Sotalol-induced torsade de pointes: management with magnesium infusion

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Summary: A 69 year old woman was treated with sotalol (320 mg daily) for intermittent atrial fibrillation. Sotalol was initially well tolerated, and reversion to sinus rhythm with sinus bradycardia occurred 4 weeks after initiation of therapy. Shortly thereafter, the patient developed recurrent syncope due to torsade de pointes. This was treated successfully with intravenous magnesium infusion and withdrawal of sotalol. Subsequently, the atrial fibrillation was adequately managed using amiodarone, with no recurrence of torsade de pointes. Development of bradycardia associated with reversion to sinus rhythm represents a potential cause of 'late' pro-arrhythmic effects of sotalol.

Introduction

Torsade de pointes (TDP) occurring during treatment with the anti-arrhythmic agent sotalol, has frequently been associated with the presence of complicating factors such as drug overdose or more commonly, hypokalaemia. We wish to report a case of sotalol-induced TDP in which no such complicating factor was clinically apparent.

Case report

A 69 year old woman was admitted for management of symptomatic atrial fibrillation of 5 days duration. There was a past history of mitral valvotomy. Rapid ventricular response to the atrial fibrillation persisted despite treatment with digoxin and verapamil. She had also been treated with frusemide and warfarin. Verapamil was ceased and sotalol commenced, initially at a dose of 160 mg daily, with her other treatment unchanged. The sotalol dosage was increased to 320 mg daily after 3 weeks. Reversion to sinus rhythm, with subsequent asymptomatic sinus bradycardia, occurred 4 weeks after initiation of sotalol treatment.

Twenty-four hours after reversion to sinus rhythm, the patient developed recurrent syncopal episodes preceded on each occasion by palpitations. Ambulatory electrocardiogram (ECG) monitoring revealed several episodes of short-lived TDP associated with pre-syncope (Figure 1). The patient was transferred to the coronary care unit and both sotalol and digoxin were ceased. An intravenous bolus of 15 mmol magnesium sulphate was administered over 1 min, with no adverse effect, followed by a 24 h infusion at a rate of 8 mmol/h. All ventricular ectopy ceased after the bolus injection, with no recurrence of TDP.

Investigations at the time of this presentation included ECG, which revealed left bundle branch block with some incremental prolongation of the QTc (0.51 s before sotalol therapy compared to 0.56 s while on sotalol therapy), serum potassium level of 3.9 mmol/l (R: 3.2–5.3) and serum magnesium of 0.87 mmol/l (R: 0.75–1.13).

Therapy with amiodarone was subsequently initiated on recurrence of rapid atrial fibrillation, and the patient was discharged without recurrence of symptoms over the next 2 years. During this time the rhythm has been controlled atrial fibrillation.

Discussion

This patient developed TDP while being treated with racemic sotalol in the usual clinical dosage. The current case represents the fourth report in the literature of this phenomenon. In the three previously reported cases, no precipitating factor other than sotalol therapy could be delineated, and plasma sotalol concentrations appeared unremarkable when measured.

The issue arises as to whether this patient might have been predisposed to TDP on the basis of a congenital prolonged repolarization syndrome. While this remains possible, the patient had...
References


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