Escitalopram Induced Torsade de Pointes and Cardiac Arrest in a Patient With Surgically Treated Mitral Valve Prolapse

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Abstract

A 54-year-old female, with a history of prosthetic mitral valve replacement due to mitral valve prolapse one year prior, was admitted after suffering a cardiopulmonary arrest. Her initial rhythm demonstrated Torsade de Pointes with the initial electrocardiogram (ECG) showing a prolonged QT interval. Laboratory test results were normal including potassium and magnesium serum levels, and imaging did not show significant abnormalities. A review of patients' medicines showed that the patient started taking escitalopram one month prior to the presentation for major depressive disorder. Selective serotonin reuptake inhibitors (SSRI) are widely prescribed and continue to be a mainstay of treatment for multiple psychiatric conditions. It is important to keep the potential cardiovascular side effects of SSRIs in mind when prescribing. Consideration of underlying cardiac conditions is vital to decrease the likelihood of poor outcomes.

Categories: Cardiology, Internal Medicine, Psychiatry

Keywords: out-of-hospital cardiac arrest, major depressive disorder, torsade de pointes, ssri, prolonged qtc interval

Introduction

Selective serotonin reuptake inhibitors (SSRIs) are a class of medications widely used in primary care for a variety of common disorders such as depression, anxiety, post-traumatic stress disorder, and chronic pain. QTc prolongation is a well-documented side effect of this class [1], with Torsade de Pointes (TdP) being less common [2]. Data on escitalopram is limited, with no demonstrated correlation between plasma level and length of QT interval [3].

Structural heart disease increases the risk of developing an arrhythmia; the presence of mitral valve prolapse (MVP) is associated with an increased risk of ventricular arrhythmias and sudden death. Arrhythmias have been known to persist even after the replacement of the defective valve [4]. Although MVP is known to be arrhythmogenic [5], there is often a trigger that sets off the arrhythmogenic cascade. We present a patient with a history of longstanding mitral valve prolapse one-year post repair. He was on escitalopram and presented with arrest due to TdP. In this patient, escitalopram likely was the provoking factor for arrhythmia. Primary care physicians should remember QTc prolongation as a potentially fatal side effect of SSRIs, particularly in patients with a history of structural heart disease, and actively monitor ECG pre and post-treatment initiation.

Case Presentation

A 54-year-old female, with a history of prosthetic mitral valve replacement due to mitral valve prolapse one year prior, was admitted after suffering a cardiopulmonary arrest. Cardiopulmonary resuscitation was initiated by bystanders and continued by emergency medical services (EMS). The initial recorded rhythm was Torsade de Pointes which degenerated to coarse ventricular fibrillation (Figure 1), requiring multiple defibrillation shocks before the restoration of sinus rhythm and return of spontaneous circulation (Figure 2). Initial 12 lead ECG showed sinus rhythm with prolonged QTc (Figure 3). Transthoracic echocardiogram showed an ejection fraction of 40-45% with a well functioning prosthetic mitral valve. Laboratory test results demonstrated normal serum electrolyte levels, including potassium and magnesium. A detailed medical and medication history was performed after patient achieved neurological recovery. The patient confirmed that she had been taking escitalopram for a month preceding the cardiac arrest for major depressive and generalized anxiety disorders. Review of her previous records revealed an ECG that was obtained prior to starting escitalopram showing normal QTc interval.

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FIGURE 1: Initial Rhythm: Torsade de Pointes

FIGURE 2: Torsade de Pointes transforming to ventricular fibrillation, followed by successful defibrillation

FIGURE 3: Post resuscitation 12 Lead EKG showing QTc prolongation

Discussion

SSRIs are commonly selected for the treatment of depression due to their efficacy and wide therapeutic index [6]. Though they are known to be associated with QTc prolongation, the incidence of significant arrhythmias is low and often occurs in patients with predisposing factors such as electrolyte abnormalities, congenital long QTc syndrome, and other structural cardiac diseases [7]. The pathophysiology of ventricular arrhythmias can be explained using the concept of an arrhythmogenic substrate that comprises a focal area that has the potential to generate abnormal electrical impulses, the triggers, and the modulators (Figure 4) [8]. Mitral valve prolapse (MVP) is often associated with a mechanical myocardial and papillary muscle injury and fibrosis that can form a focus from which aberrant electrical potentials can be generated [9]. Furthermore, Naksuk et al showed that mitral valve surgery did not necessarily decrease the burden of ventricular arrhythmias and proposed that the arrhythmogenic substrate formed following longstanding MVP is frequently not reversed by valve replacement surgery [4]. The ECG taken prior to surgical repair in our case, showed frequent ventricular ectopic beats in a pattern of ventricular bigeminy, hinting to the development of an arrhythmogenic focus associated with long term mechanical strain related to MVP.

FIGURE 4: Mechanism of Arrhythmia Development

The relationship between QTc prolongation, Torsade de Pointes, and ventricular fibrillation has been well established [10]. SSRIs, as a class, are known to cause QT prolongation, but only rarely cause TdP [11]. Escitalopram is an S-enantiomer of citalopram and the QTc prolongation effect of citalopram has largely been attributed to its metabolite didemethylcitalopram (DDCT) [12]. The mechanism of the QT prolongation involves the blocking of the inward potassium rectifier (IKr) current, prolonging phase 3 of cardiac repolarization and thus prolonging the QTc. If during this interval, there is repolarization originating from a different source, the compounding effect can lead to oscillating currents precipitating TdP. Thus, the QT-prolonging effect of escitalopram serves as the trigger to induce TdP in the susceptible area of the myocardium. In this case, the presence of an arrhythmic substrate from mitral valve prolapse and QTc prolongation from escitalopram are the factors that potentially acted synergistically to precipitate that ominous rhythm.

