Computer-Assisted Measurement of Retinal Vascular Width and Tortuosity in Retinopathy of Prematurity

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**Objectives**: To validate the accuracy of ROPtool software in measuring retinal vascular width and tortuosity in a large image set compared with expert diagnoses.

**Methods**: Tortuosity and dilation indexes generated by ROPtool were compared with 3 expert consensus grades of normal, pre-plus, or plus disease for 368 quadrants in 92 RetCam (Clarity Medical Systems, Pleasanton, California) fundus images. Sensitivity and specificity of ROP-tool software in diagnosing tortuosity and dilation sufficient for plus and pre-plus disease were calculated. These measures were compared with individual accuracies of 3 experienced pediatric ophthalmologists.

**Results**: The mean tortuosity indexes for expert-diagnosed categories of normal, pre-plus, and plus disease were 7.04, 18.73, and 34.62, respectively (P < .001), and the mean dilation indexes were 9.63, 12.05, and 13.61, respectively (P < .001). When optimal tortuosity and dilation index thresholds (from receiver operating characteristic curves) were applied, resultant sensitivity and specificity were 0.913 and 0.863, respectively, for plus tortuosity and 0.782 and 0.840, respectively, for plus dilation. These values were comparable to the performance of examiners judged against the same expert panel.

**Conclusion**: ROPtool version 2.1.5 accurately measures tortuosity and dilation of posterior pole blood vessels in RetCam images, corresponding well with expert diagnostic categories of normal, pre-plus, and plus disease and performing comparably to experienced examiners.

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this newly developed measure of vessel width performed well. Our primary objective in the present study was to validate the accuracy of ROPtool software (version 2.1.5) in measuring retinal vascular width and tortuosity using a large image set. This validation is an important step toward using ROPtool clinically at the bedside or as part of a telemedicine program. We also sought to assess (for the first time) diagnostic accuracy of ROPtool when tortuosity and dilation were considered together, as is done during clinical examination.

VALIDATION STUDY OF ROPtool ACCURACY IN ASSESSING RETINAL VASCULAR TORTUOSITY AND DILATION

ROPtool Analysis

A convenience sample of 92 RetCam fundus images was used for this larger validation study. These were a subset of images described in previous publications, none of which were in the pilot study image set. All images were analyzed using ROPtool operated by one of us without knowledge of prior grading (A.E.K.), who selected 2 major vessels in each of 4 quadrants wherever possible, with a goal of identifying the major arteriole and major venule in each quadrant. Although the program has the capacity to record operator judgments of vessel identity (arteriole vs venule), this function was not used herein. ROPtool analysis required approximately 1 to 3 minutes per image and never exceeded 6 minutes. ROPtool returned TIs and DIs for each vessel and each quadrant, and the “quadrant index” equaled the largest vessel index in that quadrant. Figure 1 shows the interface of ROPtool version 2.1.5.

Accuracy of ROPtool

Using the cutoff points generated in the pilot study, ROPtool scores were then converted into designations of plus, pre-plus, or neither for tortuosity alone and for dilation alone. Next, sensitivity and specificity of ROPtool in diagnosing tortuosity and dilation sufficient for plus and pre-plus disease categories were calculated using the consensus of 3 expert examiners (“experts”) as the reference standard. All 3 were certified investigators in the Early Treatment for Retinopathy of Prematurity
For comparison, individual accuracies of 3 additional experienced pediatric ophthalmologists (“examiners”) relative to those of the experts were calculated in the same way.

Details of Expert and Examiner Grading

The experts and examiners assessed whether each of 4 quadrants in an image (cropped to a 28-diopter indirect ophthalmoscopy view) contained retinal vascular dilation, tortuosity, or both sufficient to diagnose plus or pre-plus disease. Grade 0 corresponded to normal vasculature, 1 to pre-plus disease, and 2 to plus disease. In preparation for this process, all 6 graders were reminded of the definition of plus disease and were sent a copy of the standard photograph used in the Cryotherapy for ROP study, as well as examples of posterior pole images from the Original Article titled, “International Classification of Retinopathy of Prematurity Revisited.” The grades of the 3 experts were averaged and rounded to the nearest whole number to establish the reference standard (“truth”) for each quadrant.

ROC Curves

Using data from all 368 quadrants of 92 RetCam fundus images, ROC curves were constructed for plus and pre-plus TI and DI, plotting sensitivity vs 1 minus specificity of each potential threshold index compared with quadrant-level expert truth. Areas under the curves were then calculated. From these curves, candidate plus and pre-plus thresholds were identified for future recalibration of ROPtool.

STATISTICAL ANALYSIS

All data were collected and analyzed using commercially available software (Microsoft Office Excel 2003; Microsoft Corporation, Redmond, Washington). ROC curves were generated and analyzed using Excel 2003 and MatLab 7.7 (2008b; MathWorks, Natick, Massachusetts). For determining statistical significance of TI and DI “trends” across the 3 expert diagnostic groups, the generalized estimating equation approach was used to account for correlation within quadrants of the same eye. Because the increase over the disease groups for each outcome measure was not linear, the tests for trends were performed as tests for the difference from the normal group for each disease group (pre-plus and plus) simultaneously.

