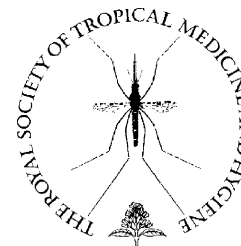




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CASE REPORT

First case of ivermectin-induced severe hepatitis

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Summary Loiasis, caused by the filarial parasite *Loa loa*, is endemic in West and Central Africa. Ivermectin has been shown to be an effective treatment of loiasis. We report the case of a 20-year-old woman originally from Cameroon who was infected by the *L. loa* parasite and developed severe hepatitis, identified 1 month after a single dose of ivermectin. Liver biopsy showed intralobular inflammatory infiltrates, confluent necrosis and apoptosis, compatible with drug-induced liver disease. To our knowledge, this is the first case of ivermectin-induced severe liver disease published in the literature.

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1. Case report

In October 2000, a 20-year-old woman was referred to the Swiss Tropical Institute, Basle, Switzerland, for evaluation of a migrating worm in the sclera of her right eye (Figure 1). She reported general body itching over several years, headache and dizziness. She had lived in Cameroon until the age of 15; she had moved to Switzerland 5 years previously and had not visited Africa since then. She had had meningitis with subsequent persistent deafness and a history of hepatitis A and B. Initial physical examination was inconspicuous except for moderate abdominal pain.

The worm that was removed from the patient's right eye was identified as an adult *Loa loa*, with a length of 2 cm. Initial *L. loa* microfilaraemia was 820/ml of whole

blood. There was an eosinophilia of 18.5% (absolute count 350/ μ l), and liver enzymes were normal. The patient was treated with albendazole (600 mg/d) for 21 d. One month after the end of therapy, the microfilaraemia dropped to 250/ml and was still at this level 3 months thereafter (Table 1). Clinical signs and liver enzymes remained normal, and no abdominal pain was reported. A single dose of 15 mg (300 μ g/kg) ivermectin, a dosage previously reported as having no major side effects (Kombila et al., 1998), was given to reduce the microfilaraemia further. At a routine follow-up 1 month after administration of ivermectin, the patient reported moderate new diffuse abdominal pain. Physical examination showed a new tenderness in the upper right quadrant; the liver was not enlarged. Laboratory investigation revealed elevated liver enzymes: ALAT 907 IU/l (normal range 7–42), ASAT 279 IU/l (normal range 5–39) and γ GT 66 IU/l (normal range 8–78); bilirubin, alkaline phosphatase, C-reactive protein, and red and white blood cell counts were in the normal range. Viral hepatitis were ruled out by means of serology (HAV, HBV, HCV, CMV, EBV). An HIV test was also negative. The patient had no his-

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Figure 1 Transocular migration of the *Loa loa* macrofilariae (2 cm) in the patient.

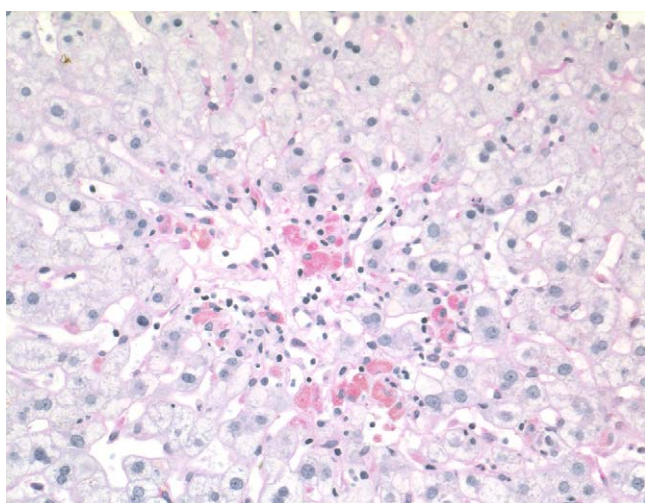


Figure 2 Liver biopsy with intralobular inflammatory infiltrates, focal ceroid loaded macrophages, perivenular confluent necrosis and intralobular apoptosis.

tory of alcohol or drug abuse. No other medications had been taken. A liver biopsy was performed, which showed predominantly perivenular intralobular inflammatory infiltrates, with conglomeration of ceroid loaded macrophages

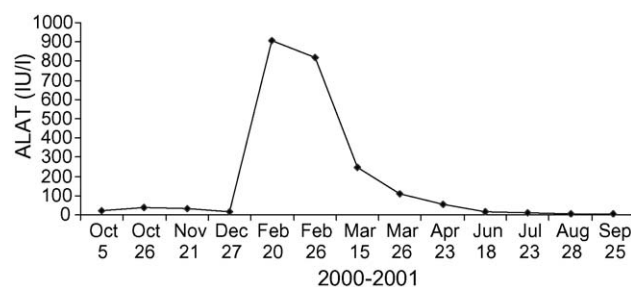


Figure 3 Follow-up of liver enzyme ALAT.

and confluent necrosis and apoptosis (Figure 2). The biopsy findings were consistent with resolving acute drug-induced hepatitis.

