Guillain-Barré syndrome associated with hepatitis A in a male homosexual

A DUNK, W J JENKINS, AND S SHERLOCK

From the Academic Department of Medicine, Royal Free Hospital School of Medicine, London

SUMMARY A 48-year-old male homosexual developed the Guillain-Barré syndrome in association with acute hepatitis. The hepatitis A virus was almost certainly transmitted sexually. Since the incidence of viral hepatitis is high in active male homosexuals, they are particularly at risk of developing such complications.

Introduction

Sexually transmitted viral hepatitis is an increasingly important problem in active male homosexuals.\(^1\)\(^2\) Although hepatitis B infection usually causes a more severe illness than hepatitis A and may cause chronic liver disease, hepatitis A is more common\(^1\) and may occasionally cause other serious complications as well as fulminant hepatic failure. We report the potentially fatal complication of Guillain-Barré syndrome with serologically confirmed hepatitis A in an active male homosexual.

Case report

A previously healthy 48-year-old man felt unwell with headache, fatigue, and myalgia. Three days later he noticed dark urine and paraesthesia in his hands and feet. The next day his legs and upper arms were weak and he became jaundiced.

He had not had any injections, tattoos, or blood transfusions and did not abuse drugs. He had not travelled abroad recently nor had any contact with jaundiced people. Nevertheless, he had been an active homosexual for 10 years. He had between 20 and 25 new partners each month, with whom he had oroanal contact.

CLINICAL FINDINGS

On admission to hospital four days after the onset of his symptoms he was slightly jaundiced, but there were no signs of chronic liver disease. There were, however, pronounced neurological abnormalities.

He was hypotonic and areflexic. Power was generally reduced in the arms and legs, particularly proximally. Joint-position sense and vibration sense were diminished in the feet, but sensory testing was otherwise normal.

LABORATORY FINDINGS

The serum liver function tests were abnormal. Serum aspartate aminotransferase was 237 IU/l (normal 5-40 IU/l) and alkaline phosphatase 458 IU/l (normal 30-110 IU/l); serum bilirubin concentration was 29 \(\mu\)mol/l (normal 5-17 \(\mu\)mol/l). Hepatitis B surface antigen (HBsAg) and antibody (HBsAb), hepatitis B core antigen (HBcAg), and IgM antibody were all absent; serological test results for recent Epstein-Barr virus and cytomegalovirus infections were negative. Hepatitis A IgM antibody measured by radioimmunoassay was detectable.

COURSE OF ILLNESS

The weakness of the limbs progressed, and three days after admission the patient was unable to walk or feed himself. The following day he developed bilateral lower motor neurone VII cranial nerve palsies and was unable to sit up. He also became dysarthric, but swallowing and the gag reflex remained normal.

A lumbar puncture showed normal pressure (100 mm Hg); the cerebrospinal fluid contained no cells, 0·3 g/1 protein, and 3·5 mmol/l (63 mg/100 ml) glucose. In view of these somewhat surprising results nerve conduction studies were performed. Motor conduction in the right median nerve was reduced at 37 m/s and the terminal latency was prolonged (6·8 ms). The F wave latency with elbow stimulation was increased (37 ms). No sensory nerve action potential was detectable. These abnormalities of conduction are consistent with the clinical diagnosis of the Guillain-Barré syndrome.

Accepted for publication 10 January 1982
The patient made a rapid recovery from hepatitis. By the eleventh day after admission the prothrombin time had improved sufficiently for a liver biopsy to be performed safely; it showed the changes of recent acute hepatitis. Meanwhile, the patient was also recovering power in the limbs and he could walk with a frame 12 days after admission. He was well enough to be discharged three days later. When he was next seen two months afterwards, power in the limbs was normal, but the VII cranial nerve palsies persisted. Five months after the initial symptoms there was further improvement, but a mild left VII cranial nerve palsy remained.

Comment

The presence of IgM antibody against hepatitis A virus identified it as the cause of the acute hepatitis in this patient. Hepatitis A is now considered to be one of the enteric infections that are transmitted sexually among homosexual men. Our patient, who was frequently the oral partner in oroanal contact, was particularly at risk, since Corey and Holmes showed that the acquisition of hepatitis A infection was correlated with this form of sexual activity.2

The progressive symmetrical polyradiculopathy, which developed in this patient in association with the acute hepatitis, is typical of the Guillain-Barré syndrome. The diagnosis is essentially clinical and supported by nerve conduction studies.3 Although the protein content of the CSF is usually high, it may be normal, especially in the early stages, and in some cases remains so.4

The major danger in the Guillain-Barré syndrome is paralysis of the respiratory muscles, but the prognosis is generally good. Approximately 75% of patients recover totally,5 and the delayed improvement of the facial nerve palsies in this patient is typical.

The Guillain-Barré syndrome is often associated with some form of bacterial or viral infection. Respiratory infections are the most common.6 The association with hepatitis is unusual. In a series of 1100 cases of Guillain-Barré syndrome Leneman6 reported 11 associated with acute hepatitis. This report6 and others7,8 preceded the development of specific serological tests to identify the hepatitis viruses. Subsequently, a few cases of Guillain-Barré syndrome associated with hepatitis B have been reported,9,10 but there has only been one previous report associated with hepatitis A.11

Because the incidence of viral hepatitis is high in active male homosexuals, they may be particularly at risk of developing such complications.

References

Guillain-Barré syndrome associated with hepatitis A in a male homosexual
A Dunk, W J Jenkins and S Sherlock

Br J Vener Dis 1982 58: 269-270
doi: 10.1136/sti.58.4.269

Updated information and services can be found at:
http://sti.bmj.com/content/58/4/269

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/