

Managing corneal defects:

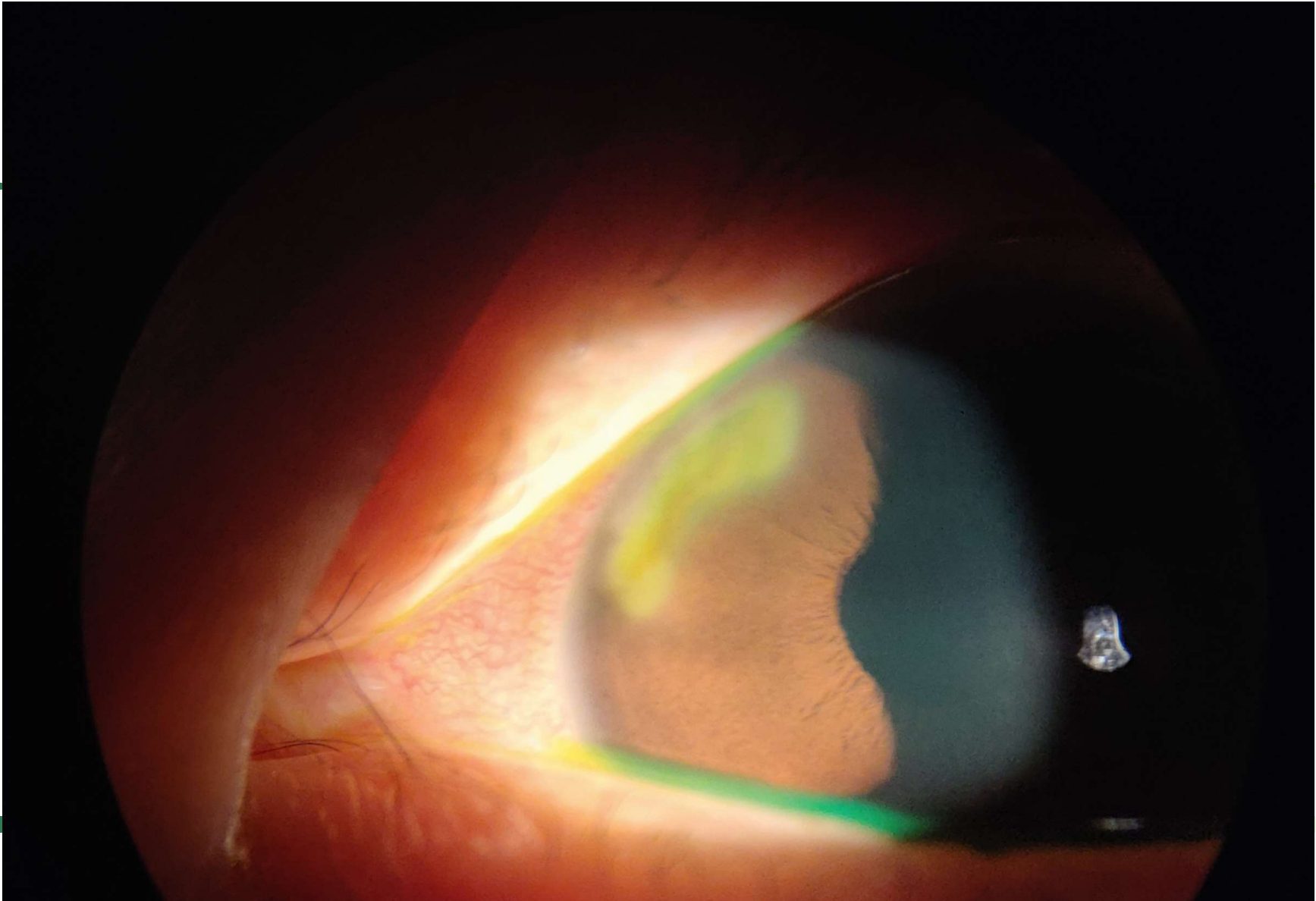
Marginal keratitis and Recurrent erosion

Case 1: JW

- Mrs JW; 48yo female non-CL wearer (prev worn 10 years prior)
 - Presented to optom with 1 day history of mild red left eye and FB sensation
 - Referred onwards for “subepithelial/anterior stromal opacification”, epithelial ulceration and mild hyperemia. Started on 2-hourly preservative free lubricants
 - “lubricant helps a bit a times but not all the time”
 - “Pain in the bright light in the morning”; photophobic
 - No recent trauma or injury
 - Feels a bit like pressure on the eye
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Examination

