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PIBIC Last Name - Yamazaki First Name - Ester Middle - Sakae

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### **PTK and Mitomycin C in Adenoviral Infiltrates : Partial Results**

**AUTHOURS:** Ester S. Yamazaki, Celina Murata, Caroline Ferraz, Norma Alleman, Denise de Freitas, Mauro Campos.

**PURPOSE:** Pre and postoperative evaluation of patients with subepithelial fibrosis caused by adenoviral infiltrates submitted to phototherapeutic keratectomy (PTK) using mitomycin C.

**PATIENTS AND METHODS:** This prospective, consecutive cases series included patients with uncorrected visual acuity (UCVA) worse than 20/40 caused by central nummular adenoviral infiltrates. They were submitted to pre and postoperative ophthalmologic examination including Ocular Responder Analyzer hysteresis®, Pentacam®, Ultrabiomicroscopy (UBM) and optical coherence tomography (OCT-Visante)®. Transepithelial phototherapeutic keratectomy using LadarWave® excimer laser with balanced saline solution mask, plus mitomycin C 0.002% for 1 minute was performed. Postoperative follow-up visits were done in day 1, 3, 7, 14, 30, 90 days.

**RESULTS:** 13 eyes of 9 patients (6 women and 3 men), with mean age 41,1 ± 14, 7 (range 20 to 65) presented preoperative mean spherical equivalent (SE): -0.12 ± 0,86 D, and +1.72 ± 1,18 D (1st month). Postoperatively, UCVA improved to equal or better than 20/40 in 76,9% and best spectacle corrected visual acuity (BSCVA) was 92,3% with gained of one or more visual lines. No patients lost visual lines. Mean corneal opacity thickness was 164,5 micron (m) at UBM, 157,7 m at Pentacam and 148,8 m with OCT preoperatively. Ablation depth was calculated considering 50m of epithelium plus 10% of total opacity, therefore mean programmed ablation was 65,14m ± 1,99. Mean central pachymetry was 491m before and 437,9 m after surgery (1st month). Pre and postoperative values of corneal total volume reduced from 55,24 to 53,29 mm<sup>3</sup> and partial volumes at 3, 5, 7 mm diameter changed to 3,54 to 3,25, 10,44 to 9,64 and 22,52 to 21,27 mm<sup>3</sup>, respectively. The coefficient of hysteresis change from 8, 91 ± 2, 25 to 7, 6 ± 1,78 (1st month).

**CONCLUSION:** Partial results in this study show that PTK with mitomycin C for fibrosis caused by adenoviral infiltrates is an option for treatment when clinical approach failure.