RESULTS

TREND FOR INCREASING ROPtool VALUES

Figure 2 shows trends for increasing ROPtool values for tortuosity and dilation (P < .001 for both) across expert-diagnosed categories of normal, pre-plus, and plus disease. The mean TIs for the 3 categories were 7.04, 18.73, and 34.62, respectively, and the median TIs were 6.27, 18.38, and 30.39, respectively. The mean DIs were 9.63, 12.05, and 13.61, respectively, and the median DIs were 9.81, 12.58, and 13.16, respectively.

ACCURACY OF ROPtool AND INDIVIDUAL EXAMINERS RELATIVE TO EXPERT CONSENSUS

The Table shows sensitivity, specificity, and overall concordance for diagnosing tortuosity and dilation sufficient for plus disease. The sensitivities of the pilot study thresholds were excellent (1.000 and 0.964 for tortuosity and dilation, respectively), but the specificities of 0.658 and 0.446, respectively, indicated many false-positive results or overcalls. Overall concordance with experts was 0.701 for tortuosity and 0.541 for dilation using these a priori pilot study thresholds. When new optimal cutoff points were chosen from the present data set, specificity and overall concordance of ROPtool improved to 0.863 and 0.870, respectively, for plus tortuosity, and to 0.840 and 0.832, respectively for plus dilation.

Figure 3 shows ROC curves for tortuosity at the plus (A) and pre-plus (B) levels and for dilation at the plus (C) and pre-plus (D) levels. Areas under the curve are robust in all 4 cases.

INTEREXPERT AGREEMENT

In the pilot study, the ophthalmologists whose consensus comprised the reference standard agreed on a designation of plus, pre-plus, or normal dilation without adjudication for 127 of 154 (82%) vessels; agreement on the degree of tortuosity was previously reported to be 82%.
The 3 experts determining truth in the larger validation study were unanimous in their 3-level tortuosity grades for 242 of 368 quadrants (66%) and in their 3-level dilation grades for 187 of 368 quadrants (51%). In 13 (4%) and 18 (5%) quadrants, their opinions of tortuosity and dilation, respectively, diverged completely, with one giving a grade of “plus,” another “pre-plus,” and the third “normal.”

**COMMENT**

ROPtool version 2.1.5 performed well in the present study, returning accurate measurements of tortuosity and dilation for high-quality RetCam fundus images. ROPtool analysis is rapid, typically requiring less than 3 minutes per image. The robust areas under the ROC curves indicate good correspondence of ROPtool with expert diagnoses of plus- and pre-plus-level retinal vessel tortuosity and dilation. We have shown herein that, given optimized plus and pre-plus thresholds, the diagnostic capabilities of ROPtool are on par with those of experienced examiners.

By contrast, ROPtool did not perform as well when the program was calibrated with thresholds determined

<table>
<thead>
<tr>
<th>Variable (^a)</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>Concordance</th>
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<tbody>
<tr>
<td>Plus Disease Tortuosity</td>
<td>Pilot study</td>
<td>1.00</td>
<td>0.66</td>
</tr>
<tr>
<td>ROPtool threshold</td>
<td>New(^b)</td>
<td>0.91</td>
<td>0.86</td>
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<tr>
<td>Expert</td>
<td>1</td>
<td>0.94</td>
<td>0.86</td>
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<tr>
<td>2</td>
<td>0.80</td>
<td>0.96</td>
<td>0.94</td>
</tr>
<tr>
<td>3</td>
<td>0.63</td>
<td>0.99</td>
<td>0.94</td>
</tr>
</tbody>
</table>

| Plus Disease Dilation | Pilot study | 0.96 | 0.45 | 0.54 |
| ROPtool threshold | New\(^b\) | 0.78 | 0.84 | 0.83 |
| Expert | 1 | 0.94 | 0.86 | 0.87 |
| 2 | 0.89 | 0.88 | 0.88 |
| 3 | 0.49 | 0.97 | 0.90 |

\(^a\) Pilot study indicates ROPtool performance when calibrated with the numeric plus and pre-plus disease thresholds determined a priori from pilot study data.

\(^b\) New indicates ROPtool performance when calibrated with optimal thresholds selected from the data set generated in the present study.