- Left eye:
 - Moderate conjunctival hyperemia
 - Epithelial ulcer and stromal infiltrate
 - No AC reaction
 - Moderate SPK
 - Mild bilateral MGD and blepharitis
 - BCVA R 6/7.5+, L 6/7.5-
 - IOP R 13.5 L 12mmHg (NCT)
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- What's your diagnosis?
 - Management?
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Management

- Diagnosis; left marginal keratitis
 - Management:
 - Maxidex qid 1 week then short taper
 - Chlorsig qid 1 week
 - Reviewed 10 days later
 - Much improved but still mild infiltrate
 - Epithelial defect nearly resolved
 - MGD and moderate SPK remains
 - Continue tapering Maxidex
 - Chlorsig continued until epithelial defect closes
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- Final resolution 4 weeks after presentation
 - No infiltrate or epithelial defect remains
 - Continue lubricants and blepharitis management
 - Mild IOP spike (26.5mmHg)
 - Resolved after ceasing steroid
 - No glaucomatous neuropathy currently evident
 - Counselling regarding future steroid eyedrop/oral intake use
 - Discharge to optometry care for annual review
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Marginal keratitis

- First described by Thygeson in 1946
 - Peripheral corneal inflammatory response
 - Bacterial antigens trigger immune response in sensitized cornea
 - Affects peripheral corneal tissue
 - Particularly susceptible to immune-mediated responses due to proximity to limbal vasculature and conjunctival lymphoid tissue
 - Antigen/antibody complexes deposited in peripheral cornea
 - Antigen from Staphylococcal bacteria in tear film and on lid margin
 - Antibody to this derived from limbal blood vessels
 - Inflammatory cascade, neutrophil recruitment and infiltrate formation
 - Eventually overlying epithelium breaks down causing defect
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Signs and Symptoms

- Symptoms variable from asymptomatic to severe including FB sensation, epiphora, burning, grittiness, photophobia
 - Findings:
 - Round or curvilinear infiltrate on the peripheral cornea
 - Clear margin between limbus and infiltrate of 1-2mm
 - Epithelial breakdown overlying with SPK or ulcer
 - Injection of bulbar conj and mild chemosis of palpebral conj
 - Lid erythema and reactive ptosis
 - Blepharitis (all forms) usually present
 - Mild AC reaction rare but does occur
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Management