There are no clear guidelines regarding electrocardiographic monitoring in patients taking SSRIs, likely due to the relative rarity of severe arrhythmias. QTc prolongation and TdP related to SSRI are infrequent and sporadic and are usually associated with toxic drug levels or drug overdose excluding people who are at higher risk [13]. A baseline ECG is usually done before initiation of antidepressant therapy with a tricyclic anti-depressant and this has been extended to SSRIs [14].

Conclusions

This case serves to highlight the danger of using QTc prolonging agents in patients with a history of structural heart disease such as mitral valve prolapse, even after corrective repair. It is imperative that physicians be aware of the adverse cardiovascular effects of the SSRI including QTc prolongation and TdP. In our opinion, an initial EKG is warranted before prescribing SSRIs in the presence of structural heart disease including MVP. We feel that surveillance ECGs should be obtained when increasing medication dosage. Patients should also be educated regarding the potential adverse effects of the medication class.

Additional Information

Disclosures

Human subjects: Consent was obtained by all participants in this study. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following: **Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

1. Drye LT, Spragg D, Devanand DP, et al.: Changes in qtc interval in the citalopram for agitation in alzheimer's disease (citad) randomized trial. PLoS ONE. 2014, 9:98426. [10.1371/journal.pone.0098426](https://doi.org/10.1371/journal.pone.0098426)
2. Tampi RR, Balderas M, Carter KV, Tampi DJ, Moca M, Knudsen A, May J: Citalopram, qtc prolongation, and

- torsades de pointes. *Psychosomatics*. 2015, 56:36-43. [10.1016/j.psym.2014.09.002](https://doi.org/10.1016/j.psym.2014.09.002)
3. Carceller-sindreu M, Diego-adeliño JD, Portella MJ, et al.: Lack of relationship between plasma levels of escitalopram and qtc-interval length. *Eur Arch Psychiatry Clin Neurosci*. 2017, 267:815-822. [10.1007/s00406-016-0758-6](https://doi.org/10.1007/s00406-016-0758-6)
  4. Naksuk N, Syed FF, Krittawong C, et al.: The effect of mitral valve surgery on ventricular arrhythmia in patients with bileaflet mitral valve prolapse. *Indian Pacing Electrophysiol J*. 2016, 16:187-191. [10.1016/j.ipej.2016.10.009](https://doi.org/10.1016/j.ipej.2016.10.009)
  5. Miller MA, Dukkipati SR, Turagam M, Liao SL, Adams DH, Reddy VY: Arrhythmic mitral valve prolapse: jacc review topic of the week. *J Am Coll Cardiol*. 2018, 72:2904-2914.
  6. Marken PA, Munro JS: Selecting a selective serotonin reuptake inhibitor: clinically important distinguishing features. *Prim Care Companion J Clin Psychiatry*. 2000, 2:205-210. [10.4088/PCC.v02n0602](https://doi.org/10.4088/PCC.v02n0602)
  7. Yekhtaz H, Farokhnia M, Akhondzadeh S: Cardiovascular considerations in antidepressant therapy: an evidence-based review. *J Tehran Heart Cent*. 2013, 8:169-76.
  8. Malik M: Arrhythmic substrate. *Heart*. 2008, 94:402-3. [10.1136/hrt.2007.124198](https://doi.org/10.1136/hrt.2007.124198)
  9. Basso C, Perazzolo marra M, Rizzo S, et al.: Arrhythmic mitral valve prolapse and sudden cardiac death. *Circulation*. 2015, 132:556-66. [10.1161/CIRCULATIONAHA.115.016291](https://doi.org/10.1161/CIRCULATIONAHA.115.016291)
  10. Yap YG, Camm AJ: Drug induced qt prolongation and torsades de pointes. *Heart*. 2003, 89:1363-72. [10.1136/heart.89.11.1363](https://doi.org/10.1136/heart.89.11.1363)
  11. Kogut C, Crouse EB, Vieweg WV, et al.: Selective serotonin reuptake inhibitors and torsade de pointes: new concepts and new directions derived from a systematic review of case reports. *Ther Adv Drug Saf*. 2013, 4:189-98. [10.1177/2042098613492366](https://doi.org/10.1177/2042098613492366)
  12. Howland RH: A question about the potential cardiac toxicity of escitalopram. *J Psychosoc Nurs Ment Health Serv*. 2012, 50:17-20. [10.3928/02793695-20120307-02](https://doi.org/10.3928/02793695-20120307-02)
  13. Alvarez PA, Pahissa J: QT alterations in psychopharmacology: proven candidates and suspects. *Curr Drug Saf*. 2010, 5:97-104. [10.2174/157488610789869265](https://doi.org/10.2174/157488610789869265)
  14. Dodd S, Malhi GS, Tiller J, et al.: A consensus statement for safety monitoring guidelines of treatments for major depressive disorder. *Aust N Z J Psychiatry*. 2011, 45:712-25. [10.3109/00048674.2011.595686](https://doi.org/10.3109/00048674.2011.595686)