An Early Presentation of Neurosyphilis Manifesting as Cranial Polyneuropathies: A Case Report

Abinayaa P. Ravichandran, Katie Reming, Keithan Sivakumar, Hussam A. Yacoub

Abstract

Neurosyphilis (NS) is a rarely reported disease, with neurological manifestations occurring in the late stages of the infection. The incidence of NS has steadily increased, particularly with increased cases of human immunodeficiency virus (HIV) infection among homosexual partners, and can occur at any stage of the disease. In this case report, we present a 53-year-old homosexual man with HIV who presented with a gradual onset of vertigo over a course of 2 weeks, followed by gradual onset of multiple cranial neuropathies. Magnetic resonance imaging of the brain revealed pathological enhancement of multiple cranial nerves. Laboratory workup revealed reactive Venereal Disease Research Laboratory dilutional titer in the cerebrospinal fluid, supporting the diagnosis of NS. He was treated with intravenous penicillin with gradual resolution of symptoms. Our case illustrates that NS can occur with no known prior evidence or manifestation of the infection. The diagnosis of NS should be entertained in individuals with a known history of HIV and unexplained multiple cranial neuropathies.

Keywords: Syphilis; Neurosyphilis; Cranial neuropathies

Introduction

In the modern era, neurosyphilis (NS) is often a challenging diagnosis due to its rarity and diverse presentation. There has been a steady reemergence in the incidence of NS over the past two decades, particularly with the increase in prevalence of human immunodeficiency virus (HIV) infection among homosexual partners. This case report describes an uncommon presentation of NS in an immunocompetent HIV-positive homosexual man who presented with multiple cranial neuropathies and headache.

Case Report

A 53-year-old homosexual man presented with a gradual onset of vertigo over a course of 2 weeks along with unilateral right-sided ear pain, fullness and hearing loss. He also experienced right-sided intermittent, throbbing headache and post-nasal drip. The patient had a known history of HIV treated with antiretroviral therapy (ART), elvitegravir-cobicistat-emtricitabine-tenofovir, with an undetectable viral load and a CD4 count of 847/mm³. Other comorbidities included hyperlipidemia, hypertension, pre-diabetes, factor-V Leiden mutation and a history of deep vein thrombosis. The patient denied recent infections, fever, visual disturbance, neck pain/stiffness, or rash. Evaluation by otorhinolaryngology revealed minimal fluid in the middle ear compartment bilaterally on otoscopic examination. Audiometric evaluation revealed bilateral sensorineural hearing loss, with the right ear worse than the left by 10 decibels. Treatment was initiated with an oral prednisone taper, intranasal fluticasone and fexofenadine with minimal improvement of symptoms.

Three weeks later, the patient presented to the emergency department with right-sided facial paralysis and paresthesia. Neurological examination revealed right-sided wide palpebral fissure and inability to close the eyelid, flattening of the nasolabial angle, and decreased sensation to light touch in the distribution of the maxillary and mandibular trigeminal nerve branches. The remainder of the examination was unrevealing. Magnetic resonance imaging (MRI) of the brain revealed abnormal enhancement of the right facial nerve (Fig. 1) and subtle enhancement of the left facial nerve at the fundus of the left internal auditory canal (Fig. 2), with no cerebellopontine angle or intracranial masses, acute infarction, or hydrocephalus. Serology testing was negative for Lyme antibodies. Syphilis screening was not performed at this time. A presumed diagnosis of Bell's palsy was made, and the facial paralysis significantly improved. The patient continued to experience facial hypoesthesia, hearing loss, vertigo and intermittent headaches.

Over the following 4 weeks, the patient continued to have worsening headaches associated with occasional vomiting. He was treated with a tapering course of prednisone along with...
oral acyclovir. Two days later, he experienced painless loss of peripheral vision. On ophthalmological evaluation, visual acuity was 20/50 on the right, with an inferior temporal quadrant visual field deficit, loss of color vision and an afferent pupillary defect. On fundoscopic examination, the patient was found to have severe edema of the right optic nerve with evidence of hemorrhage at the margin. The remaining neurological examination was unremarkable.