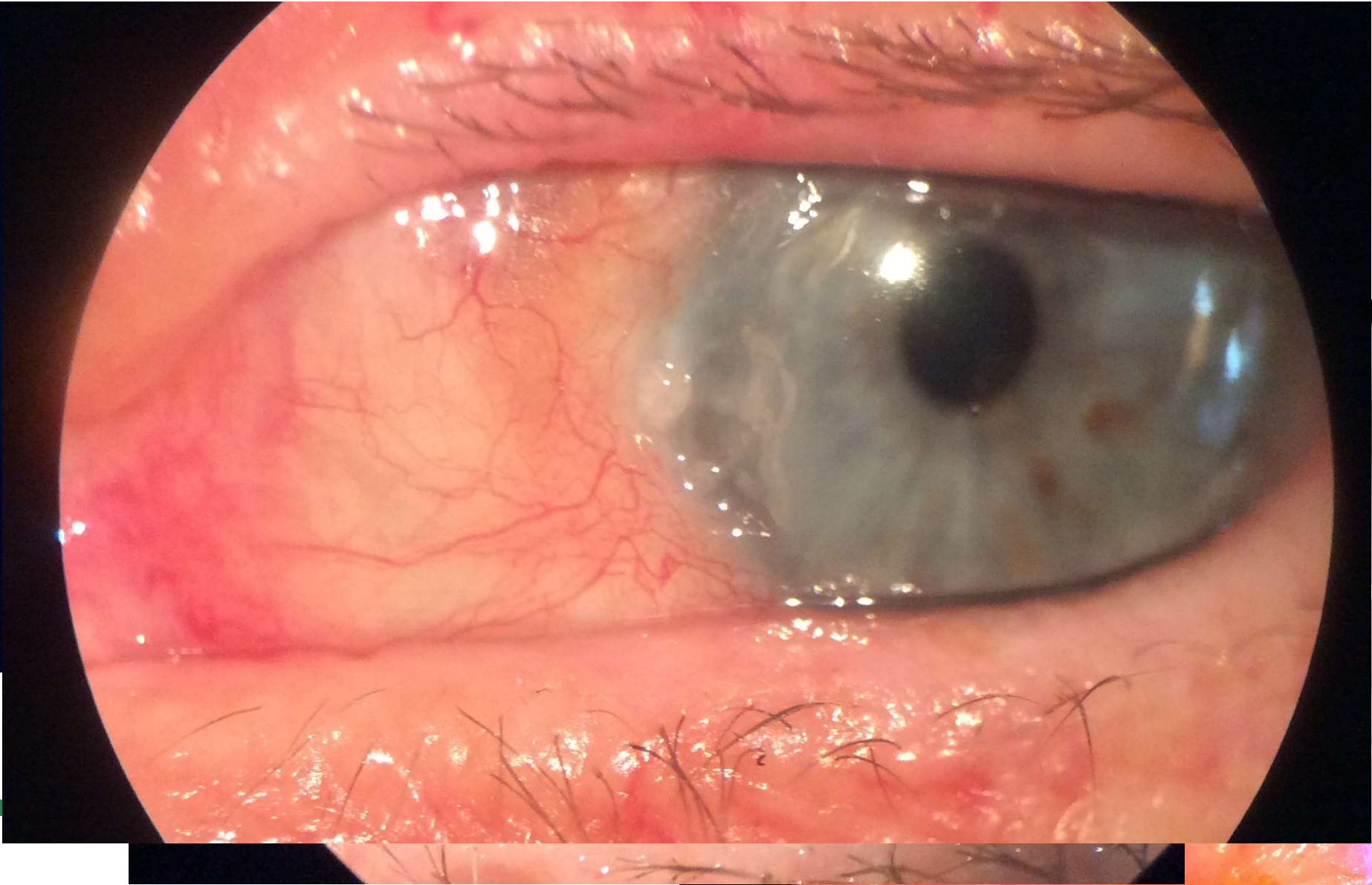
1. Extinguish the inflammatory response
 2. Treat the underlying cause
- Inflammation treated with topical steroid
 - Flarex or Maxidex are often good choices applied 3-4 times daily
 - Study: predsol more effective than no tx;
 - neomycin not more effective than no tx
 - Reduce antigen burden
 - Manage blepharitis (improve lid hygiene)
 - Add tea tree oil to lid hygiene for demodex blepharitis
 - Consider oral doxycycline/azithromycin management for ocular rosacea
 - Topical antibiotic (Chlorsig) whilst epithelium heals
 - Treat any underlying ocular surface issues
 - Consider any contributing systemic issues
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Differential diagnoses

- Microbial keratitis
 - Generally more central, unilateral, often singular large lesion
 - Severe inflammatory response with AC reaction, severe hyperemia, possible hypopyon
 - Increased symptoms with severe photophobia, decreased vision, epiphora
 - HSV keratitis
 - Branching ulcer or diffuse SPK often with less dense infiltrate and no clear zone
 - Unilateral, no blepharitis, prior history, reduced corneal sensitivity
 - Consider no topical steroid if unclear marginal keratitis vs HSV
 - Drug hypersensitivity:
 - Dorzolamide, atropine, pilocarpine, gentamicin, phenylephrine
 - CLIPU; contact lens-related
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DDx: PUK

- Peripheral inflammatory keratitis
 - Necrotizing scleritis, epithelial ulceration, corneal thinning
 - Rapid corneal melting and perforation if untreated
 - Variable presentation
 - Pain on eye movements (scleritis), hyperemia, blurred vision, photophobia, epiphora, other anterior/posterior inflammation may be present
 - often unilateral, limited to one sector of cornea, within 2mm of limbus, can extend over limbus onto sclera with no clear zone
 - Can be due to infection
 - 50% of non-infectious cases have underlying autoimmune condition
 - Review history and ask questions
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Moorens ulcer

- Rare; estimated 0.03% incidence in Chinese population study
 - Circumferential progressive and rapid thinning and ulceration
 - Can result in perforation
 - 25%-30% of cases bilateral
 - No clear zone between limbus and ulcer
 - Undermined and infiltrated leading edge of ulcer
 - Severe pain, progressive, chronic autoimmune condition
 - Sometimes idiopathic
 - Some association with prior infection with HepC, trauma, parasitic infection
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Terrien's degeneration

