Abstract 8

PROGRESSION OF PACHYCHOROID NEOVASCULOPATHY INTO ANEURYSMAL TYPE 1 CHOROIDAL NEOVASCULARIZATION OR POLYPOIDAL CHOROIDAL VASCULOPATHY

Oral

Siedlecki J.*, Schworm B., Priglinger S.

University Eye Hospital Munich, LMU ~ Munich ~ Germany

Purpose:

To describe the progression of pachychoroid neovasculopathy (PNV) into pachychoroid aneurysmal type 1 choroidal neovascularization (PAT1) / polypoidal choroidal vasculopathy (PCV).

Methods:

For this retrospective cohort study, the database of the Department of Ophthalmology, Ludwig Maximilians University, Munich, was screened for patients diagnosed with and treated for PNV with a follow-up of ≥ 2 years. Multimodal imaging, including optical coherence tomography and fluorescein and indocyanine green angiography, was reviewed for the presence of choroidal neovascularization (CNV), aneurysms within/at the margins of the CNV and sub-foveal choroidal thickness (SFCT) at first diagnosis and during follow-up.

Results:

In total, 37 PNV eyes of 32 patients with a mean follow-up of 3.3 ± 1.1 (2.0–5.2) years were included in the study. At PNV diagnosis, mean age was 59.7 ± 8.7 (38.5-78.0) years and mean SFCT was 357 ± 92 (185-589) µm. During follow-up, 5 eyes (13.5 %) developed aneurysms after a mean 3.4 ± 0.8 years (2.3-4.2 years) years, defining PAT1/PCV. Risk of PAT1/PCV conversion was 7.4 %, 13.6 % and 30.7 % at years 3, 4 and 5. Lower age at PNV diagnosis (p=0.025) and sustained choroidal thickening (p=0.0025) were identified as risk factors.

Conclusions:

PNV can develop aneurysms within its type 1 CNV, defining conversion to PAT1/PCV. In this study, Kaplan Meier estimates of risk for conversion were 7.4 %, 13.6 % and 30.7 % at years 3, 4 and 5. Younger age at PNV diagnosis and sustained choroidal thickening might represent risk factors.

