Predictive Values of Optical Coherence Tomography (OCT) Parameters in Assessment of Glaucoma progression

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SUMMARY
Goal: the purpose of the current study was to estimate the predictive values of optical coherence tomography parameters in early, developed perimetric and terminal glaucoma. Methods: 180 eyes of 120 consecutive patients were evaluated in this retrospective cross sectional pilot study. Copernicus Spectral –domain optical coherence tomography with resolution of 3 mm obtained through the optic nerve head were included. All examined eyes were divided to four groups (healthy,early, developed perimetric glaucoma and terminal glaucoma). The values of the thickness of the retinal nerve fibre layer, the size of the disk, the volume of the cup, the E/D parameter and the size of the RIM were compared in four study groups. Results: The sensitivity of RNFL was 90,0%, specificity 82,0 %, positive predictability 83,3 % and negative predictability was 89,1 %. The total accuracy was 86,0 % and area under curve (AUC) was 0,878 for RNFL indeks comparing early to developed glaucoma. The sensitivity for CUP was 78%, the specificity was 80,8 %, the positive predictability 81,2% and the negative predictability was 77,5 %. The total accuracy was 77,5 % and area under curve (AUC) was 0,792. The sensitivity for RNFL was 88,0 %, the specificity was 66,7 %, the positive predictability was 81,5% and the negative predictability was 76,9%. The total accuracy was 80,0% and the area under curve (AUC) for RNFL comparing developed to terminal glaucoma was 0,815. The increasing 0,1 unit RNFL decreases the risk of developing glaucoma from early to another developed stage of glaucoma for 6,95%. The increasing of E/D for only one unit increases the risk to develop another stage of glaucoma for 18,75 times. The increasing for only one unit of CUP increases the risk to develop terminal glaucoma for 8,47 times and increasing for 0,1 unit of the value of RIM decreases the risk developing terminal glaucoma for 9,27%. The increasing for 0,01 unit of the E/D index increases the risk for terminal glaucoma for 23,23 times. The increasing for one unit of RNFL decreasing the risk developing terminal glaucoma for 5,7%

Key words: glaucoma, progression, optical coherence tomography, optic disc morphology.

1. INTRODUCTION
The progressive glaucomatous optic disc atrophy is characterized by the optic disc changes, including enlargement of the optic disc cup and the loss of the neuroretinal rim. Morphologic changes may precede loss of function as the tests of the visual field (1, 2). Optical coherence tomography (OCT) was introduced approximately 20 years ago and has achieved an important role in present clinical diagnoses (3, 4).

With aim to diagnosed glaucoma it is fundamental to identify glaucoma progression in order to intensify the treatment before nerve fiber loss proceeds. In terminal glaucoma the deterioration can be identified by the changes in the visual field testing and the thinning of the nerve fibre layer (5).

Few data exist about long term course of OCT measurements and its relevance for detection of glaucoma progression. Wollstein et al.(6) found a loss of average RNFL thickness of 11,7 microns within 4,7 years in glaucoma patients.

2. AIM OF THE STUDY
The aim of the study is to emphasise which index of OCT testing shows the most important predictor values to indentify the glaucoma progression.
3.3. METHODS

Total number of 180 eyes of 120 consecutive patients were evaluated in this retrospective cross sectional pilot study. Copernicus SD-OCT with resolution of 3 mm obtained through the optic nerve head (ONH) were included. All examined eyes were divided to four groups. The size of the disc (size), the volume of cup (cup), cup/disc ratio (E/D), size of the rim (RIM) and the thickness of the retinal nerve fibre layer (RNFL) were observed in 50 healthy eyes, 50 eyes with the signs of initial open angle, preperimetric glaucoma, 50 eyes with developed simplex glaucoma and 30 eyes with terminal glaucoma signs (final glaucomatous atrophy). Exclusion criteria were ocular hypertension, other optic nerve neuropathies causing optic nerve atrophy and angle closure glaucoma. The diagnosis of glaucoma was made if one or more of the following morphological criteria were present: thinning or notching of the neuroretinal rim, loss of peripapillary RNFL, unfulfilled inferior-superior, nasal-temporal rule. Pathologic visual field test results associated with a pathologic optic disc configuration confirmed the diagnosis but were not necessary for the inclusion to the study.

These results were incorporated in the final statistical conclusion.

All statistical analyses were performed in SPSS for Windows v17.0 (SPSS Inc., Chicago, Il, USA). The specificity and sensitivity OCT parameters as potential markers progression of disease was investigated by ROC curve. Binary logistic regression was performed to show how changing variables can influence progression of the disease. All the tests were performed with the significance level of 0.05.

4. RESULTS

Results are presented in Tables 1-3 and Figures 1-2.

An increasing of the RIM for 1 unit decreases the risk to develop manifest glaucoma for 69,5%. The increasing of the E/D relation for 0,01 unit increases the risk developing manifest glaucoma for 17,75%. The enlargement of the RNFL index for one unit decreases the risk developing manifest glaucoma from initial to developing glaucoma (initial vs. developed perimetric glaucoma) 82,0%, the positive predictability 81,2%, negative predictability 77,5%. The total or final accuracy was 79,3%. AUC for CUP was 0,866 (Table 3).

