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## Syphilitic optic neuritis and uveitis – case report

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### ABSTRACT

We report the case of a 34-year-old male patient who presented to the Ophthalmology Department because of a decline in visual acuity in both eyes. The patient was diagnosed with anterior and posterior uveitis in the right eye and optic neuritis in the left eye, secondary to syphilis. Topical and systemic treatments were

implemented, leading to a gradual improvement in the patient's condition. Considering the rising global incidence of syphilis and the non-specific ocular manifestations it may present, the diagnosis should be considered in patients with ocular inflammatory conditions of unknown etiology.

**KEY WORDS:** syphilis, syphilitic uveitis, syphilitic optic neuritis.

### INTRODUCTION

Syphilis is a sexually transmitted infectious disease caused by the bacterium *Treponema pallidum*. The primary modes of transmission include sexual contact with an infected person and vertical transmission. Syphilis is a chronic multisystem disease characterized by a wide spectrum of symptoms and following a cyclical course with alternating periods of exacerbation and latency. Involvement of the nervous system, including optic neuritis and other ocular manifestations, can occur at any stage of syphilis infection.

### CASE REPORT

A 34-year-old male patient was referred to our Department due to uveitis in the right eye and optic neuritis in the left eye. The patient denied any comorbidities and chronic medication use.

Based on the physical examination and the medical records provided, it was determined that the onset of ophthalmic symptoms had occurred four months earlier. At that time, the patient was hospitalized at another ophthalmology center because of painless deterioration of vision in the right eye to the level of hand movement in front of the eye, which had persisted for two days. The patient reported no complaints in the left eye. Despite conducting examinations, the underlying cause of progressive vision loss in the right eye remained unidentified. Intravenous methylprednisolone treatment was initiated.

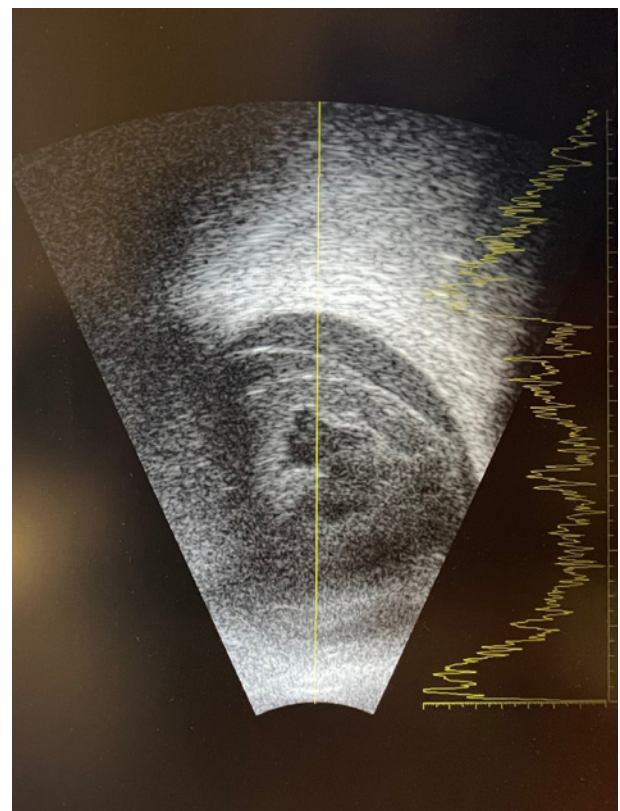


Figure 1. Ultrasound findings in the right eye

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Figure 2. Color fundus photograph of the left eye

On the day of admission to our Department, the patient's physical examination revealed loss of light perception in the right eye. The visual acuity in the left eye was 0.8. The intraocular pressure was 23 mmHg and 18 mmHg, respectively. Slit-lamp examination revealed, in the right eye, deposits on the corneal endothelium, single inflammatory cells in the anterior chamber, a non-reactive, unrounded pupil with posterior synechiae, and no fundus view. Ultrasound examination showed the presence of multiple densities in the vitreous and thickening of the choroid (Figure 1).

In the left eye, no pathologies were found in the anterior segment, but optic disc edema was observed (Figure 2). In addition, the patient's optic nerve was assessed by spectral optical coherence tomography (SOCT) (Figure 3). SOCT scan of the left macula was within the normal range (Figure 4).

Fluorescein angiography showed multiple foci of diffuse hyperfluorescence with signs of late-phase leakage (Figure 5 A-D).

