Bilateral Acute Anterior Uveitis from Sinusitis Complicated by Optic Disc Oedema in a Child: A Case Report and Review of the Literature

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Abstract

Background

To describe an unusual case in which bilateral sinusitis was followed by bilateral anterior uveitis complicated by bilateral papillitis that completely resolved after appropriate treatment of the sinusitis.

Case presentation

A 11-year-old caucasian child presented with bilateral anterior uveitis and a diagnosis of poststreptococcal uveitis was made after a complete clinical examination. After 1 week course of topical steroids and systemic clarithromycin were then administered, leading to complete remission. After 2 months from diagnosis the situation recurred bilaterally. An encephalic Magnetic Resonance Imaging (MRI) showed bilateral maxillary sinusitis and both topical steroids and systemic benzylpenicillin a bilateral optic disc edema was also detected. The treatment was then continued, leading to complete remission. After 2 months from diagnosis the situation recurred bilaterally. An encephalic Magnetic Resonance Imaging (MRI) showed bilateral maxillary sinusitis and both topical steroids and systemic clarithromycin were then administered, leading again to bilateral remission. One month after the treatment had been stopped a monolateral recurrence occurred in his right eye and MRI showed persistent inflammation in the omolateral right maxillary sinus. Clarithromycin was then switched to systemic ciprofloxacin. A complete resolution of both uveitis and disc edema along with the improvement of the sinusitis occurred. Omolateral turbinate hypertrophy with nasal valve impairment was subsequently diagnosed and no recurrences occurred only after a surgical repair of the condition.

Results

Based on our report, sinusitis should always be considered in children with anterior uveitis and/or papillitis. A prompt and adequate antibiotic treatment is essential in order to avoid potentially severe complications.

Keywords: Anterior uveitis; Antibiotic therapy; Case report; Papillitis; Sinusitis

Abbreviations

AC: Anterior Chamber; OU: Both Eyes; MRI: Magnetic Resonance Imaging; ODE: Optic Disc disc Edema; LE: Left Eye; OCT: Optical Coherence Tomography; RE: Right Eye

Background

Anterior uveitis is 10 times less frequent in children than in adults but it can be much more severe and accounts for 30-40% of cases among children with intraocular inflammation [1-5]. Optic Disc Edema (ODE) in association with sudden-onset anterior uveitis is a relatively uncommon clinical finding [6]. Both disorders were detectable in our patient. Infections are a frequent situation among children, and sinusitis complicates upper respiratory tract infections in 5-10% of cases [7]. Uveitis originating from a sinusitis is a very infrequent event.

Case Presentation

A 11-year-old caucasian boy came to our attention complaining redness and photophobia in his Left Eye (LE) for a few days. Conjunctivitis was diagnosed and tobramycin + dexamethasone ophthalmic suspension 4 times daily was prescribed. After 10 days, the treated eye completely healed but similar signs and symptoms began in the fellow eye. Again, a non-specific conjunctivitis was diagnosed and diclofenac sodium ophthalmic solution 4 times daily was prescribed. The treatment was not effective and after 1 week, slit-lamp examination revealed bilateral circumlimbal injection, small inferior keratic precipitates and flare and cells within the Anterior Chamber (AC) which were quantified as 3+ in the Right Eye (RE) and 1+ in the LE, respectively, according to a six grade semi-quantitative scoring (0, trace, 1+, 2+, 3+ and 4+). RE pupil was miotic, with posterior synechiae at 3, 5 and 7 o’clock positions. Despite the significant intraocular inflammation, the visual acuity was 20/20 in both eyes (OU), with no blurring and no pain with eye movements. Posterior segment examination showed no vitreous cells but revealed mild congestion of retinal veins. Intraocular pressure was 12 mmHg OU. The ocular motility was normal.

A 4mg dose of parabulbar Betamethasone was immediately injected (RE) and Tobramycin + dexamethasone eye drops were started 5 times daily OU.

A thorough pediatric examination revealed no systemic abnormalities.

Several clinical investigations were performed: C-reactive protein, angiotensin-converting enzyme, plasma viscosity, antinuclear antibody, antineutrophil cytoplasmic antibody and chest X-ray, all

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of which proved normal. HLA-B27 was negative. Calcium, potassium, magnesium, phosphorus and creatinine were normal. Although a throat swab was negative, the patient had recurrent sore throats and blood testing revealed an elevated ASO titer (1150 IU/mL). Rheumatoid factor was negative and CRP was at the upper limit of the standard range. Post-streptococcal uveitis was diagnosed and following a rheumatological evaluation a treatment with benzathine Benzylpenicillin 1,200,000 IU once a week was started.

After 7 days the clinical picture was further complicated by the appearance of bilateral Optic Disc Edema (ODE), which was monitored by Optical Coherence Tomography (OCT) imaging. The visual field testing was normal OU.

The treatment was continued and the patient showed a significant improvement during the following days; three weeks after the acute event the situation completely resolved and steroid treatment was then tapered.