The sensitivity for the E/D was 82,0%, the specificity 83,7%, negative predictability 81,3%. The final accuracy was

### Table 1. Independent predictors of progression glaucoma from initial to developed stage (binary logistic regression)

<table>
<thead>
<tr>
<th>Variables</th>
<th>B</th>
<th>S.E</th>
<th>Wald</th>
<th>df</th>
<th>p=</th>
<th>OR</th>
<th>95,0% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>RIM</td>
<td>-1,181</td>
<td>0,571</td>
<td>4,331</td>
<td>1</td>
<td>0,037</td>
<td>0,305</td>
<td>0,100  0,933</td>
</tr>
<tr>
<td>E/D</td>
<td>2,932</td>
<td>1,397</td>
<td>14,165</td>
<td>1</td>
<td>0,0001</td>
<td>0,073</td>
<td>0,019 0,285</td>
</tr>
<tr>
<td>RNFL</td>
<td>-0,081</td>
<td>0,17</td>
<td>22,436</td>
<td>1</td>
<td>0,00002</td>
<td>0,922</td>
<td>0,891 0,953</td>
</tr>
</tbody>
</table>

### Table 2. The independent predictors of progression terminal, absolute glaucoma from developing stage of the disease (binary logistic regression)

<table>
<thead>
<tr>
<th>marker</th>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
<th>PP (%)</th>
<th>NP (%)</th>
<th>AUC</th>
<th>CI</th>
<th>p&lt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>CUP cut off (0,56)</td>
<td>78,0</td>
<td>80,8</td>
<td>81,2</td>
<td>77,5</td>
<td>0,866</td>
<td>0,794-0,938</td>
<td>0,005</td>
</tr>
<tr>
<td>E/D cut off (0,305)</td>
<td>80,0</td>
<td>82,9</td>
<td>83,7</td>
<td>81,3</td>
<td>0,899</td>
<td>0,838-0,959</td>
<td>0,005</td>
</tr>
</tbody>
</table>

### Table 3. The sensitivity and specificity OCT parameters in assessment of early stages glaucoma (incipient vs. control eyes). PP – positive predictive value, NP – negative predictive value, CI – confidence interval, AUC– area under curve

<table>
<thead>
<tr>
<th>marker</th>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
<th>PP (%)</th>
<th>NP (%)</th>
<th>AUC</th>
<th>CI</th>
<th>p&lt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>RNFL cut off (99,50)</td>
<td>90,0</td>
<td>82,0</td>
<td>83,3</td>
<td>89,1</td>
<td>0,878</td>
<td>0,803-0,954</td>
<td>0,005</td>
</tr>
</tbody>
</table>

Figure 1. The sensitivity and the specificity of the OCT parameters in an assessment the initial to developing glaucoma (initial vs. developed perimetric glaucoma) PP – positive predictive value, NP – negative predictive value, CI – confidence interval, AUC– area under curve
AUC - area under curve

CI - confidence interval

NP - negative predictive value

1,58 microns per year in progressors and of -0.34 microns per year in nonprogressors. The rate of change found using Stratus OCT. The rate of progression of -1.58 microns per year was significant in glaucomatous eyes. It may represent focal damage in glaucoma causing some pores to drastically lose diameter thus increasing the the variability in the structure in optic nerve head.

Lee et al. (8) performed a trend-based analysis using Stratus OCT. The rate of progression of -1.58 microns per year was significant in glaucomatous eyes. It may represent focal damage in glaucoma causing some pores to drastically lose diameter thus increasing the the variability in the structure in optic nerve head.

DARC (Detection of apoptotic retinal cells) is going to be the new technique which utilizes the unique optical properties of the eye to directly visualize retinal ganglion cell death. It gives glaucoma physicians the opportunity to detect glaucoma earlier and monitor the response to treatment in a visual and quantifiable manner (12, 13).

Werkmeister et al. (14) emphasised that imaging in glaucoma patients focuses on ONH, the RNFL and the RGC layer. Among these options, the imaging of RGCs is the most promising approach because the loss of RGCs is directly associated with the characteristic glaucomatous visual field loss. The understanding of the structure-functional relationship in glaucoma has also significantly increased (15, 16).

6. CONCLUSION

In conclusion, we demonstrated the importance of the structural changes of optic nerve head with aim to assess the management and treatment of progressive glaucoma eye in spite of good functional results and no thinning of RNFL. Glaucomatous eyes typically had thinner RNFL but the statistically different structural ratios inside the optic nerve head. Therefore it is important to consider the changes inside the optic disc under the regular clinical glaucoma follow up.

It is an exciting time for physicians and over the next decade will certainly see advances in early detection (RNFL) and efficacious treatments and neuro-protection (structural changes of ONH).

CONFLICT OF INTEREST: NONE DECLARED.

REFERENCES


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