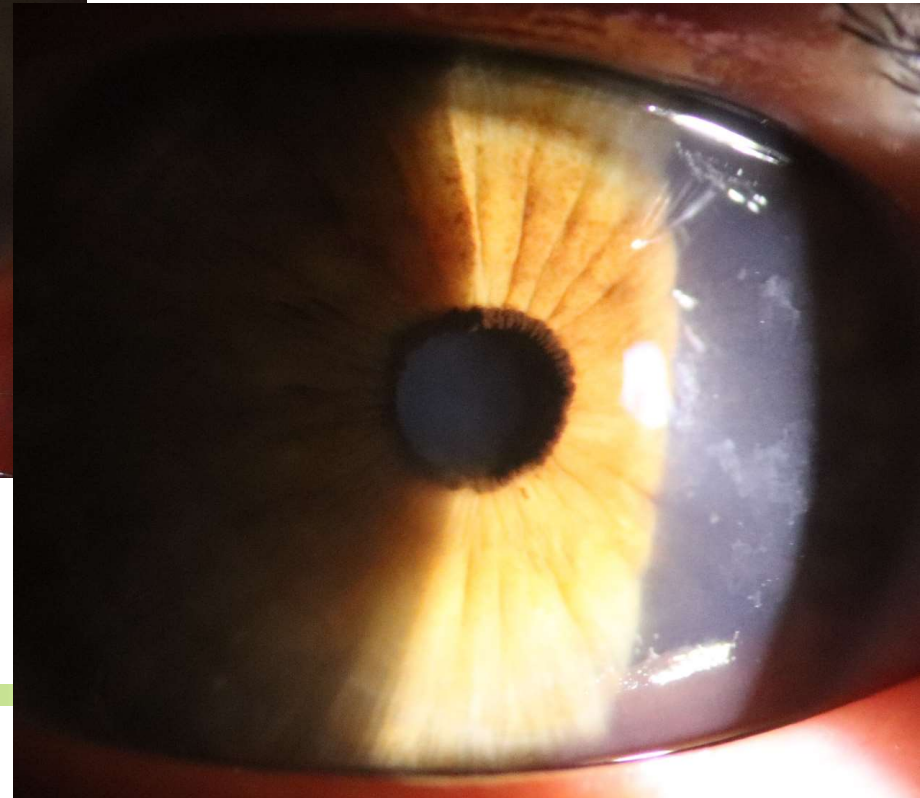
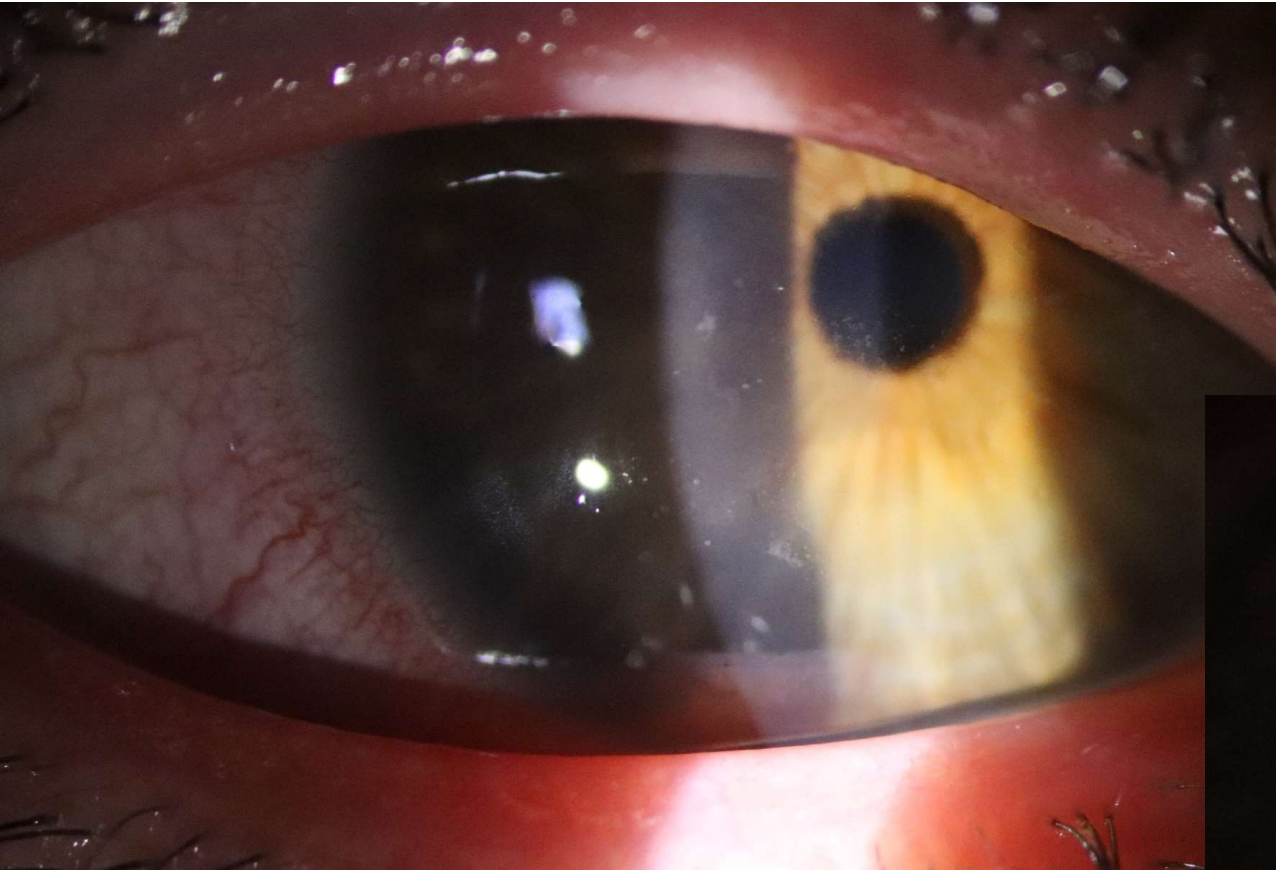
- Bilateral painless degeneration of peripheral cornea
 - Vision loss due to increasing corneal astigmatism
 - Eye white and quiet
 - Idiopathic
