Antiarhythmic Medication Induced Multi-Organ Dysfunction

Abstract
Propafenone is a Class 1 C drug indicated for reduction of episodes of paroxysmal atrial fibrillation. There is a paucity of literature to guide the escalation of propafenone in a patient who has been on long-term therapy. Multi-organ toxicity following dose escalation within the recommended therapeutic dose range appears to be a rare event. We report a patient who presented with severe systolic heart failure six weeks after propafenone dose escalation. We discontinued propafenone therapy and the patient had full clinical and echocardiographic recovery.

Keywords: Propafenone; Cardiotoxicity; Hepatotoxicity; Dose escalation

Introduction
Propafenone used in the treatment of atrial fibrillation and supraventricular tachycardia is a class IC anti-arrhythmic medication which is generally safe at when administered in the therapeutic range. Under rare circumstances, toxicity involving the cardiovascular system can occur.

Case Report
A 77-year-old Caucasian female with history of chronic atrial fibrillation (AF) on propafenone 150 mg three times daily was admitted for uncontrolled ventricular rates and dyspnea on exertion. Physical exam was relevant for irregularly irregular heart rate at 145 beats per minute (bpm). Her initial electrocardiogram (ECG) revealed atrial fibrillation at a rate of 145 beats per minute (Figure 1). Labs and transthoracic echocardiogram (TTE) were in normal limits. Her propafenone was increased to 225 mg three times daily which did not improve her AF. She was then successfully electrically cardioverted to normal sinus rhythm two days later and was discharged on this increased dose of propafenone.

She was admitted 6 weeks later with dyspnea and palpitations. Physical exam revealed irregularly irregular heart rate. Labs showed elevated liver enzymes (AST: 1049 U/L, ALT:577 U/L) and international normalized ratio (INR) of 23. Labs for viral hepatitis and autoimmune hepatitis were negative. ECG revealed AF with heart rate of 111 bpm with left bundle branch block (Figure 2). TTE was significant for ejection fraction of 20 % and paradoxical septal wall motion. Propafenone toxicity was high on the differential was causing the present clinical picture and it was discontinued along with warfarin which she was on for anticoagulation.

After discontinuation of the propafenone, she improved clinically with normalization of transaminitis, INR, ECG showing rate of 88 bpm with loss of left bundle branch block and TTE showing 60-65% ejection fraction on day 10 of her admission. She was discharged and has been doing well at her outpatient follow up appointments.

Discussion
Propafenone (2′-[3-(propyl amino)-2-(hydroxy)-propoxy]-3-phenylnpropophenone hydrochloride) is a Vaughn Williams Class 1 C antiarrhythmic drug with proven efficacy in reduction of episodes of supraventricular tachycardia (SVT) and AF [1,2]. Guidelines on AF management state that propafenone may be initiated as an out-patient if care is taken to ensure the patient has no structural heart disease, no sinus node or atrio-ventricular node disease and no Brugada pattern on EKG [3].

Propafenone increased the median free interval without recurrence of arrhythmia to 44 days in the ERAFT (European Rythmol/Rytmonorm Atrial Fibrillation Trial) trial [1], a mean of 98 days in the PSVT study [2] and over 300 days in the RAFT (Rythmol Atrial Fibrillation Trial) trial [4]. Propafenone has also been shown to be effective in converting patients with recent onset of atrial fibrillation with reported success rates of conversion between 72 and 94 % [5,6]. A recent meta- analysis found that unlike flecainide, propafenone does not have an adverse effect on mortality and it is not pro-arrhythmic [7].
In the study of patients with PSVT or atrial fibrillation, congestive heart failure occurred in 1.9% of patients [2]. Reduction in ejection fraction is confined to those with preexisting reduced ejection fraction [8]. Liver toxicity is a rare side effect of propafenone. In a review of nine cases reported in literature propafenone toxicity was noted a median of 4 weeks after propafenone initiation [9]. A nonlinear relationship between dose and steady state concentrations has been noted. This means a large increase in plasma concentration can result from a small dose increase [10]. We did not come across recommendations when adjusting the dose of propafenone on patients who have been on chronic therapy. We did however find two other cases in literature of marked reduction in systolic function in patients who had been on long term therapy with propafenone and had undergone a medication dose increase [11,12].

Our patient is noteworthy for a few reasons. Her ejection fraction fell from normal to an estimated 20%. From our review of literature, it appears such a precipitous decline in ejection fraction with propafenone dose increases in the usual therapeutic range

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is rare. The most remarkable finding in her ECG at the time of maximal toxicity (when her QRS interval was 176) is the marked left bundle branch block. This caused marked dyssynchrony noted on her TTE.

We believe that our patient's cardiac and hepatic dysfunction was caused by propafenone considering the sub-acute worsening of her clinical picture after propafenone dose increase, and subsequent improvement after discontinuation of the medication.

Conclusion

Propafenone is an effective drug in converting patients with recent onset AF and reducing recurrences of AF. Although rare, propafenone toxicity should be considered in patients having symptoms of systolic heart failure and hepatotoxicity soon after a dose increase. Monitoring for three days post dose increase may be insufficient in judging the risk of toxicity. It may be reasonable to review ECGs prior to any exposure to propafenone when making dose adjustments much later.

References