Diclofenac-induced hepatotoxicity

D. Schapira, L. Bassan, A.M. Nahir and Y. Scharf

The B. Shine Department of Rheumatology and the Department of Pathology, Rambam Medical Center, and Faculty of Medicine, Technion-Israel Institute of Technology, Haifa, Israel.

Summary: Two patients developed clinical, biochemical and histopathological signs of a liver hypersensitivity reaction following treatment with diclofenac. Hepatic side effects of this drug are very rare. The relevant literature is reviewed.

Introduction

Clinical and laboratory side effects of non-steroidal anti-inflammatory drugs are well known. Hepatotoxicity is one of the rare side effects of aspirin (Seaman et al., 1974), indomethacin (Kelsey & Scharyj, 1967), naproxen (Victorino et al., 1980), phenylbutazone (Benjamin et al., 1981), sulindac (Whittaker et al., 1982) and other drugs (Zimmerman, 1981).

Diclofenac sodium (Voltaren) is a potent and widely used non-steroidal anti-inflammatory and analgesic compound. It is classified among the most powerful drugs of this kind, while being one of the best tolerated (Brunner & Krupp, 1976). Diclofenac and its metabolites are excreted in both urine and bile. Experiments showed that the main route of drug elimination is different in various species, renal excretion being the most important in man and rhesus monkey and biliary excretion the most important in rat and dog. The lack of enterohepatic recycling in man probably accounts for the reduced gastrointestinal toxicity of this drug (Meness et al., 1978). Hepatic side effects are very rare. We report two patients who developed acute hepatotoxicity shortly after the initiation of treatment with diclofenac.

Case reports

Case 1

A 68 year old woman had diffuse joint pains and recurrent knee effusions. Treatment with diclofenac 100 mg/day brought relief. However, 2 weeks later, low grade fever and lassitude appeared together with pathological values of bilirubin, alkaline phosphatase and transaminase. The drug was withdrawn and the pathological, clinical and laboratory findings disappeared. Other non-steroidal anti-inflammatory drugs failed to improve the joint pains and diclofenac was reintroduced. A week later fever and lassitude returned together with a diffuse maculopapular skin rash, clinical jaundice and abnormal liver function: bilirubin 88.9 μmol/l (normal < 20.0) (54.7 μmol/l direct), aspartate transaminase 90 IU/l (normal 5–42), alkaline phosphatase 274 IU/l (normal 20–85). The erythrocyte sedimentation rate was 30 mm/h, the white cell count 15 × 10⁹/l (42% eosinophils). Hepatitis B surface antigen (HB,Ag), antismooth muscle and antimitochondrial antibodies were absent. There was a mild elevation of the IgG, the serum levels of IgM and IgA were normal. Immunological tests for collagen diseases and serological tests for various bacterial or viral agents were negative.

Scintigraphy revealed no hepatomegaly. The microscopical examination of the skin biopsy revealed normal epidermis. In the upper dermis a mixed infiltrate around the small blood vessels was seen (Figure 1). The infiltrate consisted mainly of lymphocytes but a few polymorphonuclear cells were also present. The bone marrow biopsy showed a few granulomas without necrosis consisting of Langhan's type giant cells, epitheloid cells and many eosinophils (Figure 2).

Permission for liver needle biopsy was not obtained. The drug was discontinued once more. The clinical jaundice and the skin rash rapidly disappeared. Both liver function and blood count returned to normal within 2 months.

Case 2

A 70 year old woman was admitted for recurrent inflammation of the wrists, the knees and the finger joints. The erythrocyte sedimentation rate was 47 mm/
hour, the complete blood count, liver function tests, blood proteins and protein electrophoresis were normal, and rheumatoid and antinuclear factor were absent. Calcium pyrophosphate deposition disease with diffuse pseudogout attacks was diagnosed and treatment with diclofenac, 150 mg/d was initiated. Five days later a diffuse maculopapular skin rash and pruritus appeared and icteric sclerae were noticed. The stools were acholic and the urine was dark. The bilirubin was 82 μmol/l (44.5 μmol/l direct), the aspartate transaminase 100 IU/l, the alkaline phosphatase 173 IU/l and the lactic dehydrogenase 626 IU/l (normal 115–480). The white blood cells were 8.8 × 10⁹/l (6% eosinophils). The liver technetium scan was normal and abdominal echography failed to disclose extrahepatic biliary obstruction. HBsAg, antimitochondrial, antismooth muscle and antiparietal cell antibodies were absent.

Liver needle biopsy revealed slight fatty metamorphosis of the liver cells. There was centrilobular confluent hepatocyte loss (Figure 3a) associated with mixed lymphocytic and polymorphonuclear infiltrate. The hepatocytes showed ballooning and there were a

Figure 1 Case 1. Vasculitis in upper dermis (H & E × 125).

Figure 2 Case 1. Bone marrow granuloma (H & E × 500).

Figure 3 Case 2. (a) Liver biopsy showing centrilobular confluent hepatocyte loss (H & E × 325). (b) Portal tract with giant cell Langhans' type surrounded by mononuclear infiltration (H & E × 325).
few acidophilic bodies. A single portal non-caseating granuloma was found (Figure 3b) with one Langhan's type giant cell and mononuclear infiltration around it. No microorganisms were demonstrated.

The stains for iron and HbAg (Shikata technique) were negative. Drug-induced hepatitis was diagnosed. The withdrawal of diclofenac brought a gradual clinical and laboratory remission within 1 month.

Discussion

Minor abnormalities of liver function (Shiokawa et al., 1972; Nasution, 1976; Trang et al., 1976; Ciccolunghi et al., 1978, 1979; Ciucci, 1979; McMahon & Cash, 1979), as well as isolated cases of jaundice (Dunk et al., 1982; Babany et al., 1983) have been reported in patients treated with diclofenac. Concurrent therapy with other drugs or previous liver and biliary tract disease in most of the cases made the diagnosis of diclofenac-induced hepatic damage uncertain. In our two cases the correlation between the drug and the liver damage is most probable.

Unfortunately in the first case a liver needle biopsy was not performed, but remission after removal of the drug as well as the hypersensitivity type of drug reaction as shown by the skin vasculitis, eosinophilia and histological picture of the bone marrow incriminate diclofenac as responsible for the liver injury. In the second case histological findings in the liver biopsy strongly suggest an allergic drug reaction and the diagnosis is supported by concomitant pruritus, skin rash and eosinophilia.

In conclusion, diclofenac, like other non-steroidal anti-inflammatory drugs, can cause hepatocellular damage. Fortunately, this side effect is very uncommon but nevertheless the diagnosis of drug-induced hepatotoxicity should be considered in patients who develop hepatic damage while receiving the drug.

References


Diclofenac-induced hepatotoxicity.

D. Schapira, L. Bassan, A. M. Nahir and Y. Scharf

doi: 10.1136/pgmj.62.723.63

Updated information and services can be found at:
http://pmj.bmj.com/content/62/723/63

Email alerting service

These include:
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/