 - Fine yellow-white stromal opacities separated from limbus by clear zone
 - No epithelial defect
 - May look like arcus senilis
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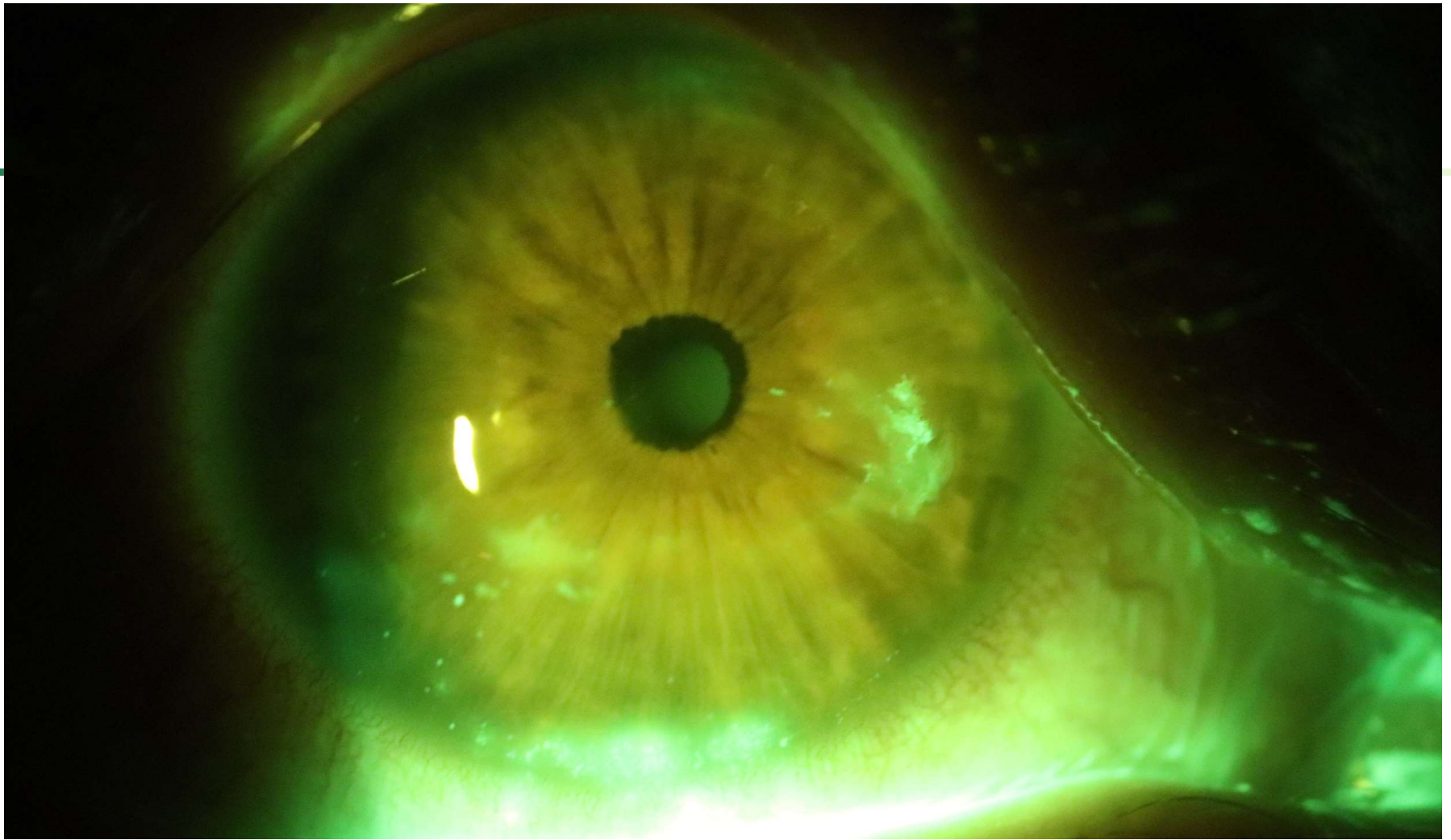
Case 2: RCE

