

Letter to the Editor



Zoster sine herpete: a disease that ophthalmologists should be aware of

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TO THE EDITOR

We read with great interest the review by Zhou et al. [1] in the article "Zoster sine herpete: a review". Herpes zoster ophthalmicus is a well-known consequence of varicella zoster virus (VZV) reactivation, occurring in 10-20% of herpes zoster cases, and may cause potentially sight-threatening complications anywhere from the anterior segment through to the optic nerve, retina, and central nervous system [2]. Although less reported in the literature, zoster sine herpete (ZSH) also presents with intraocular manifestations which, if misdiagnosed and left untreated, can result in adverse visual outcomes.

As mentioned by Zhou et al., corneal involvement may occur in the form of keratitis [1,3]. Although there is no classic ophthalmic presentation, it appears in both the literature and anecdotally that the most common case is a raised intraocular pressure associated with anterior uveitis [4-6]. Slit lamp examination may show inflammatory deposits on the posterior corneal surface, anterior chamber inflammation, trabecular meshwork pigmentation, iris atrophy, and/or adhesions between the iris and the lens (posterior synechiae), causing a distorted pupil. Inflam-

mation may persist as chronic anterior uveitis, which can lead to uveitic glaucoma, a potentially sight-threatening complication [5]. In rare cases, acute anterior uveitis with a hyphaema has been the clinical presentation of ZSH [7,8].

We acknowledge that the key to diagnosing ZSH is having a high level of suspicion and subsequent laboratory testing. Polymerase chain reaction (PCR) testing for VZV DNA in the aqueous humour is essential for definitive diagnosis and subsequent management with antiviral therapy. A higher viral load in aqueous sampling is significantly associated with uveitic complications of iris atrophy and pupil distortion [4], which suggests that ongoing VZV replication will perpetuate inflammation and increases the risk of associated complications. This highlights the importance of early initiation and appropriate duration of antiviral therapy to minimise this risk.

Limited published evidence on ZSH in general, and even more so in an ophthalmology context, means that there are no established guidelines on recommended management and therapy. Our recommendation is that ZSH should be considered in patients presenting with anterior uveitis with elevated intraocular pressure, but particularly in those with slit lamp findings of iris atrophy, or pupil dis-

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404 Al-Ani and Niederer

tortion. Performing an anterior chamber tap for VZV PCR is necessary to make a definitive diagnosis, but treatment with antiviral therapy in cases of high clinical suspicion should not be delayed if PCR testing is unable to be performed.

CONFLICT OF INTEREST

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