LETTER TO THE EDITOR

Comment re: Unilateral solitary choroidal granuloma as presenting sign of secondary syphilis

Chiara M. Eandi · Enrico Bertelli · Emmett T. Cunningham Jr.

Received: 23 February 2013 / Accepted: 11 March 2013 / Published online: 27 March 2013 © Springer-Verlag Berlin Heidelberg 2013

Dear Editor:

We read with interest the paper by van der Vaart and colleagues [1] entitled "Unilateral solitary choroidal granuloma as presenting sign of secondary syphilis," and were struck by the similarity of the color fundus images of the two presented cases to the findings in Acute Syphilitic Posterior Placoid Chorioretinopathy (ASPPC) [2]. Our recently reported case series and comprehensive review of the literature concluded that ASPPC lesions, like the two presented, tend to be yellow-white, flat, round or oval-shaped, and often involving the macula [2]. While detailed descriptions of additional imaging studies were not presented by van der Vaart and associates, we have found ancillary imaging to be quite useful in the diagnosis of ASPPC [2]. Fluorescein angiography, for example,

The authors have no financial interest in any aspect of this manuscript and have full control of all primary data.

C. M. Eandi (⊠)
Department of Surgical Science, Eye Clinic, University of Torino, Via Asti 17,
10131 Torino, Italy
e-mail: ceandi@gmail.com

E. Bertelli Department of Ophthalmology, Bolzano Central Hospital, Bolzano, Italy

E. T. Cunningham Jr. Department of Ophthalmology, California Pacific Medical Center, San Francisco, CA, USA

E. T. Cunningham Jr. Department of Ophthalmology, Stanford University School of Medicine, Stanford, CA, USA

E. T. Cunningham Jr. West Coast Retina Medical Group, San Francisco, CA, USA tends to show early hypofluorescence with progressive late hyperfluorescence, often with an irregular or 'leopard skin' pattern. Indocyanine green typically shows hypofluorescence corresponding to the macular lesion in both the early and late phases, although late hyperfluorescence can be observed as well. In the acute setting, Spectral Domain-Ocular Coherence Tomography (SD-OCT) tends to show some degree of disruption of the outer retinal hyper-reflective bands associated with the external limiting membrane, the photoreceptors, and/or the photoreceptorretinal pigment epithelium (RPE) junction. Irregular nodularity of the photoreceptor-RPE junction can also be observed. Lastly, enhanced depth imaging (EDI) OCT and B-scan ultrasonography show no thickening of the choroid or scleral in ASPPC, and effectively rule out choroiditis and scleritis as a cause of the lesion. All of these changes typically normalize completely following treatment with neurosyphilis doses of intravenous penicillin for 10 to 14 days. Given the diameter of the lesions presented by van der Vaart and colleagues, one would expect the choroid to be thickened if choroiditis were, in fact, the underlying cause. Perhaps this is what the authors intended when they wrote that the "granulomatous choroidal lesions were singular and rounded in appearance, with height noted also on OCT." A more detailed description of findings noted on imaging studies performed by the authors would be of interest.

It is noteworthy that both of the patients reported by van der Vaart and associates [1] did quite well following a 2week treatment with intravenous penicillin, despite the fact that one was infected by the human immunodeficiency virus (HIV; the status of the second patient was not reported). While a number of case reports and small clinic-based series have suggested that HIV co-infection can worsen the severity and/or outcome of ocular syphilis, our experience and acuity at last visit in 35 affected eyes in 23 HIV-positive patients to 58 affected eyes in 37 HIV-negative patients, and found no meaningful differences in either severity of clinical

presentation or vision outcome [2]. Similarly, Amaratunge and associate reviewed 41 original reports on syphilitic

uveitis in the English language literature published from

1984 to June, 2008, including 93 HIV-positive and 50

HIV-negative patients [4]. They found that only one of the

50 HIV-negative patients (2 %) had isolated anterior

or intermediate uveitis, compared to 27 of the 93 HIV-

positive patients (29 %; p=0.000023, Fisher's exact test).

Given that isolated anterior or intermediate uveitis tends to

be less likely to cause permanent vision loss than posterior

or panuveitis, this large retrospective review would seem to

similarly suggest that HIV co-infection alone does not

put patients at increased risk for a more severe, vision-

threatening uveitis at presentation. The notion that HIV co-

References

 van der Vaart R, Greven C, Manning R, Haines N, Kurup SK (2011) Unilateral solitary choroidal granuloma as presenting sign of secondary syphilis. Graefes Arch Clin Exp Ophthalmol 249:1575–1577

Graefes Arch Clin Exp Ophthalmol (2013) 251:2289-2290

- Eandi CM, Neri P, Adelman RA, Yannuzzi LA, Cunningham ET Jr, on Behalf of the International Syphilis Study Group (2012) Acute syphilitic posterior placoid chorioretinitis: report of a case series and comprehensivereview of the literature. Retina 32:1915–1941
- 3. Cunningham ET Jr, Eandi CM, Wender JD (2013) To the Editor (letter). Retina 33(2):451
- Amaratunge BC, Camuglia JE, Hall AJ (2010) Syphilitic uveitis: a review of clinical manifestations and treatment outcomes of syphilitic uveitis in human immunodeficiency virus-positive and negative patients. Clin Experiment Ophthalmol 38(1):68–74