Blood work revealed a white blood cell count (WBC) of 9.1 × 10⁹/L and CD4 T cell count of 847/mm³, with an undetectable HIV viral load. Autoimmune testing was positive for antinuclear antibodies, with a titer of 1:80, and a rheumatoid factor titer of 1:211. Extensive immunological workup including anti-cytoplasmic antibody, double stranded DNA antibody, anti-Ro, anti-La, Smith auto-antibody, SCL 70 antibody and myeloperoxidase antibody was unrevealing. Serological testing was reactive for rapid plasma reagin (RPR), with titers of 1:256. West Nile virus, viral hepatitis and a repeat Lyme assay were negative. Cerebrospinal fluid (CSF) analysis revealed a clear, colorless fluid with an opening pressure of 17 cm H₂O. Protein was elevated at 84 mg/dL. CSF red blood cell count was 0, and WBC count was elevated at 104/mm³ with a lymphocytic predominance of 96%. Venereal Disease Research Laboratory (VDRL)-CSF dilutional titer was reactive at 1:4, suggestive of NS. Complete meningocencephalitis panel was unremarkable. MRI of the brain revealed expanded abnormal enhancements involving multiple cranial nerves, including the trigeminal nerve bilaterally (Fig. 3), the fundal and labyrin-

thine portions of the right facial nerve, and geniculate ganglion. Subtle enhancement was also seen at the fundus of the left internal auditory canal and right side of the optic chiasm (Fig. 4).

Treatment was initiated with IV penicillin G, 4 million units every 4 h for 14 days. Within 24 h, the patient developed nausea, sinus tachycardia and a low grade fever. A Jarisch-Herxheimer reaction was suspected and symptoms resolved with the use of acetaminophen and ondansetron. The patient was discharged on hospital day 5 with home infusion for the remainder of the antibiotic treatment. The facial paralysis completely resolved, with residual facial hypoesthesia, hearing loss and vision loss.

Over the course of several days, the patient reported subjective improvement of vision in the right eye by approximately 30%. Ophthalmological evaluation revealed a visual acuity of 20/50 in the right eye with an afferent pupillary defect, absence of color vision and pale optic nerve on fundoscopic examination. Evaluation of the left eye was unrevealing. Ocular coherence tomography demonstrated damage to the peripheral retinal nerve fiber layer sparing the central vision. Hearing and facial weakness completely resolved with no residual deficits.

**Discussion**

Historically, NS has been described as a late manifestation of
the disease, occurring decades after the primary infection as tabes dorsalis and general paresis. NS is rarely reported in the United States with the availability and effectiveness of penicillin. The incidence of primary and secondary syphilis in the United States has increased substantially over the past two decades from 2.12 to 10.8 per 100,000, with men who have sex with men (MSM) being the most at risk population [1, 2].

Treponema pallidum (T. pallidum), the spirochete bacterium that causes syphilis, can invade the central nervous system (CNS) at any stage of the infection [3, 4]. The majority of individuals remain asymptomatic, with evidence of bacterium eradication without any pharmacotherapy [4]. Less than 2% of patients develop NS, with symptoms occurring months to decades after the initial infection [5].

Aseptic meningitis is one of the earliest manifestations of NS, with a prototypical constellation of symptoms including headache, photophobia and neck stiffness. This can progress to ocular syphilis and cranial neuropathies [3]. In a case series published by the Center for Disease Control, approximately 75% of patients had visual symptoms, ocular manifestation and headache, whereas only 6% and 12% had cranial neuropathies and acute meningitis, respectively [6]. Vascular complications such as stroke and hemorrhage have been reported in approximately 4% of patients [6, 7]. Neurological manifestations of syphilis are more than twice as likely to occur in cases of HIV co-infection, with the prevalence of NS being 2.1% in HIV-positive versus 0.6% in HIV-negative patients [8].

is also evidence of a positive correlation between the degree of immunosuppression and the likelihood of developing NS [3].

