



POSTER PRESENTATION

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PreS-FINAL-2282: Amaurosis as a presenting sign of antiphospholipid syndrome secondary to systemic lupus erythematosus - case report

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Introduction

Antiphospholipid syndrome (APS) secondary to systemic lupus erythematosus (SLE) can be recognised in children with arterial or venous thrombosis. Amaurosis due to thrombosis of central retinal vein is rarely presenting manifestation of SLE with secondary APS.

Objectives

To present APS secondary to SLE with aggressive ophthalmological onset in 17 years old female.

Methods

We report a patient with unilateral amaurosis due to thrombosis of central retinal vein. Amaurosis was a reason for her urgent admission at Ophthalmology. She was transferred to Pediatric rheumatology department as suspected SLE. The patient had rapidly developing disease. Eleven days after the attack of retinal vein thrombosis, she became febrile with malar rash, facial ulcer, neurological symptoms (right Mingazzini positive), arterial hypertension, haematological abnormalities, proteinuria and immunological disorders. Head MRI-MRA was performed and subocclusion of left medial cerebral artery was found. The diagnosis of APS secondary to SLE was established.

Results

The patient significantly improved with aggressive immunosuppressive and prompt anticoagulant therapy but ophthalmological complication have been improved slowly with uncertain prognosis.

Conclusion

The patients with SLE related symptoms have to be referred to rheumatologist immediately because APS secondary to SLE may have aggressive thrombotic onset and cause serious organs damages.

Disclosure of interest

None declared.

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