant retinas show mature vasculature by 38 weeks’ post-conceptional age. It would therefore seem prudent that each high risk infant receive, where available, at least one additional retinal examination by an ophthalmologist at approximately 38 weeks’ post-conceptional age to document retinal vascularisation into zone III. However, in countries where ophthalmological consultation is not routinely available and the incidence of cicatricial ROP is high, examinations by non-ophthalmologists might represent the only screening a high risk infant receives before hospital discharge.

This study was supported in part by an unrestricted grant to the Storm Eye Institute from Research to Prevent Blindness, Inc., New York, USA.


Non-ophthalmologist screening for retinopathy of prematurity

Richard A Saunders, Margaret L Donahue, Jerry E Berland, Eric L Roberts, Billy Von Powers and Philip F Rust

Br J Ophthalmol 2000 84: 130-134
doi: 10.1136/bjo.84.2.130

Updated information and services can be found at:
http://bjo.bmj.com/content/84/2/130

These include:

References
This article cites 17 articles, 0 of which you can access for free at:
http://bjo.bmj.com/content/84/2/130#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

Epidemiology (1068)
Paediatrics (358)
Retina (1608)

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/