In the biochemistry panel, a number of abnormalities were found, including ESR (30 mm/h), doubtful IgM antibody test and positive IgG antibody test for *Borrelia burgdorferii* and HSV, and reactive WR-RPR result. Western blot assay for Lyme disease along with TPHA and FTA-ABS tests for syphilis were ordered for further verification.

The results of other laboratory tests (Quantiferon-TB, ANA, ANCA, RF, antibodies to *Toxoplasma gondii*, *Toxocara* spp., *Bartonella henselae*, HIV, HCV, HBV) were negative.

Contrast-enhanced MRI of the brain revealed an incidental finding – an arachnoid cyst in the right posterior cranial fossa, measuring 76 × 54 × 33 mm, deforming the right cerebellar hemisphere. Lung X-ray and abdominal ultrasound findings were normal. Neurological consultation revealed bilaterally positive Sterling's and Rossolimo's signs, and Babinski's sign tendency on the left side.

During the patient's stay in the Department, topical agents (dexamethasone drops, mydriatics) were applied to both eyes. In addition, the patient received intravenous steroid therapy (methylprednisolone at a dose of 1 g/day for a total of 5 days). A decrease in visual acuity to 0.5 was noted in the left eye.

After one week, on the day of discharge, the best-corrected visual acuity in the left eye was 0.8. The patient's intraocular pressure was within the normal range. Physical examination revealed a raised optic disc with blurred margins (Figures 6, 7). Steroid therapy was maintained in the oral form (prednisone).

After receiving the examination results confirming the diagnosis of syphilis, oral antibiotic therapy (doxycycline 200 mg) was initiated. The patient was referred to the Neurology Outpatient Clinic to determine eligibility for lumbar puncture and potential continuation of treatment following the protocol for neurosyphilis. In addition, the patient was advised to schedule consultations at the Infectious Diseases Outpatient Clinic and the Dermatology Outpatient Clinic.

The patient remains under the care of ophthalmology and neurology specialists. The last examination, six months after hospitalization in the Ophthalmology Department, showed no sense of light in the right eye and visual acuity of the left eye at 1.0. Physical examination of the right eye revealed a clear cornea, clean anterior chamber, unrounded pupil, posterior synechiae, and pigment deposits on the lens. No abnormalities are found in the anterior segment and fundus of the left eye (Figure 8). The findings of the follow-up SOCT scan of the optic nerve were normal (Figure 9).

## DISCUSSION

The incidence of syphilis worldwide is rising, especially among high-risk populations. Those most at risk include homosexual men and HIV-positive individuals [1, 2]. Ocular manifestations are rare, found in only 1-5% of neurosyphilis cases [3]. They can occur at any stage of the disease, but are most common (4.6%) in secondary syphilis [4]. Approximately 75% of patients are male [2]. The mean age of onset of ocular symptoms is 48.6 years [5].

Syphilis can involve all ocular structures, but the most common manifestation is uveitis. Panuveitis occurs in 41% of patients with ocular syphilis, while optic nerve is involved in 22% [6]. The most common symptoms reported by patients include reduced visual acuity, ocular pain and headache, photophobia, red eyes, and increased lacrimation. In some cases, the disease may progress asymptotically [3, 7-10]. Possible complications include the development of cataract, glaucoma, epiretinal membrane, optic nerve atrophy, or retinal detachment [8].

The first-line drug in the treatment of syphilis is crystalline penicillin at a dose of 18-24 million units/day for 10-14 days. An alternative is oral doxycycline at a dose of 200 mg twice a day for two to four weeks [1, 4].





