Spontaneous intercalated corneal epithelial folds in thyroid eye disease: a case report

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Abstract
We observed an under-recognized, ethnic-related sign in thyroid eye disease in a 41-year-old Chinese woman. Spontaneous corneal epithelial folds form after fluorescein staining as a result of tear film disruption related to orbital tension and upper lid pressure on the cornea. They occur in anatomically predisposed eyes similar to acquired lower lid epiblepharon. Like their chorioretinal counterparts, presence of corneal epithelial folds prompt for further workup, even in East Asian patients whose thyroid eye changes may appear less impressive.

Key words: Asian continental ancestry group; Epithelium, corneal; Graves ophthalmopathy

Case presentation
In June 2018, a non-smoking 41-year-old Chinese woman presented with a 3-month history of worsening periocular swelling and tightness. She had a 4-year history of stable Graves disease after radioactive iodide treatment, and she was taking 50 mg thyroxine supplement. On examination, visual acuity of each eye was 20/30, with normal Ishihara and pupillary responses and visual field on confrontation. Intraocular pressure was 13 mm Hg and 11 mm Hg at primary gaze and 17 mm Hg and 19 mm Hg on upgaze for right and left eye, respectively. Spontaneous, intercalated corneal epithelial (CE) folds were evident only after fluorescein eyedrop staining and under cobalt-blue lights (Figure 1) before applanation tonometry. She had bilateral upper lid puffiness, retraction, and increased resistance to retropulsion. Despite having mild exophthalmos of 18 mm bilaterally on Hertel exophthalmometry, normal eye movement, and a low clinical activity score of 1 on eyelid swelling, she was referred for further oculoplastic evaluation that showed no chorioretinal fold or optic disc swelling. The patient’s anterior chamber depth, intraocular pressures, anterior segment optical coherence tomography, and corneal topographies (Figure 3) were all normal.

Magnetic resonance imaging demonstrated minimally enlarged or inflamed extraocular muscles with predominant fat expansion (Figure 2). Combined intravenous pulse methylprednisolone with orbital radiotherapy was proposed as compassionate treatment for progressive and symptomatic thyroid eye disease despite a low clinical activity score, which was well tolerated.

At the 3-month follow-up, periocular swelling and exophthalmos improved, but CE folds persisted. Patient was offered options of staged rehabilitative orbital decompression, upper lid recession, and blepharoplasty for residual deformities. At the 21-month follow-up, CE folds were static.

Discussion
To the best of our knowledge, there has been only one case report describing corneal stromal (deep) striae (folds) persisting after endoscopic decompression for optic neuropathies in an Asian woman with thyroid eye disease. These corneal stromal striae were evidence of raised orbital
Folds form at the front (cornea) or back (chorioretina) of the eye when tissue layers buckle back as vertical lines. Chorioretinal folds are another sign indicating raised orbital pressure in thyroid eye disease. We propose that CE folds or stromal striae are more prevalent among the anatomically predisposed eyes, similar to acquired lower lid epiblepharon in thyroid eye disease. In East Asians whose upper lid creases are low or absent, progressive fat expansion beneath a tight orbital septum exerts pressure on the most superficial (ie epithelial) layer of the cornea leading to formation of CE folds.

In our patient, CE folds did not interfere with visual acuity, analogous to microstriae after LASIK, although contrast sensitivity, higher order aberration or Maddox rod effect was not evaluated. Similar to chorioretinal folds, there is no treatment guideline for CE folds per se in thyroid eye disease. Soft tissue signs improved although CE folds persisted after combined pulse methylprednisolone and radiotherapy. For patients with no demonstrable visual consequence, continuing management of any associated ocular surface problem (eg dry eye) is an acceptable option. In patients with symptomatic periocular deformities (including upper lid swelling, retraction, and exophthalmos), structural changes after orbital decompression, upper lid recession, and blepharoplasties may lead to resolution of CE folds. Persistent chorioretinal folds, attributed to scleral remodeling, was reported despite complete removal of retrobulbar tumor.
Conclusion

The presence of CE or chorioretinal folds in patients with thyroid eye disease may prompt further oculoplastic evaluation or radiological workup, especially in East Asian patients whose periorbital deformities or disease activities are often minimal or subtle.

Author contributions

Concept or design: All authors
Acquisition of data: All authors
Analysis or interpretation of data: All authors
Drafting of the article: All authors
Critical revision for important intellectual content: All authors

All authors had full access to the data, contributed to the study, approved the final version for publication, and take responsibility for its accuracy and integrity.

Conflicts of interest

As an editor of the Journal, KKL Chong was not involved in the peer review process for this article. Other authors have disclosed no conflicts of interest.

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Ethics approval

The patient was treated in accordance with the Declaration of Helsinki. The patient provided written informed consent for treatments and procedures and for publication.

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