

References

- 1 Cukras C, Agrón E, Klein ML, Ferris 3rd, FL, Chew EY, Gensler G *et al*. Natural history of drusenoid pigment epithelial detachment in age-related macular degeneration: Age-Related Eye Disease Study Report No. 28. *Ophthalmology* 2010; **117**: 489–499.
- 2 Parodi MB, Virgili G, Evans JR. Laser treatment of drusen to prevent progression to advanced age-related macular degeneration. *Cochrane Database Syst Rev* 2009; Issue 3: (Art. No. CD006537)10.1002/14651858.CD006537.pub2).
- 3 Margolis R, Ober MD, Freund KB. Disappearance of drusen after rhegmatogenous retinal detachment. *Retinal Cases & Brief Reports* 2010; **4**: 254–256.
- 4 Gallego-Pinazo R, Suelves-Cogollos AM, Dolz-Marco R, Arevalo JF, García-Delpech S, Mullor JL *et al*. Intravitreal ranibizumab for symptomatic drusenoid pigment epithelial detachment without choroidal neovascularization in age-related macular degeneration. *Clin Ophthalmol* 2011; **5**: 161–165.
- 5 Kishore K, Jain S, Sharma YR, Kashyap B. Disappearance of drusen after intravitreal anti-VEGF injections for sub-macular hemorrhage (SMH) secondary to neovascular macular degeneration. *IOVS* 2012; **53**, (ARVO e abstract 2912).
- 6 Krishnan R, Lochhead J. Regression of soft drusen and drusenoid pigment epithelial detachment following intravitreal anti-vascular endothelial growth factor therapy. *Can J Ophthalmol* 2010; **45**(1): 83–84.
- 7 Holz FG, Staudt S. Disappearance of soft drusen following macular hole surgery. *Retina* 2001; **21**: 184–186.
- 8 Stefánsson E. Physiology of vitreous surgery. *Graefes Arch Clin Exp Ophthalmol* 2009; **247**(2): 147–163.

S Dithmar, S Pollithy and T Ach

Department of Ophthalmology, University Hospital Heidelberg, Heidelberg, Germany
E-mail: stefan.dithmar@med.uni-heidelberg.de

Case report was presented at annual meeting of the German Society of Ophthalmology 2012 (DOG 2012).

Eye (2013) **27**, 779–781; doi:10.1038/eye.2013.35;
published online 5 April 2013

Sir,
Acute cilio-choroidal effusion due to acetazolamide: unusual posterior involvement (OCT aspects)

Acetazolamide, a sulphonamide-derived medication, frequently used in glaucoma and after cataract surgery, can very rarely cause idiosyncratic reaction, and few reports are present in literature.^{1–3}

We report ocular coherence tomography (OCT) scans of the posterior pole in a case of ciliary body oedema after the drug administration causing bilateral angle-closure glaucoma (ACG). In our case we found a massive choroidal effusion with posterior retinal folds and papillary oedema, never described before in literature.



Figure 1 Fundus photo showing evident disc swelling and retinal folds.

Case report

The drug reaction occurred in a 71-year-old white man who was prescribed a single oral dose of acetazolamide (250 mg) after cataract surgery and IOL implantation under local anesthesia.

On examination, both eyes showed congestion and oedema of the inferior bulbar conjunctiva, heavy cloudy cornea, very shallow anterior chamber. IOL was shifted forward. Intra ocular pressure was 52 mm Hg in right eye and 60 mm Hg in left eye. A diagnosis of ACG was made. Fundus examination was characterized by bilateral peripheral choroidal detachment and papillary swelling (Figure 1). Posterior OCT scans confirmed papillary oedema together with retinal folds (Figure 2) and nerve fiber layer thickening.

Acetazolamide has been suspected to be the cause of the secondary ACG, and after it was discontinued the effusion receded rapidly.

Comment

Few cases of acute secondary ACG with choroidal effusion and anterior shift of the iris-lens diaphragm have been associated with acetazolamide compared with other sulphonamides.^{1–3}

With regard to the posterior involvement there are only few reports of retinal folds attributed to topiramate and hydrochlorothiazide.^{4,5}

Papillary oedema has never been associated with sulpha drugs. Posterior involvement with retinal folds and papillary oedema due to acetazolamide has never been described before.

OCT was able to document this effusion caused by the absence of any barrier in the prelaminar region that could inhibit the diffusion of fluid from the choroid into the papilla and peripapillary region.

Retinal folds were caused by the choroidal effusion contained by the barrier of retinal pigment epithelium and the inextensible scleral coat.