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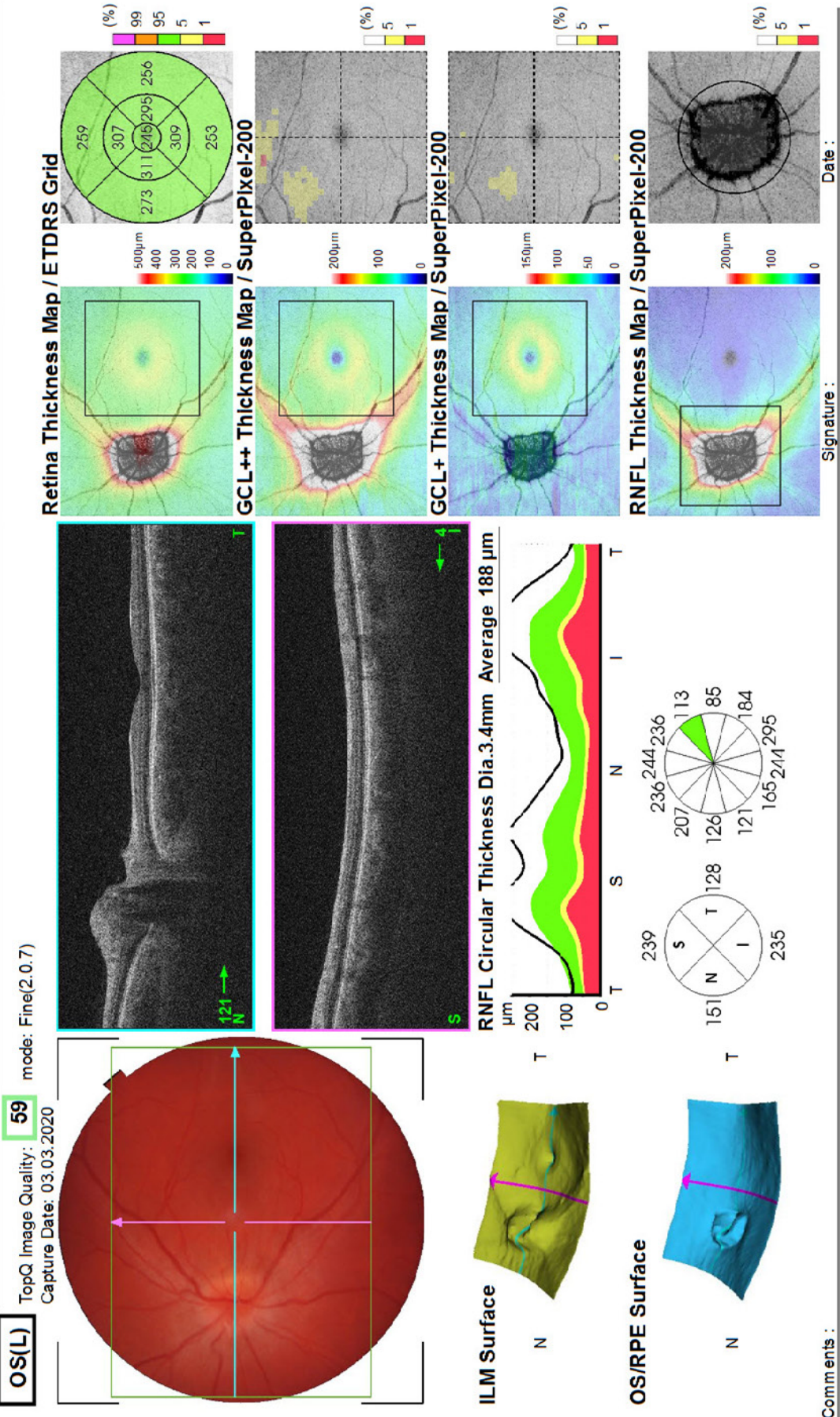