- Young man with 1 day history of pain on waking
 - Used VitaPos during the day and lubricants
 - Vision 6/7.5-
 - Inferior corneal RCE partially reepithelialized
 - No infectious keratitis visible
 - BCL inserted. Gel at night. Topical lubricants
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Review

- Vision worsened 6/12
 - Red and inflamed
 - Sloppy mobile epithelium
 - Debrided worst area
 - BCL reapplied
 - Flarex and chlorsig qid
 - Plenty of lubricants
 - Review 2 days
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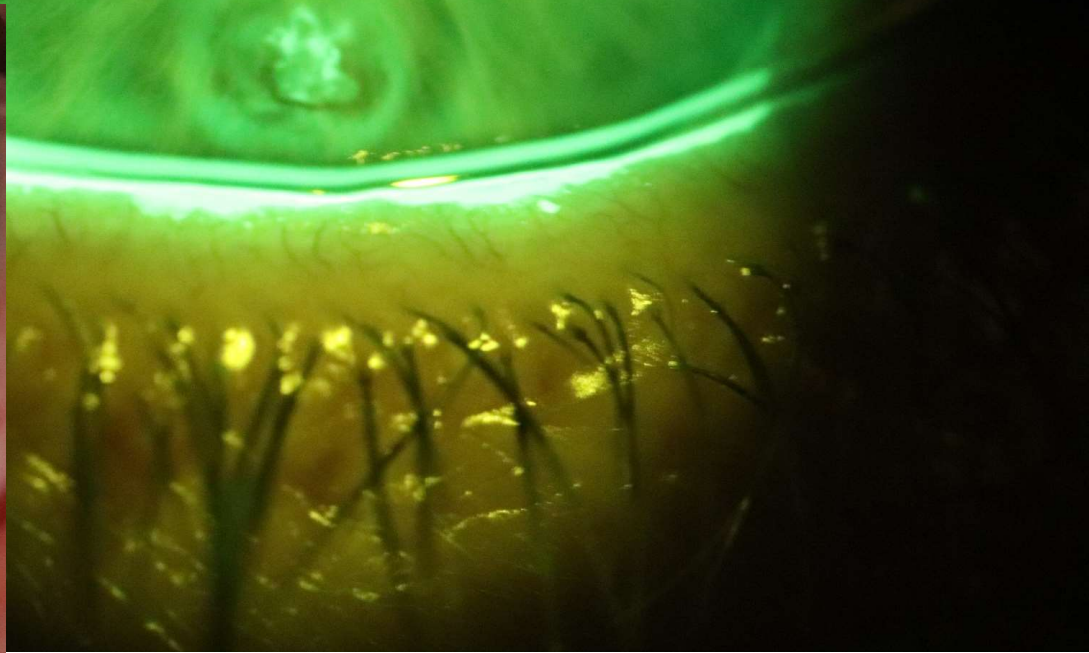
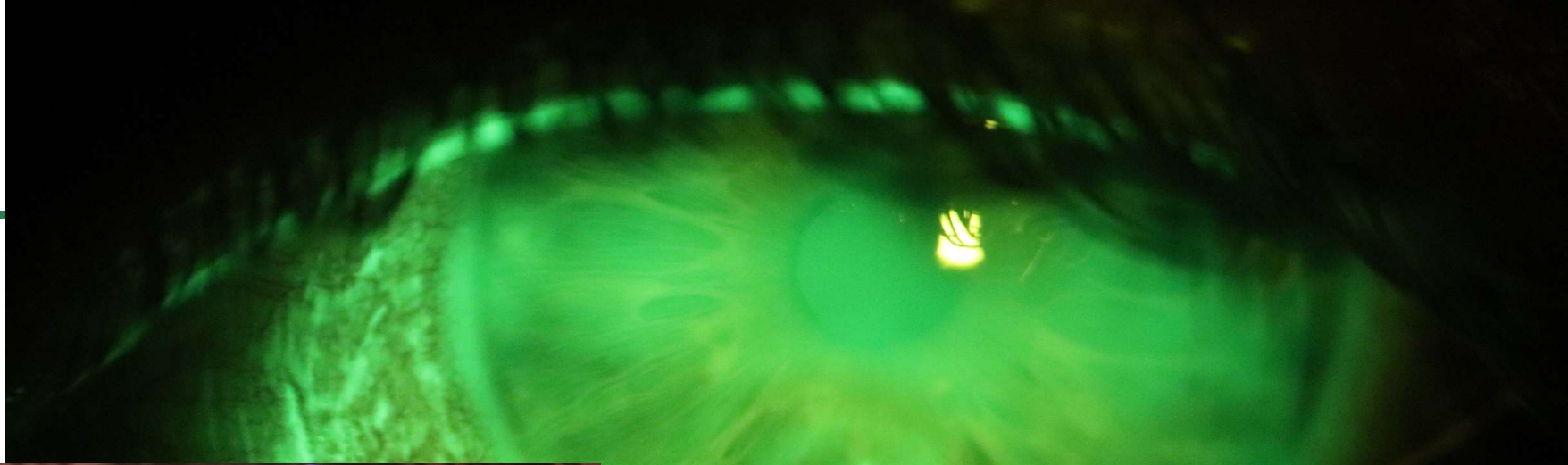