**Figure 3.** Receiver operating characteristic curves assessing the diagnostic accuracy of ROPtool. Tortuosity index (A and B) and dilation index (C and D) agreement with expert diagnoses is shown for plus disease (A and C) and for pre-plus disease or worse (B and D). AUC indicates area under the curve. Arrows show optimal thresholds to maximize sensitivity while minimally compromising specificity (tortuosity index of 17.00 and dilation index of 10.50 for plus disease and tortuosity index of 10.50 and dilation index of 10.50 for pre-plus disease).
a priori from pilot study data. This disparity is likely due to disagreement between different groups of experts used for the pilot study and the present study. Chiang et al previously reported that a panel of 22 experts agreed on 3-level categorization of plus disease, similar to that in the present study, only 12% of the time. In their study the mean weighted $\kappa$ statistics for individual experts compared with the overall panel indicated only fair ($\kappa$ range, 0.21-0.40) to moderate ($\kappa$ range, 0.41-0.60) agreement. Therefore, it is not surprising that our reference standard shifted when determined by a panel of 3 experts instead of the 2 of us used for the pilot study. Among our examiners, we also observed a wide range of sensitivities for detecting abnormality, 0.630 to 0.935 for plus-level tortuosity and 0.491 to 0.945 for plus-level dilation. Although human variability represents a fundamental limitation in using expert consensus as diagnostic truth, human opinion remains the standard approach in clinical care for ROP. Given the capacity of ROPtool for consistent measurement, this computer program can be calibrated to match any desired thresholds for plus and pre-plus disease, even if these vary among different groups of clinical experts.

While established thresholds of disease severity are necessary for diagnosis and intervention, the spectrum of indexes in ROPtool may eventually allow these to be tailored to each clinical circumstance. For instance, in a telemedicine screening program, sensitivity might be the most important factor. In this circumstance, it is critical to avoid missing disease that threatens a child’s vision, even at the expense of increased false-positive results. The disadvantage of false-positive results is that they require the ophthalmologist to examine an infant in the nursery who does not need treatment, despite the use of photoscreening. We believe that these additional examinations are a reasonable price to pay for avoiding wholly unacceptable false-negative results. Thresholds for referral could be tailored to accommodate a primary provider’s baseline diagnostic capabilities simply by calibrating ROPtool. If desired, excellent sensitivity for eyes with plus disease requiring treatment or with pre-plus disease requiring close ophthalmologic supervision could be attained. On the other hand, if ROPtool was used to give a bedside “second opinion,” balancing sensitivity and specificity would be more important. Furthermore, new ROP therapies on the horizon such as antivascular endothelial growth factor pharmacotherapy might be used at a different level of disease severity than laser treatment. Optimal points for intervention could be located along a spectrum of cutoff points to produce a multipurpose diagnostic tool. In the future, we might even move away from the notion of 1 or 2 cut points in ROP management altogether, considering instead a progression of anatomical changes in retinal vessels that can be quantified.

Finally, results of this study give new insight into the human diagnostic process. For tortuosity and dilation, there were trends for increasing index values from one diagnostic category to the next. Of note, tortuosity measurements covered a much larger range than dilation measurements. This finding indicates more marked contrast between normal vessel curvature and pathologic tortuosity, which a human can appreciate visually. Although the trend for increasing tortuosity from normal to plus disease is clearly demonstrated (Figure 2A), we see only minimal difference in median vessel width from pre-plus to plus disease (Figure 2B). One possibility is that ROPtool cannot distinguish pathophysiologic changes in vessel dilation, but the marked difference between normal and greater-than-normal (plus or pre-plus) dilation indexes suggests otherwise. A second possibility is that the range of pathophysiologic change in vessel width from normal to pre-plus or plus level is narrower than that for vessel tortuosity in ROP. A third possibility is that clinicians appreciate increasing venous dilation but require a sufficiently tortuous arteriole to consider a quadrant plus disease, no matter how dilated nearby vessels appear. Therefore, expert categories of plus dilation and pre-plus dilation have considerable overlap of ROPtool values. Clinicians typically consider tortuosity and dilation together when judging plus disease, and further study is necessary to identify the best method for mathematically combining tortuosity and dilation measurements for overall diagnoses that reflect expert truth.

This study should be viewed in light of some limitations. First, our convenience sample size was limited in number and was composed entirely of RetCam fundus images, which are not available to all care providers. Second, experts graded images as plus, pre-plus, or neither; using a scale with finer gradations (eg, 0-9) may have highlighted subtler diagnostic distinctions. Third, the effects of image quality on ROPtool measurement remain to be clarified; when we isolated images on which ROPtool performed less well, there were no clear similarities between them with regard to image quality. Fourth, because our primary objective was evaluation of the accuracy of ROPtool, we did not compare consensus of experts viewing images with results of indirect ophthalmoscopy; such an analysis would someday have a role in justifying the use of telemedicine screening.

In conclusion, ROPtool version 2.1.5 accurately measures tortuosity and dilation of posterior pole blood vessels in RetCam images, corresponding well with expert diagnostic categories of normal, pre-plus, and plus disease. In addition, we observed trends in tortuosity and dilation indexes from one expert category to the next that may provide insight into human (not just computer-assisted) ROP diagnosis. Going forward, we plan to investigate the best method to mathematically combine vessel width and tortuosity, which will greatly facilitate bedside or telemedicine diagnosis of plus disease.

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REFERENCES