Figure 3. SOCT scan of the left optic nerve

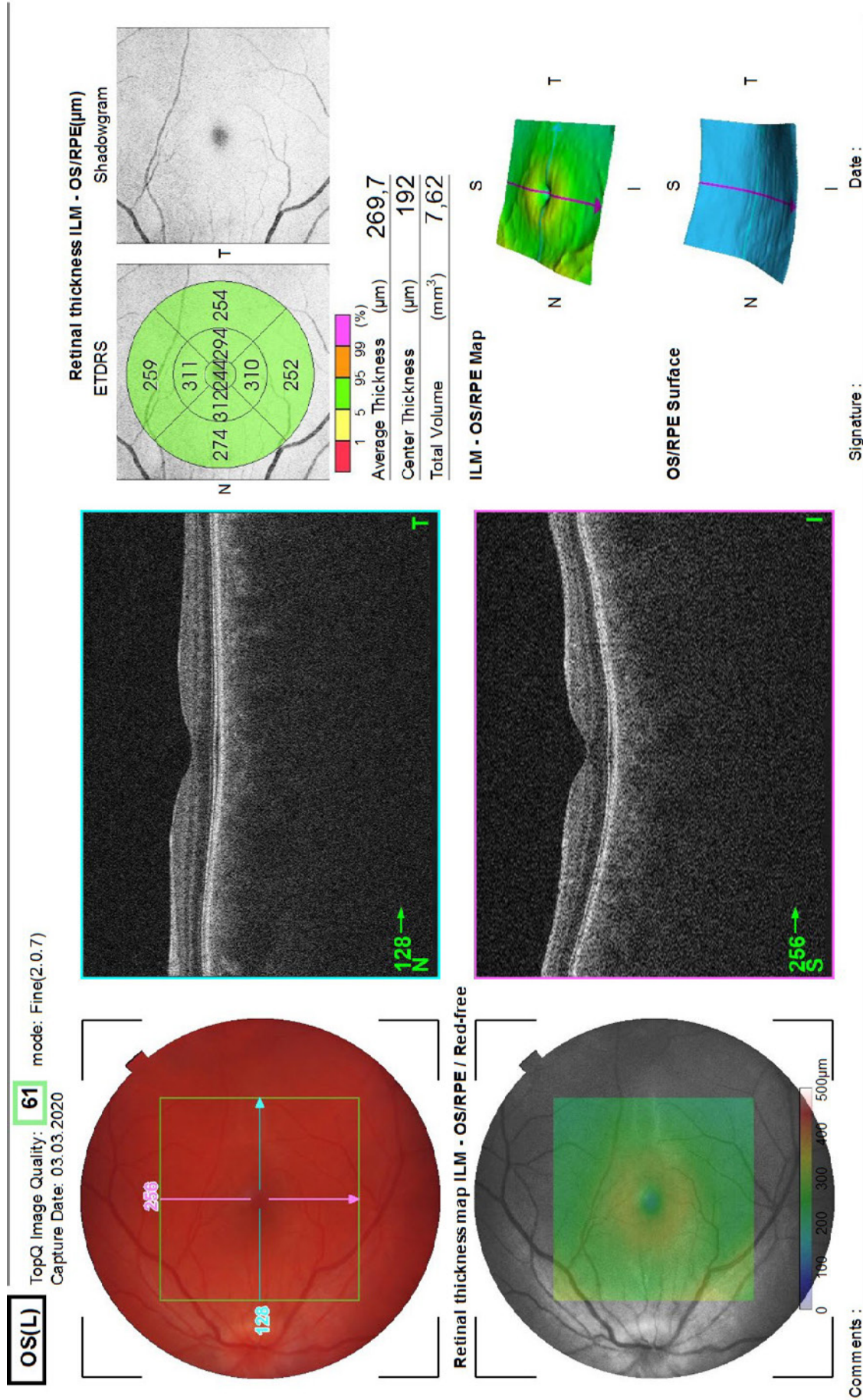


Figure 4. SOCT scan of the left macula



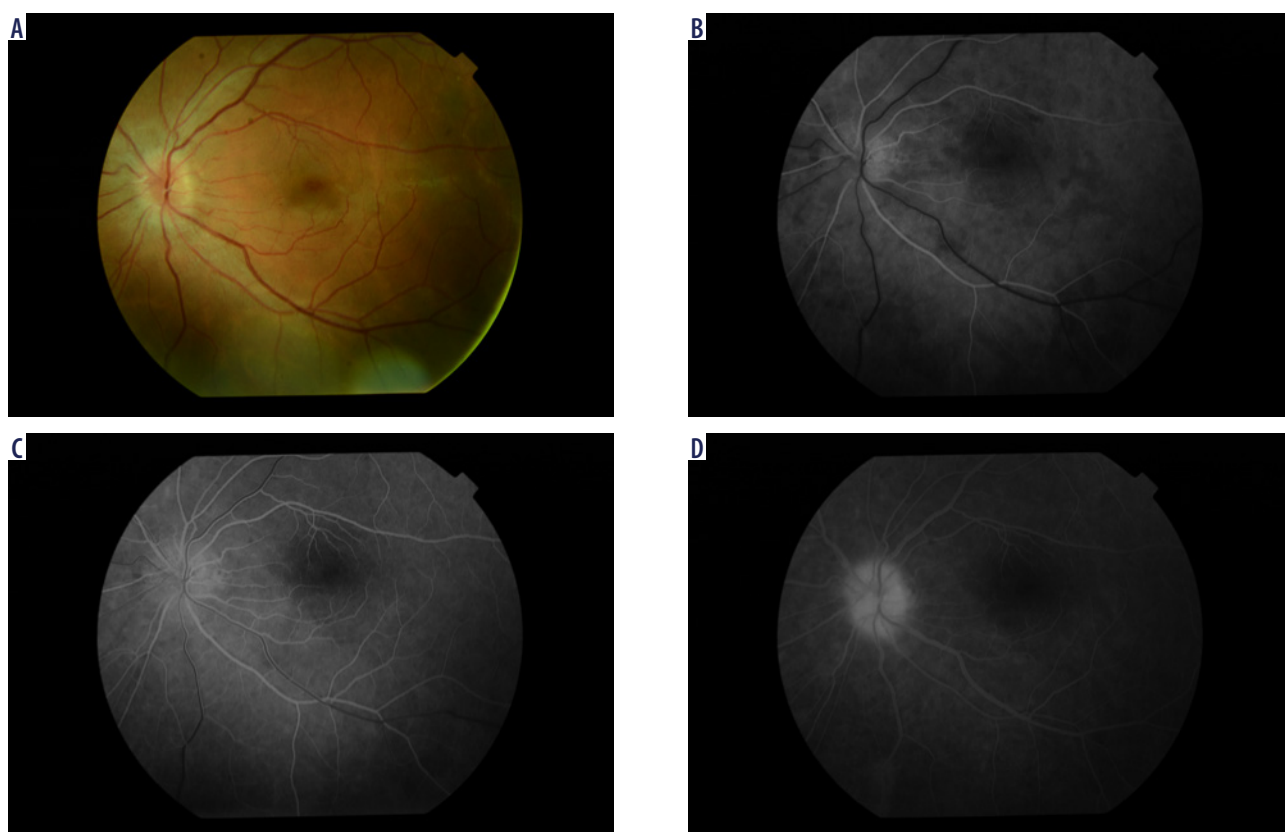


Fig. 5 A-D. Fluorescein angiography findings



Figure 6. Color fundus photograph of the left eye



Print Date : 24.03.2020

