Intrahepatic Cholestatic Hepatitis after Ajmaline and Flecainide Application in a Patient with Ventricular Tachycardia and Cyp2d6 Polymorphism

Case Report

Intrahepatic cholestatic hepatitis is a rare but therapy-limiting side effect of antiarrhythmic drugs, in literature only few reports are related to the subscription of ajmaline [1-3]. As potential causes for development of intrahepatic cholestatic hepatitis drug-triggered autoimmune reactions as well as direct toxic effects are recently discussed. However, the mechanism of this severe hepatic side effect remains unclear.

Here we present a 53-year-old male who was admitted to our emergency department with sudden-onset palpitation and dizziness without angina or dyspnoea. The initial ECG and the vital parameters revealed a hemodynamic stable ventricular tachycardia (VT) with 210 bpm. Tachycardia showed a right bundle branch block and left anterior hemiblock with a superior axis. The laboratory testing showed a slightly elevated troponin I (0.41 µg/l [ref.<0.04 µg/l]) and mild cholestasis (γGT 202 U/l [ref. < 56 U/l]) without hepatitis. A pre-hospital i.v. application of 300 mg amiodarone and 5 mg metoprolol by the emergency physician was effectless, whereas an application of 30 mg ajmaline i.v. induced a conversion to normofrequent sinus rhythm.

At the second day after admission the patient was referred to coronary angiography and right ventricular electrophysiological examination. A coronary heart disease could be excluded; no tachycardia could be induced during electrophysiological testing. The echocardiography showed a mild to moderate impaired left ventricular function with a global hypokinesis (LV-EF 40-45%). At the third day after admission a VT-storm occurred, which was successfully terminated several times (13 times) with i.v. bolus application of 30 mg ajmaline for each VT. Therefore we conducted an emergency left ventricle electrophysiological examination, which revealed a flecainide-sensitive idiopathic focal ventricular tachycardia located in the inferolateral wall (Figure 1). Subsequently we performed a successful catheter-assisted endocardial ablation by use of 3D-mapping. After the procedure we started oral flecainide (300 mg/d) for long-term therapy as relapse prevention, as the patient’s hemodynamics was impaired due to VT-storm. Hereafter no recurrence of ventricular tachycardia was observed. A post-procedural magnetic resonance tomography of the heart showed a marked late-enhancement in projection to the ablation region with a LV ejection fraction of 42%. After all, as a secondary prophylactic step and a safety procedure concerning a relapse of VT we implanted an ICD-system (day 14 after admission).

Seven days after discharge the patient complained of pruritus, jaundice, fatigue and pain in the right upper abdomen for the duration of three days. Laboratory testing revealed the constellation of cholestatic hepatitis with strikingly elevated cholestatic enzymes (γ-glutamyl transferase (γGT) 2037 U/l [ref.<56 U/l], Alkaline phosphatase (AP) 394 U/l [ref.<127 U/l]) and bilirubinemia (3.7 mg/dl [ref.<1.2 mg/dl]), accompanied by a significant eosinophilia (0.7/nl [ref.<0.45/nl]). Infective (viral hepatitis A, B, C) and autoimmune (IgM, IgG, ANA, AMA-M2) causes of hepatitis were excluded. Ultrasound examination of the abdomen showed slightly hyperechoic parenchyma of the liver, whereas intra- and extrahepatic bile ducts were normal.

In consideration of the predominantly elevated cholestatic enzymes, the eosinophilia and the patients’ history we suspected drug-induced cholestatic hepatitis due to ajmaline or flecainide.

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hepatitis due to antiarrhythmic medication, probably related to ajmaline. Despite metabolization by CYP2D6, the long-term medication with flecainide was necessary to avoid a relapse of potential life-threatening ventricular tachycardia, so that we did not interrupt this therapeutic strategy. Nevertheless we reduced the flecainide-dose adjusted to the reduced enzyme-activity. Compared to ajmaline, flecainide is supposed to have minor clinical significance regarding CYP2D6 metabolism [5-8]. Consistent with this assumption a continuation of flecainide therapy did not hinder recovery of the cholestatic hepatitis, even though a prolongation cannot be excluded.

Our case demonstrates that a continuation of dosage-reduced flecainide is reasonable in an intermediate metabolizer of CYP2D6 while subsiding cholestatic hepatitis. As mentioned above our patient had a mild cholestasis at admission, which is associated with a higher risk for development of drug-induced cholestasis [9].

Furthermore, genotype testing of Cytochrome P450 2D6 (CYP2D6) revealed a heterozygous genotype with a defective CYP2D6 allele (CYP2D6*5) resulting in a decreased enzyme activity accordingly to an “intermediate metabolizer”.

Ajmaline and flecainide are well-known antiarrhythmic drugs and indicated for emergency treatment of supraventricular and ventricular tachycardia. These antiarrhythmic drugs are metabolized in the liver via the cytochrome P450 enzyme CYP2D6 [5,6]. The polymorphism of this enzyme results in poor, intermediate, efficient or ultrarapid metabolizers of CYP2D6 drugs. About 5-10% of the European Caucasian population are poor metabolizers and nearly 10-17% are intermediate metabolizers [7].

Drug-induced cholestasis may occur under conditions of increased drug concentrations or genetic alterations such as CYP2D6 polymorphism. In accordance to the few reported cases of ajmaline-induced cholestatic hepatitis [1,2], our patient had a similar clinical course characterized by a latency of nearly 3 weeks until occurrence of symptoms and liver enzyme activation. Previously, Mellor et al. supposed that the mechanism of ajmaline-induced cholestatic hepatitis may relate to CYP2D6 polymorphism. Recently, to the best of our knowledge and in comparison to previous reports we demonstrate for the first time that a CYP2D6 polymorphism leads to intrahepatic cholestatic hepatitis due to antiarrhythmic medication, probably related to ajmaline. Despite metabolization by CYP2D6, the long-term medication with flecainide was necessary to avoid a relapse of potential life-threatening ventricular tachycardia, so that we did not interrupt this therapeutic strategy. Nevertheless we reduced the flecainide-dose adjusted to the reduced enzyme-activity. Compared to ajmaline, flecainide is supposed to have minor clinical significance regarding CYP2D6 metabolism [5-8]. Consistent with this assumption a continuation of flecainide therapy did not hinder recovery of the cholestatic hepatitis, even though a prolongation cannot be excluded.

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Furthermore, genotype testing of CYP2D6 is not routinely implemented in the clinical setting for risk stratification concerning the development of severe hepatic side effects following treatment by antiarrhythmic drugs. In consideration of the clinical situation the evaluation of CYP2D6 polymorphism in patients at higher risk for liver side effects and medication with antiarrhythmic agents, which are metabolized by this enzyme could be helpful for prevention of adverse drug reactions. This algorithm needs to be proven in further studies. Nevertheless a strict clinical and laboratory observation of these patients in an outpatient manner is necessary.
References