2nd review

- Improving, VA 6/10=
 - Symptoms improved
 - Eye quiet and white
 - New BCL inserted
 - Review 3 days and trial without BCL
 - Given size and mobility of erosion may require superficial keratectomy
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Case 3

- 2-3 times weekly wakes with pain in the eye
 - Using Hylo-forte and VitaPos ointment daily
 - History; 2019 fingernail injury to eye
 - 6/15 acuity today
 - Gr 1+ AC reaction and moderate hyperemia
 - RCE visible
 - Started Maxidex and Chlorsig qid
 - BCL inserted
 - Continue Hylo-forte
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Follow-up

- 1/52 review; vision 6/6, mild staining at centre of prior defect
 - Faint stromal haze
 - AC quiet
 - Cease Chlorsig, Maxidex bd , Vitapos at night without fail and Hylo-forte during day
 - BCL removed
 - 1/12 review; vision 6/7.5-
 - No recurrence. Looks great
 - Lubricate and Vitapos without fail.
 - Cease maxidex
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Recurrent corneal erosion

- Repeated epithelial defects sustained usually on waking due to poor adhesion of epithelium to basement membrane
 - Typically lasts minutes to hours but can last days
 - Redness, photophobia, blurred vision, epiphora
 - Diagnosed with clinical signs and history
 - Conjunctival injection, new or healing epithelial defect
 - Typically unilateral but may be bilateral
 - History of prior trauma
 - Px describes waking some mornings with significant pain as soon as they open their eyes
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Risk factors and epidemiology

- Most common subsequent to minor shallow corneal trauma (45-64%)
 - Second most common cause underlying EBMD (19-29%)
 - Other factors include corneal dystrophy, denegeration (band keratopathy), prior ocular surgery (cataract, corneal transplant, refractive laser)
 - Co-morbidities that increase risk include;
 - Ocular rosacea and MGD
 - Blepharitis and dry eye
 - Diabetes
 - Common onset in px aged 40s and 50s
 - Slight female predominance
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Mechanism and pathophysiology