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Technician :

Name : kamil blelecki

Gender : Male

Fixation : OS(L) Wide

OS(L)

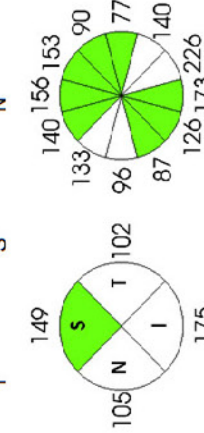
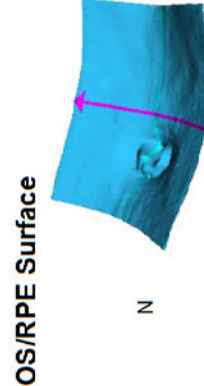
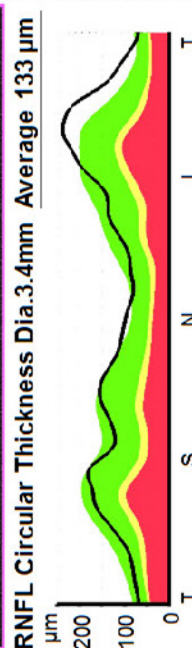
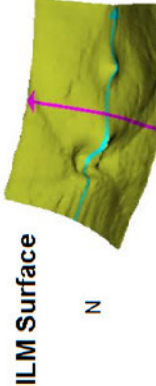
