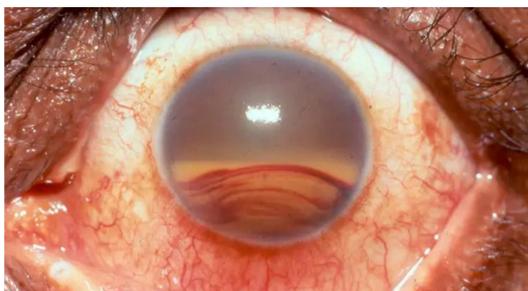


Just eye pain? No, it's UGH syndrome

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BACKGROUND

- Uveitis-Glaucoma-Hyphema (UGH) or Ellingson syndrome is a rare but serious complication that can occur after cataract or any intraocular surgery.
 - It is a result of the subluxation or malposition of the intraocular lens, damaging adjacent ocular structures and leading to microhyphema, hyphema, anterior chamber inflammation, and elevated intraocular pressures.^{2,4}
- These patients often have uncomplicated cataract implants with episodes of blurry vision occurring weeks to months following surgery.
- If left untreated, UGH syndrome can result in nerve damage, macular edema, and permanent vision loss.
- Current treatment options include intraocular lens exchange, topical and systemic medication, and cyclophotocoagulation, the placement of a capsular tension ring to redistribute zonular tension and anti-vascular endothelial growth factor (anti-VEGF) therapy.
- Incidence of UGH is approximately 0.4 to 1.2.³



Images are not from patient case; they are from American Academy of Ophthalmology.^{5,6}

SUMMARY OF CASE

- A 71-year-old female with end-stage renal disease on dialysis, diabetes mellitus type 2, was transferred for altered mental status after reportedly missing dialysis treatment the prior week.
- CT head without contrast showing no signs of intracranial bleed or stroke.
- Her encephalopathy improved the following morning, alerting the team to her right eye pain.
- Collateral information revealed patient had been complaining of eye pain and headaches earlier in the week and that patient underwent recent cataract surgery 2-3 months ago.
- Physical examination revealed right supraorbital edema with injected conjunctiva of the right eye, tearing, redness and difficulty opening her eyelid without significant pain
- Ophthalmology was consulted and found an elevated IOP in the 50-60s, 25% hyphema in the anterior chamber, with intact extraocular muscles, highly suggestive of UGH syndrome, given the patient's recent cataract surgery.
- Patient was started on IV acetazolamide, topical Combigan and dorzolamide OD which resulted in improved ocular pain, but no notable improvement in intraocular pressure.
- Given our current facility did not have access to ophthalmological surgical intervention, the decision was made to transfer the patient to another hospital to undergo evacuation of the hyphema.

DISCUSSION

- UGH syndrome typically presents weeks to months post cataract surgery, with ocular pain, photophobia, or persistent blurry vision.
- Treatments includes surgery or medical management, but surgery tends to have better outcomes.
- In this case we attempted to treat the patient medically given our limited access to local ophthalmological surgical intervention.
- The combination of IV acetazolamide, topical Combigan and dorzolamide OD alleviated the patient's ocular discomfort but in the end was ineffective at reducing her intraocular pressures.

CONCLUSIONS

- UGH syndrome is a rare but serious complication following cataract surgery and should be closely monitored in patients post-operatively.
- UGH syndrome can appear at any time, particularly in patients with predisposing factors such as pseudoexfoliation, prior vitreoretinal surgery, or trauma.
- In UGH syndrome, the results of surgical intervention are superior to that of conservative treatment, however, surgery does not guarantee resolution.¹

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