Dear Editor,

Minocycline, a derivative of tetracycline, is an antibiotic commonly employed in the treatment of acne vulgaris for both its effectiveness and the long half-life (11-13 hours), which allows for a once to twice a day dosing schedule (1), and it accounts for 40% of the total tetracycline use (2). We describe a case of acute hepatitis caused by this antibiotic.

Case report

A 80 years old lady came to our attention in February 2007 due to dysphagia and increase of liver enzymes (AST 194 U/L, ALT 308 U/L). She had been treated with minocycline, 100 mg/day, from February 2005 to March 2006, for perioral dermatitis.

Upper endoscopy revealed a rabbit-bone impacted in the proximal third of the esophagus, removed with grasping forceps. Serological markers for hepatitis were negative; mild anemia (Hb 12.8 g/dl) and leukopenia (3,300/mmc) with a modest increase of gammaglobulines (20.4%) were noted. Anti-nuclear antibodies were present (1:160, homogenous pattern) while anti-mitochondrial, anti-liver-kidney and anti-endomyosial antibodies were negative. No Ig deficit was present.

The patient was discharged with a follow-up program. After one month, liver enzymes increased (AST 652 U/L, ALT 721 U/L, GGT 107 U/L), as well as anti-nuclear antibodies (1:320), and the liver function was mildly impaired (prothrombin time 75% with INR of 1.22, bilirubin level 2.2 mg/dl with direct bilirubin 0.7 mg/dl, cholinesterases 4939 U/L).

Liver biopsy revealed intact lobular architecture with diffuse dilation of portal spaces, inflammatory infiltrate prevalently represented by lymphocytes and monocytes, and sept development. Ductular degeneration area, focal cholangitis, severe piece-meal necrosis, several areas of lobular necrosis (zone 3) and perivenular necrosis with portal-central bridges were found (Fig. 1). Prednisone, 50 mg/day, was started with a low tapering schedule and, after one year, the patient showed a complete normalization of liver enzymes. In June 2008, steroid therapy was stopped and the patient is presently enjoying good health.

Discussion

Drug-induced hepatitis is an uncommon entity in clinical hepatology but does represent a significant proportion of acute hepatitis. Since the clinical picture is similar to viral hepatitis, the diagnosis is usually delayed. In our case, despite the patient had been treated with minocycline for about one year, it was not suspected, possibly due to the relatively short time from the withdrawal of the drug (one month) to the appearance of symptoms. The diagnosis was made through careful review of the clinical history and laboratory tests, together with histological examination of liver biopsy. The early detection of this condition is important to prevent irreversible liver damage and to discontinue the drug as soon as possible. Further studies are needed to better understand the mechanism of drug-induced hepatitis and to improve the diagnostic approach.
self-limited liver disease (3). The exact mechanism is not clearly understood, but many antibiotics, or their active metabolites, can bind to cellular macromolecules forming neo-antigens that appear extraneous to the immune system; the immune responses against such self components can trigger auto reactions to their unmodified counterpart and/or other auto antigens through epitope spreading (4).

Hepatotoxicity of tetracyclines was first described soon after the introduction in 1950, with minocycline being the most widely incriminated molecule. Since then, other cases have been described (5,6); it may be estimated that the overall literature report of minocycline hepatitis includes less than 200 cases.

Three types of minocycline-induced hepatitis have been described: typical toxin-mediated hepatic injury; fulminant hepatic failure as part of an allergic, idiosyncratic reaction chronic active hepatitis (CAH) detected by biopsy with positive ANA antibodies.

Given the possibility of liver injury related to minocycline use, it is important for dermatologists and pediatricians to be aware of this important side effects of minocycline therapy. Currently, there is no way to predict which patients are at risk to develop autoimmune hepatitis after exposure to the drug. A careful drug history should always be considered in any patient with chronic hepatitis because minocycline is often wrongly considered a cosmetic medicine and, in this particular type of patients, may determine an irreversible liver failure. An early recognition and discontinuation of the drug is important to aid recovery and to avoid the use of steroid and/or immunosuppressive agents.

Elderly patients may have an indolent progressive disease that is clinically asymptomatic. An acute, even fulminant, presentation may reflect a "de novo" disease or an exacerbation of a pre-existent chronic process. Centrilobular (zone 3) necrosis reflects an early histological stage that may evolve later in classical interface hepatitis. Prednisone (also in combination with azathioprine) is the safest effective initial treatment and all elderly patients with severe disease should be treated vigorously even though immunosuppressive medications, especially when used in combination are associated with an increased risk of opportunistic infections.

In conclusion, physicians should be aware of liver injury related to minocycline, since this side effect, although generally benign, may sometimes evolve in serious, even fatal hepatic insufficiency. Thus, a prompt recognition of potential liver injury related to the use of this drug is mandatory, in order to start as soon as possible an adequate therapeutic regimen and avoid more serious complications.

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References