OCT shows the extensive posterior pole involvement and the resolution of the rare adverse reaction

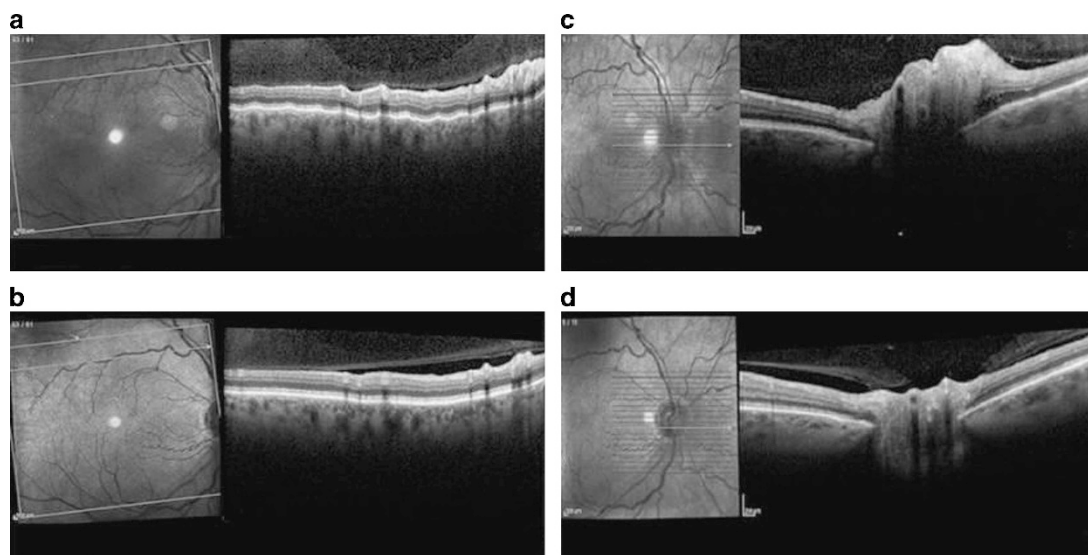


Figure 2 High-resolution OCT images of the posterior pole during acetazolamide administration with retinal folds (a) and after the resolution of the effusion (b). High-resolution OCT images of the posterior pole during acetazolamide administration with papillary oedema (c) and after the resolution of the effusion (d).

manifestations after the drug administration was discontinued.

Conflict of interest

The authors declare no conflict of interest.

References

- 1 Mancino R, Varesi C, Cerulli A, Aiello F, Nucci C. Acute bilateral angle-closure glaucoma and choroidal effusion associated with acetazolamide administration after cataract surgery. *J Cataract Refract Surg* 2011; **37**: 415–417.
- 2 Senthil S, Garudadri C, Rao HBL, Maheshwari R. Bilateral simultaneous acute angle closure caused by sulphonamide derivatives: a case series. *Indian J Ophthalmol* 2010; **58**: 248–252.
- 3 Parthasarathi S, Myint K, Singh G, Mon S, Sadasivam P, Dhillon B. Bilateral acetazolamide-induced choroidal effusion following cataract surgery. *Eye* 2007; **21**: 870–872.
- 4 Guier CP. Elevated intraocular pressure and myopic shift linked to topiramate use. *Optom Vis Sci* 2007; **84**: 1070–1073.
- 5 Roh YR, Woo SJ, Park KH. Acute-onset bilateral myopia and ciliochoroidal effusion induced by hydrochlorothiazide. *Korean J Ophthalmol* 2011; **25**: 214–217.

R Malagola, R Giannotti, L Pattavina and L Arrico

Department of Ophthalmology, University of Rome “La Sapienza”, Rome, Italy
E-mail: romualdo.malagola@uniroma1.it

Eye (2013) **27**, 781–782; doi:10.1038/eye.2013.41;
published online 5 April 2013

Sir, Spontaneous haemorrhage in an eyelid hidrocystoma in a patient treated with clopidogrel

Hidrocystomas are benign cystic tumours of the sweat glands that frequently occur in the periocular region. Rarely, more serious pathology such as basal cell carcinoma or malignant melanoma may resemble and be mistaken for hidrocystoma.^{1,2} We report a very unusual presentation of spontaneous bleeding within a hidrocystoma, mimicking a malignant melanoma.

Case report

A 62-year-old woman presented with a lesion on the medial aspect of the left upper lid, which was slowly enlarging over 2 years and darkened 6 months earlier. She admitted to having excessive sun exposure in the past, and her past medical history included hypertension, type II diabetes, and peripheral vascular disease. Her regular medications were insulin, metformin, simvastatin, ramipril, and clopidogrel.

Examination revealed a curious lesion on the medial aspect of the left upper lid (Figure 1a). The lesion, which appeared cystic and lobulated, was mostly skin coloured with surface telangiectasia and purple–blue discoloration inferiorly. Although a benign lesion was suspected, the unusual appearance prompted excisional biopsy. Histopathological examination showed a papillary hidrocystoma with apocrine differentiation, featuring small lakes of blood within the myxoid material occupying the lumen (Figure 1b).

Comment

Apocrine hidrocystomas usually present as slow-growing, solitary, or multiple small, tense,