Reversible Posterior Leucoencephalopathy Syndrome Following Duloxetine: A Case Report and Review of Literature

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Abstract

Reversible Posterior Leucoencephalopathy Syndrome (RPLS) is a disorder characterized by headaches, confusion, and/or decreased level of consciousness, seizures, and visual changes. A 58-year-old female was found unresponsive, with generalized rigidity and urinary incontinence for unknown duration. On presentation, she was febrile, tachycardic, tachypneic, and hypertensive. Neurological exam showed increased muscle tone and brisk deep tendon reflexes. Brain MRI revealed gyriform edema and white matter hypodensities in parieto-occipital regions consistent with RPLS. Discontinuation of duloxetine, one of her home medications, led to resolution of symptoms. We report this case to discuss the syndrome of RPLS and propose the role of duloxetine in causing hypertension and cerebral vasoconstriction leading to RPLS. To our knowledge, this is the first reported case of RPLS resulting from the use of duloxetine.

Keywords: Reversible posterior leucoencephalopathy syndrome; Duloxetine; Cerebral vasoconstriction; Adverse reactions

Introduction

Reversible posterior leucoencephalopathy syndrome (RPLS) as a distinct entity was first described in 1996 by Hinchey et al. This disorder is characterized primarily by headaches, confusion/decreased level of consciousness, seizures, and visual changes [1]. RPLS, as we know today, is a clinical radiologic syndrome linked to a wide variety of etiologies. By far the most common etiology linked to the pathogenesis of RPLS is hypertension [2, 3]. Other factors implicated in the development of RPLS include eclampsia [4], immunosuppressive medications [1, 5], and a variety of other drugs [5-7]. The pathogenesis of RPLS remains controversial but mainly involves loss of cerebral autoregulation and endothelial dysfunction [1]. Diagnosis is established by neuroimaging. Typical findings are symmetrical white matter edema in the posterior cerebral hemispheres, particularly the parieto-occipital regions [8]. Treatment of RPLS includes rapid lowering of blood pressure, antiepileptics, and withdrawal of offending agent [9, 10].

In our case, we discuss the syndrome of RPLS and we propose the possible role of duloxetine in causing hypertension and cerebral vasoconstriction leading to RPLS. To our knowledge, there has been no reported case of RPLS secondary to the use of duloxetine in the English literature.

Figure 1. MRI image of the brain showing gyriform edema and white matter hypodensities in parieto-occipital regions.
Case Report

A 58-year-old Caucasian female was brought to the emergency room by Emergency Medical Services (EMS) after being found with generalized body rigidity, urinary incontinence, and unresponsiveness for an unknown duration of time. The patient was intubated for airway protection by the EMS personnel. She was last seen normal the day prior to admission. Her past medical history was significant for hypertension, chronic pain syndrome, depression, and hypothyroidism. She had attempted suicide (overdose of antidepressants) twice in the past. Her home medications included duloxetine, zolpidem, cyclobenzaprine, pregabalin, oxycodone, amlodipine, and benazepril. She was working as a nurse at a psychiatric institute. There was no history of smoking, alcohol abuse, or drug abuse. At the time of initial presentation, her vital signs were blood pressure 220/126 mmHg, pulse 135/min, respiratory rate 42/min, temperature 103 degrees Fahrenheit and O2 saturation 97% on the ventilator. Physical exam revealed a well nourished, disheveled Caucasian lady who was intubated. Neurological exam was significant for bilaterally symmetric and reactive pupils, positive gag reflex, spontaneous movement in all extremities, increased muscle tone, brisk and symmetric deep tendon reflexes, downgoing bilateral plantar reflexes, and no signs of meningism or clonus. Higher mental functions could not be assessed as she was sedated. Rest of the systemic exam was normal. Initial laboratory data showed mild leucocytosis, normal complete metabolic panel, and mildly elevated troponins and CPK. Based on the initial presentation and concern for possible meningoencephalitis, the patient was started on a broad spectrum antibiotic coverage with vancomycin, rocephin, and ampicillin. Blood cultures, urine culture, and CSF gram stain and culture were obtained prior to initiation of antibiotics. On the second day of hospitalization, acyclovir was also added to the regimen for possible herpes simplex encephalitis. During the rest of the hospital course, the patient’s cultures came back sterile, herpes simplex PCR was negative and there was no growth of organisms from the CSF as well. EEG done on the patient showed a generalized slowing consistent with an encephalopathy. CT of the head done without contrast was unremarkable. MRI of the head revealed gyriform edema and white matter hypodensities in parieto-occipital regions (Fig. 1, 2). Considering the negative infective etiology and the MRI findings the diagnosis of RPLS was considered. Antibiotics were discontinued and supportive care was provided to the patient. The patient’s home medications that had been continued in hospital were reviewed again and duloxetine was discontinued, which led to complete resolution of symptoms within 5 days. The patient was discharged home to be followed up as an outpatient. Subsequently, the patient has had follow-up neuroimaging 6 weeks post her discharge with complete resolution of the initial MRI findings (Fig. 3, 4).

Discussion

Reversible Posterior Leucoencephalopathy Syndrome (RPLS) is a clinical radiologic syndrome that was first described as a separate entity in 1996 [1]. Some other names used for this syndrome include Posterior Reversible Encephalopathy Syndrome (PRES), hyperperfusion encephalopathy, and brain capillary leak syndrome. RPLS is becoming an increasingly recognized neurologic disorder. The exact mechanism by which this condition is
Paroxetine, citalopram, fluoxetine, and sertraline are among the SSRIs reported to be linked to RCVS in this case series. Duloxetine, although has a very similar pharmacological profile as the other SSRIs, differs from the class in being a potent reuptake inhibitor of norepinephrine as well [15]. We believe in our case, the use of duloxetine resulted in serotonin syndrome which led to the sequential development of hypertension, RCVS, and finally RPLS. To our knowledge, there has been no evidence in the literature to suggest the role of duloxetine in causing RCVS or RPLS.

The presenting clinical features of RPLS include headaches, seizures, visual disturbances including cortical blindness and visual hallucinations, and alterations in mental status including confusion and stupor [1, 9]. The headache can range from moderate to severe and may not respond to analgesics [9]. Seizures can be the initial presenting complaint as was the case in our patient as well [1]. Seizures are usually of the tonic-clonic type. Hypertension is accompanied with the neurologic symptoms most of the time and may precede the symptoms by a few hours [9].

The diagnosis of RPLS is established by neuroimaging. Typical findings are symmetrical white matter edema in the posterior cerebral hemispheres, particularly the parieto-occipital regions [8]. Involvement of cerebellum and brainstem is common. The calcarine and paramedian parts of the occipital lobe are usually spared which might help in differentiating from bilateral posterior cerebral infarctions [1]. Although depicted well on CT, the findings are best seen on MRI. The most common abnormalities seen on MRI are confluent areas of increased signal on proton density and T2-weighted images [8].

Early diagnosis and treatment is essential for the patients’ prognosis. Discontinuation of the offending agent and control of blood pressure are pivotal to reverse the fatal consequences [13]. Hypertension is a frequent feature in cases of RPLS and patients can improve dramatically with lowering of blood pressure [9]. Most seizures respond well to phenytoin and the drug can be tapered off quickly with reversal of neurological symptoms and imaging findings [9]. The prognosis of RPLS is good and symptoms are reversible if diagnosis is made promptly and remedial measures are taken.

References

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