NS may present earlier and occur more frequently in patients with HIV [8, 9], and often has a fulminant course with overlapping disease stages compared to patients without an HIV co-infection [10]. Our patient presented with a headache and multiple cranial neuropathies, in the absence of fever or other signs of meningeal irritation. He did not have any symptoms suggestive of primary or secondary syphilis, supporting the notion that CNS infection with T. pallidum can occur at any stage. A prospective cohort study reported that syphilis could adversely affect the HIV viral load and CD4 count even in patients receiving ARVT, therefore increasing the risk of transmission [11].

A study in Los Angeles reported a 2.1% increase in the incidence of NS among individuals infected with HIV compared to 0.6% in the non-HIV cohort between 2001 and 2004. The authors speculated a greater risk of developing NS with serum RPR titers > 1:32 and/or CD4 count < 350/mm³ [8]. Another study by Ghanem et al in 2008 reported that a CD4 < 350/mm³ is a strong predictor of developing both symptomatic and asymptomatic NS. The CD4 count in this cohort ranged between 72 and 345/mm³ in individuals with NS [12].

Our patient’s HIV infection was well controlled with ARVT, indicated by an undetectable viral load and a CD4 T-cell count of 847/mm³. Although Ghanem et al reported a synergistic relationship with increased risk of NS in HIV patients with a
low CD4 count [12], our patient developed NS despite a non-immunocompromised state.

The increased susceptibility to NS in individuals with HIV may not be predominantly proportional to the degree of immunosuppression [3]. This suggests that being immunosuppressed rather than immunocompromised may have played a role in our patient’s increased susceptibility to acquiring NS. Successful treatment with anti-retroviral therapy has been posited to effectively downregulate both the innate and acquired immune system’s ability to respond to *T. pallidum* [13]. This mechanism may help explain the relatively rapid progression to NS in a previously asymptomatic patient with a well-controlled HIV infection.

Diagnostic workup for NS includes serological testing, CSF analysis and concomitant neuroimaging. Patients with reactive non-treponemal tests should undergo confirmatory treponemal assays as false positives to the former have been reported in many HIV patients. CSF VDRL is highly specific but not sensitive. A positive CSF VDRL in the setting of clinical signs and symptoms is diagnostic of NS. If VDRL is negative, CSF fluorescent treponemal antibody absorbed (FTA-ABS) testing coupled with serology and clinical signs are required to establish the diagnosis [14]. MRI of the brain often reveals enhancing or non-enhancing cortical lesions, a pattern suggestive of encephalitis, with or without adjacent meningeal enhancement and cerebral infarctions [15]. The varied clinical presentation in our patient prompted repeat imaging, which revealed interval progression and multiple cranial nerve enhancements. Magnetic resonance angiography and spinal cord imaging are warranted if there are signs of vasculitis and posterior column involvement, respectively. Alternative diagnoses of conditions with overlapping imaging features including viral or other bacterial infections, demyelination, or inflammatory diseases should be entertained.

Parenteral penicillin G is the treatment of choice for NS. The dosage, duration and formulation vary among the different stages and systemic manifestations. Jarisch-Herxheimer reaction, an allergic febrile response to treatment, occurs most frequently in the early stages of NS, presumably due to extensive bacterial burden. NS patients with underlying HIV are treated similarly to those without the co-infection. Follow-up with repeat serologies at 6 and 12 months is recommended in individuals with clinical resolution of symptoms [14].

Acknowledgments
None to declare.

Financial Disclosure
None to declare.

Conflict of Interest
None to declare.

Informed Consent
Informed consent was obtained from the patient verbally and there are no identifying features in the paper.

Author Contributions
All authors contributed equally to the production of this manuscript. The entire manuscript was reviewed and edited by all individuals listed.

Data Availability
The authors declare that data supporting the findings of this study are available within the article. All data were extrapolated from referenced studies and those studies may be reviewed for specific details pertaining to their data collection.

References
6. Symptomatic early neurosyphilis among HIV-positive men who have sex with men, four cities, United States, January 2002-June 2004. Centers for Disease Control and
Prevention. Atlanta, GA.


