A 31 year old white man was referred by his general practitioner to the cardiology outpatient department for investigation of a persistent sinus tachycardia. In general he felt well, with no recent history of fever, chest pain, cough, or symptoms suggestive of thyrotoxicosis. On direct questioning there was a history of gradually worsening exertional dyspnoea over several months. His only past medical history was of chronic schizophrenia for which he had been taking clozapine for six years. An electrocardiogram demonstrated sinus tachycardia, voltage criteria for left ventricular hypertrophy, and a prolonged QTc. Echocardiographic findings were consistent with a dilated cardiomyopathy. Serious cardiac complications of clozapine use are rare but have been reported previously. It is important to note that sinus tachycardia may be the only obvious clinical sign, and that complications can manifest months or even years (as in this case) after starting the drug. Patients on clozapine should be informed of potential cardiac symptoms and doctors should maintain a high degree of clinical suspicion throughout the duration of treatment.

A 31 year old white man was referred for investigation of a persistent sinus tachycardia. His only significant past medical history was of chronic schizophrenia for which he had been taking clozapine for six years. An electrocardiogram demonstrated sinus tachycardia, voltage criteria for left ventricular hypertrophy, and a prolonged QTc. Echocardiographic findings were consistent with a dilated cardiomyopathy. Serious cardiac complications of clozapine use are rare but have been reported previously. It is important to note that sinus tachycardia may be the only obvious clinical sign, and that complications can manifest months or even years (as in this case) after starting the drug. Patients on clozapine should be informed of potential cardiac symptoms and doctors should maintain a high degree of clinical suspicion throughout the duration of treatment.

In the absence of any other likely aetiology a presumptive diagnosis of clozapine-induced cardiomyopathy was made and he was admitted for further management. The Committee on Safety of Medicines (CSM) was informed via the yellow card reporting scheme. His clozapine was converted to olanzapine 10 mg, and he was started on a combination of bumetanide, ramipril, and bisoprolol. After seven days there was subjective improvement in his exercise tolerance and there was no clinical evidence of cardiac failure. A predischARGE ECG demonstrated a normal QTc interval. He was discharged on the combination of β-blocker, loop diuretic, and angiotensin converting enzyme inhibitor in addition to his olanzapine.

At his six week follow up appointment there was further subjective improvement. He tolerated his drug regimen well and his mental state remained stable. On examination his sinus tachycardia had resolved and there was no evidence of cardiac failure.

His echocardiogram also demonstrated a significant improvement. Although the left ventricle remained dilated the overall systolic function had improved considerably with an estimated ejection fraction of 43%.

DISCUSSION

Clozapine is a dibenzodiazepine antipsychotic used in the treatment of resistant schizophrenia. Minor cardiovascular side effects, namely orthostatic hypotension and sinus tachycardia, are relatively common and have been previously described. There is also a well recognised 1% risk of agranulocytosis during the first year of treatment. Of great concern are the lesser reported serious and potentially fatal cardiac complications. In particular both myocarditis and dilated cardiomyopathy have been reported. To date the CSM has received 30 reports of cardiomyopathy and 40 cases of myocarditis associated with clozapine use. Killian et al presented 23 cases of myocarditis and cardiomyopathy temporally associated with this drug. Between 1989 and 1999 the United States Food and Drug Administration received reports of 28 cases of myocarditis (including 18 deaths) and 41 cases of cardiomyopathy (including 10 deaths) in patients receiving clozapine. Bayesian analysis as applied to the World Health Organisation drug monitoring database has demonstrated that clozapine is significantly more frequently reported in relation to myocarditis and cardiomyopathy than other drugs (including other antipsychotics). Estimation of the incidence of myocarditis in clozapine use ranges between one in 10 000 to one in 500. Interpretation of case reports is complicated by the often non-specific symptoms and signs of myocarditis and cardiomyopathy. ECG changes reported are also non-specific and include sinus tachycardia, T-wave inversion, prolonged QTc, and ST flattening/depression. It must be borne in mind that many of the cases reported have not had definitive evidence for their diagnoses.

It is, however, very likely that there is a causal link between clozapine and myocarditis cardiomyopathy. Several cases have been associated with a peripheral and/or cardiac eosinophilia suggestive of an IgE-mediated hypersensitivity reaction. The greatest risk of cardiac involvement most commonly occurs

Abbreviations: CSM, Committee on Safety of Medicines; ECG, electrocardiogram; QTc, corrected QT interval
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