

Patients with a minimum corneal thickness less than 400 μm generally belong to advanced (stages III and IV) KC and were obviously excluded being out from CXL safety guidelines.³

In conclusion, I sincerely thank Kate et al for their valuable comments and for giving me the opportunity to provide 2 important final clinical recommendations:

1. Pediatric KC must be treated at diagnosis without awaiting progression. Progression will occur in almost 90% of the patients within 1 year according to the literature⁴ and, in my experience, in almost 100% of the pediatric patients engaged in eye-rubbing and suffering from allergy.⁵
2. Allergy, eye-rubbing, and pediatric age increase the risk of faster KC progression and visual acuity deterioration, increasing the risk of corneal transplant. These patients require immediate CXL treatment without awaiting progression, closer follow-up, adequate medical therapy, and careful monitoring of the results.⁵

Financial disclosures/conflicts of interest: None reported.

Cosimo Mazzotta, MD, PhD*,†

*Siena Crosslinking Center; and

†Department of Medicine, Surgery and Neurosciences, Ophthalmology Unit, Siena University, Siena, Italy

REFERENCES

1. Mazzotta C, Traversi C, Baiocchi S, et al. Corneal collagen cross-linking with riboflavin and ultraviolet A light for pediatric keratoconus: ten-year results. *Cornea*. 2018;37:560–566.
2. Krumeich JH, Daniel J, Knülle A. Live-epikeratophakia for keratoconus. *J Cataract Refract Surg*. 1998;24:456–463.
3. Raiskup F, Spoerl E. Corneal crosslinking with riboflavin and ultraviolet A: part II. Clinical indications and results. *Ocul Surf*. 2013;11:93–108.
4. Chatzis N, Hafezi F. Progression of keratoconus and efficacy of pediatric [corrected] corneal collagen cross-linking in children and adolescents. *J Refract Surg*. 2012;28:753–758.
5. Mazzotta C, Traversi C, Mellace P, et al. Keratoconus progression in patients with allergy and elevated surface matrix metalloproteinase 9 point-of-care test. *Eye Contact Lens*. 2018;44(suppl 2):S48–S53.

Management of Type 2 Bubble Formed During Big-Bubble Deep Anterior Lamellar Keratoplasty

To the Editor:

We read with interest the article by Goweida et al¹ regarding the management of Descemet “type 2” bubbles encountered during deep anterior lamellar keratoplasty (DALK). We would like to contribute 2 further points.

First, we would like to propose a fourth technique to those already succinctly described in the article, which is essentially a combination of “Descemet Membrane (DM) Baring DALK” and “Microbubble Incision DALK” techniques. After type 2 bubble creation, we perform a manual dissection until we reach the plane devoid of microbubbles, which we assume to be the pre-Descemet layer. Ever mindful of “The enemy of good is perfect”, we then perforate the ceiling of the bubble under viscoelastic protection to avoid its sudden collapse, inject viscoelastic into the bubble cavity to distance its floor and remove only the central 4 mm of pre-Descemet layer. This diameter is sufficient to clear an optical zone large enough to prevent any visual disturbance even in scotopic conditions, and in fact postoperatively, we have not recorded any subjective complaints from any patient.

Second, we agree that even in the presence of an intact DM, double anterior chamber formation can indeed occur after a type 2 bubble. In fact, having recently performed multivariate regression analysis of intraoperative factors in almost 600 patients undergoing DALK surgery (article in press), we found that the occurrence of a type 2 bubble was the single highest predictive factor of postoperative formation of a double anterior chamber, independent of intraoperative DM perforation. For this reason, if a type 2 bubble is obtained, we

suggest intracameral air fill for at least 2 hours at the end of DALK surgery as a prophylactic measure.

Financial disclosures/conflicts of interest: M. Busin has received (2006–2016) reimbursement of travel expenses and royalties from Moria (Antony, France). The remaining authors have no funding or conflicts of interest to disclose.

James Myerscough, FRCOphth*,†,‡
Asaf Friehmann, MD*,§
Cristina Bovone, MD*,†,¶
Michael Mimouni, MD||
Massimo Busin, MD*,†,¶

*Department of Ophthalmology, Southend University Hospital, Southend, United Kingdom

†Department of Ophthalmology, Ospedale Privato “Villa Igea”, Forlì, Italy

‡Istituto Internazionale per La Ricerca e Formazione in Oftalmologia (IRFO), Forlì, Italy

§Sackler Faculty of Medicine, Tel Aviv University, Tel Aviv, Israel

¶Department of Morphology, Surgery and Experimental Medicine, University of Ferrara, Ferrara, Italy

||Department of Ophthalmology, Rambam Health Care Campus and Ruth Rappaport Faculty of Medicine, Technion-Israel Institute of Technology, Haifa, Israel

REFERENCE

1. Goweida MB, Ragab AM, Liu C. Management of type 2 bubble formed during big bubble deep anterior lamellar keratoplasty. *Cornea*. 2019; 38:189–193.

Reply:

We thank Myerscough et al for their interest in our article. Regarding exposing Descemet membrane by excising 4 mm diameter of the central prepupillary stroma, we agree this technique is beneficial in cases with deep scarring and advanced macular dystrophy, where the pre-Descemet stroma (PDS) is scarred or opacified by glycosaminoglycan deposits.¹ However, we would not