The patient became asymptomatic after a few days. Four months after the onset of hepatitis, all liver enzymes had returned to normal values (Table 1 and Figure 3). As microfilaraemia still persisted (19 mf/ml), diethylcarbamazine (DEC) was given for 29 d (total 7293.75 mg) and was well tolerated without any elevation of the liver enzymes. One week after initiating DEC treatment and in all subsequent examinations no microfilariae could be detected in the peripheral blood and no clinical sign of loiasis appeared.

2. Discussion

The filarial parasite *L. loa* is endemic in West and Central Africa, with an estimated 13 million people infected (Toure et al., 1998). Typical manifestations are transocular migration or subcutaneous swelling ('Calabar swelling'), caused by migrating adult nematodes (macrofilariae). Other common symptoms are generalized pruritus, fatigue and arthralgias. Hepatic dysfunction has not been reported. In exceptional cases, complications such as endomyocardial fibrosis and renal complications arise.

Diethylcarbamazine is the standard treatment of loiasis but is associated with an elevated risk of encephalitis in patients with high microfilaraemia (Carne et al., 1991). Ivermectin is used for treatment in many diseases, including loiasis, scabies, onchocerciasis and strongyloidiasis. In the treatment of onchocerciasis with ivermectin, where *L. loa* is endemic, encephalitis has been reported (Boussinesq et al., 1998; Ducorps et al., 1995; Gardon et al., 1997).

Table 1 Laboratory findings

	Date						
	5/10/00	27/12/00	20/2/01	26/03/01	23/04/01	13/7/01	23/7/01
Haemoglobin (g/dl)	11.8	13.6	13.2	12.5		12.4	12.1
Leucocytes ($10 \times 3/\mu\text{l}$)	12.7	7.0	8.0	10.5		8.7	9.5
Eosinophilia (%)	18.5	9.5	14.0	6.6		4.5	6.5
ALAT [7–42 IU/l]	21	35	907	111	54	23	13
ASAT [5–39 IU/l]	23	23	279	47	27	20	5
γ GT [66–93 IU/l]	22	42	66			27	27
Alkaline phosphatase [38–126 IU/l]	58	40	61	57	43	46	38
Bilirubin [2–24 $\mu\text{mol/l}$]	11	8	22	16	20	21	13
Microfilaraemia (<i>Loa loa</i>) (/ml)	820	220	9			19	0

In our patient the results of the liver biopsy were compatible with drug-induced hepatotoxicity. At the time of the diagnosis of drug-induced liver disease, the patient had received two drugs in the previous 5 months: albendazole and ivermectin. Albendazole, which has good effect against *L. loa* microfilariae, was initially chosen over standard therapy with DEC because of the mentioned elevated risk of encephalitis in patients with high microfilaraemia. Albendazole can produce hepatotoxicity, but in most cases this is associated with the treatment of echinococcosis, in which elevations of enzymes occur as the cysts are destroyed (Horton, 1989). There are a few cases of drug-induced hepatitis in the literature (Morris and Smith, 1987). In our case, the interval of several months between treatment with albendazole and onset of symptoms, as well as the normal liver enzymes following this treatment, are reasons to think that albendazole was not the cause of this event.

The shorter interval between administration of ivermectin and the onset of symptoms, liver histology compatible with hepatocellular toxicity, and no history of other agents causing hepatitis all point to ivermectin as causative agent of the drug-induced liver disease in our patient. Idiosyncratic drug-induced liver disease occurs in 1–10 in 10 000 patients taking a specific medicine. A review of published literature has not revealed any cases of severe ivermectin-induced liver disease, not even in escalating high doses of ivermectin (Guzzo et al., 2002). However, transient mild to moderate elevation of liver enzymes has been described (Biour et al., 2004; Zaha et al., 2002). Ivermectin is used for mass treatment against onchocerciasis, and also in *L. loa* endemic regions. However, a case like ours probably would not have been recognized in an area in Central Africa, where liver function tests are not routinely carried out, especially in the absence of jaundice. Furthermore, in a place where viral hepatitis is common, suspicion of drug-induced hepatitis would be rather low. Therefore, in the absence of controlled large clinical trials there is the potential to miss such variants of hepatitis.

3. Conclusion

We consider ivermectin to be the cause of drug-induced hepatitis in our patient. To our knowledge, this is the first case of ivermectin-induced severe liver disease published in the literature. In patients suffering from abdominal pain after treatment with ivermectin, liver cell damage should be considered.

Conflicts of interest statement

The authors have no conflicts of interest concerning the work reported in this paper.

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