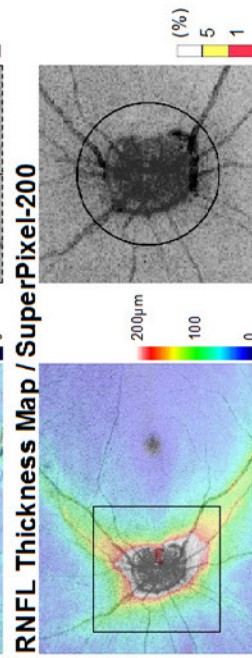
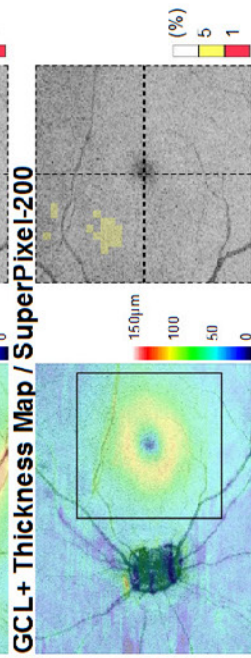
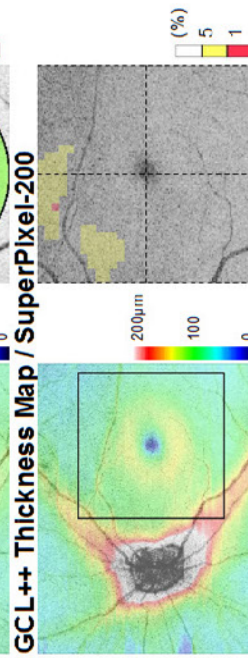
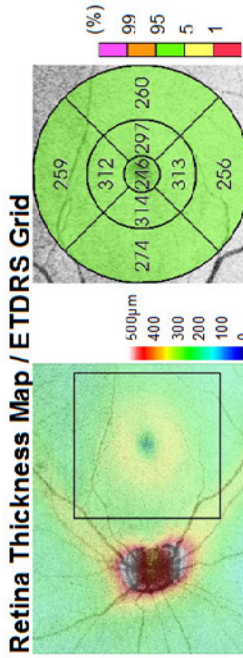
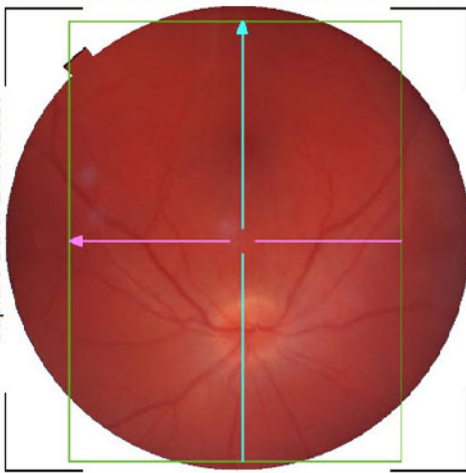
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Figure 7. SOCT scan of the left optic nerve



