Peripheral nerve involvement is a frequent complication of Human Immunodeficiency Virus (HIV) infection. Among cranial nerves, the seventh is the most commonly affected. However, facial diplegia is a rare clinical entity in the HIV infected population. We recently observed a case that is of particular interest, since the patient developed facial diplegia and unilateral vestibular neuritis revealing HIV infection.

**Case Report**

The patient is a 60-year-old heterosexual male with a past medical history of morbid obesity and chronic obstructive pulmonary disease. He had been in his usual state of health until approximately four weeks prior to admission, when he developed low grade fever, rhinorrhea, sore throat and diffuse arthralgias. Fever resolved promptly after treatment with amoxicillin and clavulanic acid. Later on he was admitted to our institution with a five-day history of constant dizziness, unsteadiness and imbalance. By the time of admission he additionally experienced perioral numbness followed suddenly by lack of facial movements, with failure of both eyes to close and difficulty in sucking, smiling and talking. Neurological examination revealed bilateral peripheral facial palsy, right-deviation tendency in past pointing and Quix tests, and mild unsteadiness with gait deviation to the right. The remainder of physical and neurological examination was normal, including intact reflexes. Laboratory studies disclosed an erythrocyte sedimentation rate of 51 mm/h. White blood cell count was 12.4 x 10^9 cells/l, with 35.6% neutrophils, 48.6% lymphocytes and 12.3% monocytes. Red blood cell and platelet counts were within normal limits. Autoantibody screening tests and blood biochemistry parameters, including angiotensin converting enzyme, were also normal. Cerebrospinal fluid (CSF) contained 30 cells/µl with 93% lymphocytes, protein 1.64 g/l and glucose 2.6 mmol/l (serum glucose 6.7 mmol/l). Cytology was negative. No organism was seen on Gram-stained smear, and cultures of CSF for bacterial, mycobacterial, viral and fungal organisms were also negative. Polymerase chain reaction and serologies for citomegalovirus, varicella zoster virus, Epstein-Barr virus, type 1 and 2 herpes simplex virus, Toxoplasma gondii, Cryptococcus, Brucella, Treponema pallidum, and Borrelia burgdorferi were negative in both serum and CSF. Human Immunodeficiency Virus testing was positive by ELISA and was confirmed by Western blot. CD4+ cell count was 252 cells/µl and CD8+ cell count was 5027 cells/µl, with CD4/CD8 ratio at 0.05. Viral RNA load was 197 000 copies/ml. A gadolinium enhanced brain magnetic resonance imaging showed normal findings. Electrophysiological study confirmed severe impairment of trigemino facial reflex, with no motor evoked potentials recorded from the orbicularis oculi muscle of both sides after stimulating the supraorbital nerve. No evidence of a generalized demyelinating polyneuropathy was found on nerve conduction studies. Since there was no evidence of any other underlying systemic illness, the diagnosis of right vestibular neuritis and facial diplegia in relation to HIV acute seroconversion syndrome was made. After five days unsteadiness and facial palsy steadily improved, with complete resolution of neurological signs within the following week. At three-months follow-up the patient remained free of all neurological features.

**Discussion**

Bilateral facial palsy, defined as the appearance of paresis of the contralateral facial nerve within 30 days of the onset of the first side, is due to a systemic cause in a much higher incidence than unilateral palsy. The diseases most commonly associated with facial diplegia are Guillain-Barre syndrome, multiple ideopathic cranial neuropathies, brainstem encephalitis, Melkerson-Rosenthal syndrome, syphilis, sarcoidosis, Lyme disease, bacterial meningitis, poliomylitis, Epstein-Barr Virus and other herpesvirus infections, leukemia, meningeal carcinomatosis, pontine tumours, Moebius syndrome, ideopathic intracranial hypertension and, more rarely, HIV infection. Although both unilateral and bilateral facial paralyses occur with a greater frequency in HIV infected patients than in the general population, bilateral palsy is exceedingly rare even among them. Seventh nerve involvement is the most frequent cranial neuropathy associated with HIV. As in our patient, it mainly occurs in the early stages of the disease, in the setting of lymphocytic pleocytosis and antibodies directed against HIV in CSF, supporting the hypothesis of a direct lesion of the facial nerve by the neurotropic virus; however, an immunologically mediated inflammation of the cranial nerves has also been suggested as a possible pathophysiologic mechanism, given the similarities with other demyelinating peripheral neuropathies associated with HIV seroconversion. Other causes of facial palsy such as Ramsay-Hunt syndrome, syphilis, type-2 herpes...
simplex infection, mononeuritis multiplex, and lymphomatous or sarcomatous infiltration of the nerve, must be considered in the later stages6,7. Our case is original in that facial palsy appeared and resolved simultaneously in both sides, in a complete symmetrical way. Moreover, its association to vestibular neuritis revealing HIV infection is a very rare clinical picture, which could have been easily misdiagnosed due to its autolimited course. To the best of our knowledge, this is the first report of simultaneous facial diplegia and vestibular neuritis revealing HIV infection. This diagnosis could be made since the extensive search for other causes was negative, serological testing for HIV infection was positive and all symptoms resolved spontaneously.

Without a high index of suspicion, physicians may miss the diagnosis of acute HIV infection. Symptoms, if present, are nonspecific and resolve spontaneously without treatment. Moreover, patients generally have no findings on physical examination except for possible lymphadenopathy. This report emphasises that cranial nerve involvement such as facial palsy or vestibular neuritis can be the first symptom of HIV infection. Therefore HIV should be considered in the investigation of these entities, more concretely facial diplegia, especially in patients with high-risk behaviours even in absence of the mononucleosis-like syndrome. Given that patients are typically highly infectious during acute HIV due to a very high viral load8, the identification of these patients, who are unaware of their seropositive status, is clearly important from a public health perspective.

REFERENCES