- Trauma results in inflammation and disruption of adhesion complexes (hemidesmosomes and anchoring fibrils)
 - Upregulation of MMP-2 and -9 further destabilises
 - Corneal epithelium drying overnight increases force of adhesion between epithelium and lid
 - Shearing forces on waking rip off the already weak epithelium
 - Presence of increased bacterial lipase combining with meibum
 - Produces toxic free fatty acids which disrupt healing and predispose px to erosions
 - Corneal trauma more commonly results in RCE macroerosions
 - Hours to days to heal
 - Signs include large epithelial defect, pain and oedema
 - EBMD more commonly related to RCE microerosions
 - Minutes to hours to resolve
 - SPK or no epithelial defect visible on exam; check fellow eye for EBMD
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DDx of RCE

- Be wary of RCE patients with AC activity
 - Highly unusual in RCE although does occur
 - Consider if infection present (microbial keratitis)
 - If signs of inflammation are greater than expected for RCE
 - Or symptoms (pain, reduced VA, photophobia) more intense than expected
 - Carefully assess for dendritic ulceration
 - Especially where there is no definite epithelial erosion
 - Consider other causes:
 - Neurotrophic ulcer, trauma, sterile ulcer (CLIPU, marginal keratitis)
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Treatment of RCE

- Goals of treatment:
 - Pain relief
 - Promote epithelial healing
 - Prevent recurrences
 - Consider patient symptoms, rate of recurrence and pathophysiology
 - 60% of patients treated medically still have symptoms
 - Rate of healing of complex adhesions is 3-6 months
 - Prophylactic treatment should continue for this period
 - Acute treatment can be ceased once initial erosion is healed
 - Treat underlying comorbidities (MGD, rosacea, blepharitis)
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Management: conservative

- Ointment at night and lubricate
 - Hylo-forte, theratears or other non-preserved at least 4-6 times daily
 - VitaPos or similar without fail every night
 - Consider cycloplegia or BCL for pain management
 - May require oral analgesia
 - BCL
 - Excellent for pain management
 - Prevent shearing forces of lid on waking and scaffold for healing
 - Flat base curve (not too tight) and high O₂ transmissibility
 - 6-12 weeks for restoration of adhesion complexes
 - Consider hypertonic ointment at night
 - Creates temporary osmotic gradient to dehydrate cornea and increase epithelial adhesion
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- Debridement
 - Weck cell or spud and remove all loose epithelium
 - Remove loose epithelium to the point of tight adherence
 - Leave a margin of 1-2 mm from limbus to protect stem cells
 - Topical anaesthesia and avoid sharp instruments
 - BCL, cycloplentolate, lubricate, antibiotic and bedtime ointment 5-6 months
 - Debridement and debur
 - Reduces major recurrences significantly
 - 5-10 seconds buffing of basement membrane until smooth
 - Don't penetrate Bowman's membrane
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- Surgical options

- Punctate adhesions

- Applied with needle through loose epithelium to anterior stroma
 - 0.5-1mm apart and 1-2mm into normal tissue
 - Don't puncture stroma; visualize with NaFl
 - Not across pupil margin due to scarring

- PTK

- Superficial keratectomy

Evidence for treatment

- Cochrane review:
 - No high quality evidence
 - BCL vs lubricants:
 - No significant difference in time to resolution of total participants with improvement
 - PTK vs alcohol delamination
 - Similar to resolution
 - Increased risk of recurrence for patients treated with PTK
 - Transepithelial vs subepithelial excimer laser ablation
 - Possibly increased risk of recurrence for trans vs sub but overlapping confidence intervals
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- Hong Kong study, 48px

- Diamond bur keratectomy vs debridement only

- Recurrences less common in those treated with the bur
 - Similar corneal haze in both groups

- 4% major recurrence in px with diamond bur tx vs 56% in debridement px

- 20% minor recurrence in px with diamond bur tx vs 65% in debridement px

- No px with diamond bur tx required re-treatment vs 52% with debridement

- Concluded simple debridement can speed resolution but doesn't prevent further episodes

Conclusions

- Think about the goals of treatment
 - Educate the patient on
 - The potential outcomes and likelihood of complete resolution
 - The probability of requiring long-term lubricant management
 - Think about the pathophysiology of RCE when selecting treatment
 - Consider a step-wise systematic approach
 - Refer if severe recurrences are continuing
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