Figure 8. Color fundus photograph of the left eye

## CONCLUSIONS

Given the increased incidence of syphilis in recent years and the fact that ocular manifestations of the condition can mimic disease entities of various etiologies, it is important to include syphilis in the differential diagnostic process. In view of the broad range of symptoms, potential multiple

organ involvement and associated diagnostic challenges, syphilis remains a disease requiring close collaboration of medical professionals of various specialties.

## DISCLOSURE

The authors declare no conflict of interest.





Print Date : 30.06.2020

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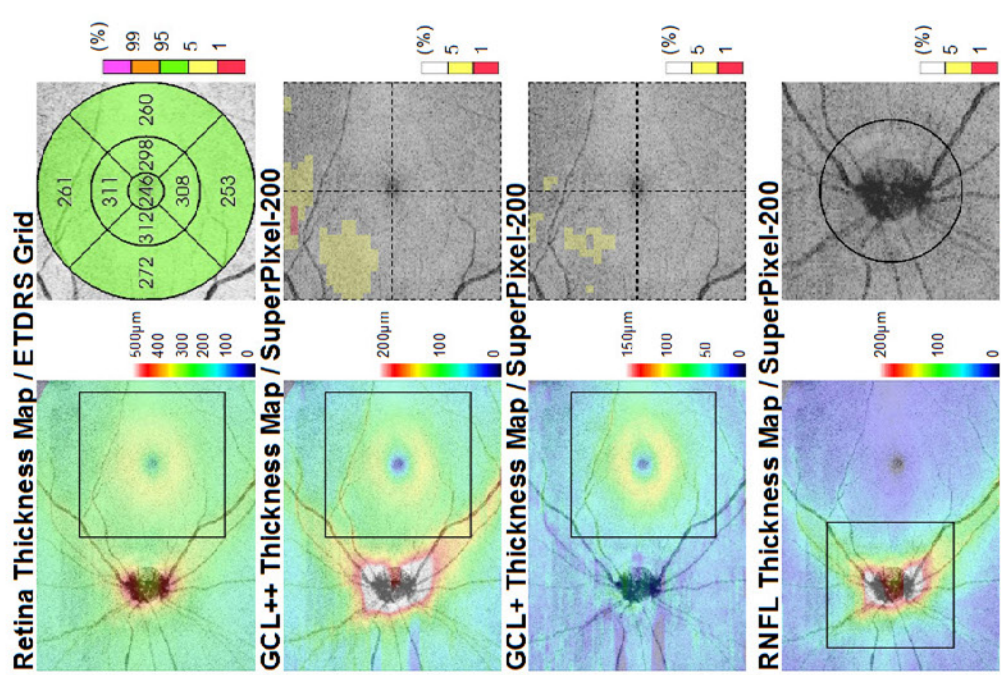
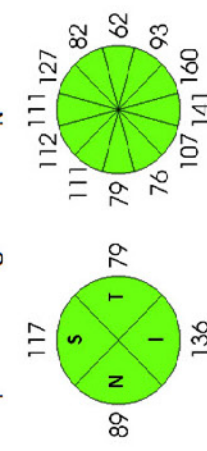
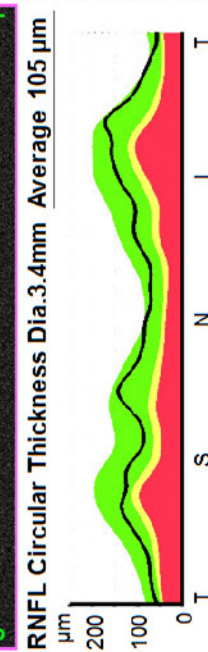
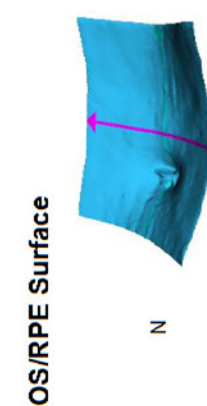
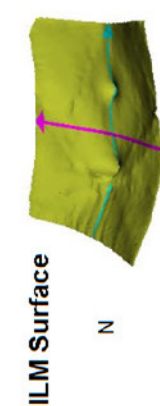
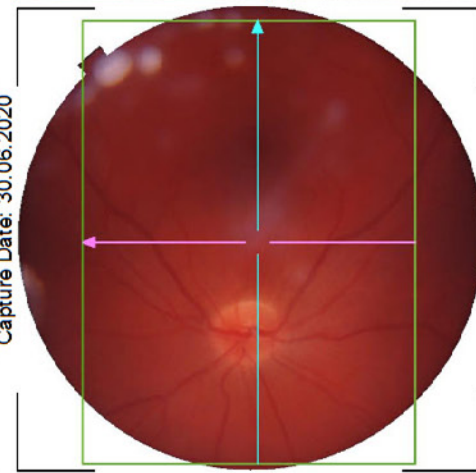
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Figure 9. Results of SOCT scan of the optic nerve

## References

1. Zajkowska J, Drozdowski W, Grygorczuk S. Kiła ośrodkowego układu nerwowego – trudności diagnostyczne. *Neurologia po Dyplomie* 2014; 9: 26-39.
2. Furtado JM, Arantes TE, Nascimento H, et al. Clinical Manifestations and Ophthalmic Outcomes of Ocular Syphilis at a Time of Re-Emergence of the Systemic Infection. *Sci Rep* 2018; 8: 12071.
3. Koripalli S, Rueda Prada L, Gummadi P, et al. A Rare Case of Neurosyphilis with Ocular Involvement in a Patient with HIV Infection and New Onset Syphilis. *Cureus* 2019; 11: e4034.
4. Kuo A, Ziaee SM, Hosseini H, et al. The Great Imitator: Ocular Syphilis Presenting as Posterior Uveitis. *Am J Case Rep* 2015; 16: 434-437.
5. Klein A, Fischer N, Goldstein M, et al. The great imitator on the rise: ocular and optic nerve manifestations in patients with newly diagnosed syphilis. *Acta Ophthalmol* 2019; 97: e641-e647.
6. Mathew RG, Goh BT, Westcott MC. British Ocular Syphilis Study (BOSS): 2-year national surveillance study of intraocular inflammation secondary to ocular syphilis. *Invest Ophthalmol Vis Sci* 2014; 55: 5394-5400.
7. Macovei ML, Georgescu RD. Papillitis in Neurosyphilis. *Rom J Ophthalmol* 2019; 63: 406-411.
8. Etheridge T, Bowen RC, Raven M, et al. Ocular Syphilis: Clinical Manifestations and Treatment Course. *WMJ* 2019; 118: 191-195.
9. Chen JJ, Bhatti MT, Bradley E, et al. Incipient Syphilitic Papillitis. *Neuroophthalmology* 2019; 44: 11-15.
10. Lutchman C, Weisbrod DJ, Schwartz CE. Diagnosis and management of syphilis after unique ocular presentation. *Can Fam Physician* 2011; 57: 896-899.