After 2 months from diagnosis the child had a recurrence of bilateral uveitis and ODE, despite the continuation of penicillin therapy. An encephalic Magnetic Resonance Imaging (MRI) was then performed. The exam (Figure 1a) showed normal globes and optic nerves bilaterally with no signs of orbital inflammation but revealed a maxillary sinusitis which was more severe on the right side, despite the absence of subjective symptoms like cheek pain or rhinorrhea. A topical corticosteroid (dexamethasone 0,1% four times daily) was restarted along with clarithromycin 250 mg BID for 7 days. Again, the uveitis promptly improved and the topical steroid treatment was gradually tapered in 4 weeks. One month after the treatment had been stopped, a monolateral relapse occurred in his RE, while LE showed no signs of active inflammation. The ODE was still detectable, although the OCT revealed a reduction in OU.

A further MRI of the maxillary sinuses was performed (Figure 1b), which revealed a persistent inflammation in the posterior portion of the right maxillary sinus. The presence of a residual inflammation in the right sinus was related to a microbial resistance to clarithromycin. It was then decided to change the antibiotic and ciprofloxacin 250 mg BID for 7 days was started along with the topical steroid.

After about 1 month, no signs of active inflammation were detectable and the OCT showed a further reduction of the ODE in OU and a bilateral resolution of the sinusitis was confirmed by a further MRI (Figure 1c). The OCT, which had improved during the administration of ciprofloxacin, completely disappeared after 3 weeks. The patient was completely asymptomatic during the following 6 years when he returned to our attention complaining again about redness, photophobia and sore throat. Optic disc swelling was still detectable, and ODE was still present in the right eye.

Nevertheless, the use of benzyl penicillin did not prevent relapse and MRI led us to suppose that the origin of the ocular inflammation was the bilateral asymptomatic maxillary sinusitis. The literature on uveitis as a complication of sinusitis is very scant and little is known about the mechanisms that lead anterior uveitis to cause ODE, which has been reported to occur in 7-29% of chronic uveitis but it is not reported as a complication of anterior chronic uveitis [4,11,12]. In our case, both cells and flare were very evident, but were not proportional to the severity of the disease. Indeed, the correlation between flare and ODE is uncertain, even though flare is normally associated with idiopathic, associated to systemic diseases or the result from a variety of infectious agents. In our case, the initial diagnosis of post-streptococcal syndrome uveitis was based on the bilateral involvement and non-granulomatous inflammation in the AC. Moreover, a familial predisposition to streptococcus infections was ascertained and, even though the boy did not show rheumatic fever, the anti-streptococcal lysin O titer was high and sore throat was recurrent. Optic disc swelling has also been described in the course of poststreptococcal syndrome uveitis [10].

**Discussion**

Uveitis is about 10-fold less common in children than in adults [2]. Young children frequently do not complain the typical symptoms nor show the physical findings seen in older patients [8]. From an epidemiological point of view, uveitis seems to be rarer in children than in adults, it is less common in girls than in boys and anterior uveitis occurs more frequently than other forms [9]. Uveitis may be idiopathic, associated to systemic diseases or the result from a variety of infectious agents. In our case, the initial diagnosis of post-streptococcal syndrome uveitis was based on the bilateral involvement and non-granulomatous inflammation in the AC. Moreover, a familial predisposition to streptococcus infections was ascertained and, even though the boy did not show rheumatic fever, the anti-streptococcal lysin O titer was high and sore throat was recurrent. Optic disc swelling has also been described in the course of poststreptococcal syndrome uveitis [10].

![Figure 1](image-url)
loss resulting from ischemic or edematous retina [15]. Concerning our case we suppose the following mechanism: the sinusitis induced severe inflammation causing a massive release of proinflammatory cytokines that reached the orbit and involved the pre-laminar portion of the optic nerve. Common complications of ocular inflammation, such as retinal edema, may be due to the effects of cytokines [16], which could have produced flare in the AC and a partial axoplasmic block secondary to intra-axonal edema in the optic nerve. Recent studies on the role of cytokines in uveitis support our hypothesis [17]. Conversely, a rise in cerebrospinal fluid pressure in the cranial subarachnoid space and its transmission to the sheath of the optic nerve was not essential in our case. Indeed, when the inflammation of the sinus improved, both eyes completely healed and no more relapses occurred after the condition predisposing to sinusitis was finally resolved.

It should be emphasized that ODE due to anterior uveitis appears to have a more favorable prognosis than post-sinusitis papillitis which may lead to severe visual complications [6,18,19] and this can explain the good visual outcome of our patient.

Finally, our report highlights the importance of antibiotic treatment in order to shorten the duration of infection and illness, to reduce mucosal damage and to prevent contiguous infectious involvement of the orbit. In our patients we used clarithromycin and ciprofloxacin. The former was used because it probably exerts its beneficial effect not only by inhibiting or killing bacterial pathogens, but also by down-regulating pro-inflammatory mechanisms. The latter antibiotic was used in order to destroy any remaining clarithromycin-resistant bacteria. Quinolone-induced arthropathic toxicity has largely restricted the use of ciprofloxacin in children and adolescents. It is now known, however, that this drug does not engender an important risk of arthropathy or tendinopathy in humans [20]; indeed, the treatment has been effective and devoid of side effects in our case.

Conclusion

In conclusion, our case report suggests that a chronic sinusitis should always be considered the differential diagnosis of in children with anterior uveitis and/or papilledema. Prompt and appropriate antibiotic treatment is essential in order to avoid potentially severe complications and to obtain a complete healing.

Authors’ contributions

SCS conception and data collection; AB and CET: article revision; all authors read and approved the final manuscript

References