Overactive Bladder Successfully Treated with Duloxetine in a Female Adolescent

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Overactive bladder (OAB) is defined as urgency, usually with frequency and nocturia, and with or without urge incontinence. Duloxetine, an antidepressant that inhibits reuptake of serotonin and norepinephrine, is indicated for the treatment of stress urinary incontinence in Europe. In this paper, we present a case of a 17-year-old female patient with OAB and depressive symptoms who was successfully treated with duloxetine. This case suggests duloxetine can be an option for patient with OAB, and it also highlights the need for further studies of duloxetine’s use in the treatment of OAB.

KEY WORDS: Duloxetine; Overactive urinary bladder; Treatment; Stress urinary incontinence; Depression.

INTRODUCTION

Overactive bladder (OAB) is a urological condition defined by a symptom complex, which includes urgency, usually with frequency and nocturia, and with or without urge incontinence.1) Frequency is generally defined as urinating more than eight times a day.

Duloxetine, a dual reuptake inhibitor of serotonin and norepinephrine, is approved for the treatment of major depressive disorder, generalized anxiety disorder, diabetic peripheral neuropathic pain, fibromyalgia, and chronic musculoskeletal pain in the US.2) In Europe, duloxetine is also indicated for the treatment of stress urinary incontinence in women.3,4) Although studies have suggested that duloxetine may be effective in treating OAB, its use in this condition is still not indicated.5,6) More importantly, the effectiveness of duloxetine in OAB patients with purely irritative symptoms (i.e., OAB without stress or urge incontinence) remains unknown. In this paper, we report a case of OAB without stress and urge incontinence complicated by depressive symptoms in an adolescent female that was successfully treated with duloxetine.

CASE

A 17-year-old female patient, a high school junior, with no known medical or urogynecological history, presented with urgency and frequency for 2 years. Her frequency was very severe; she had to urinate every 30 to 60 minutes during the day. She also exhibited fear of incontinence, although she had no actual incontinence. These urinary symptoms and related anxiety prevented her from concentrating in class, interfered with daily living, and caused her embarrassment. She stated, “When I get the urge, I must go right away. I can’t even get on a bus because I am afraid that I might get the urge while I am on it and embarrass myself.” Occasionally, she also had to wake up one to two times during the night because of nocturia.

Before the patient was referred to our psychiatric department, she was treated in our urological department for 5 months. Initial physical examination, urinalysis, and X-rays of the lower abdomen yielded no abnormal findings. Under the urologist’s care, she was prescribed solifenacin succinate (5 mg/day) for 3 weeks, oxybutynin chloride (5 mg/day) for 3 weeks, fesoterodine fumarate (8 mg/day) for 3 months, and propiverine hydrochloride (20 mg/day) for 4 weeks. However, her OAB symptoms remained unimproved, leading her to worry that the symptoms were not treatable. She could not sleep at night because of these worries. This anxiety later developed into depression, and she felt agitated and suicidal. Thus, she was referred to our psychiatric department.
Upon referral, she was treated with escitalopram (10 mg/day) for 5 weeks under impression of depression related to OAB. However, neither her depression nor her OAB showed any signs of improvement. She was then switched to fluoxetine (10 mg/day) for 2 weeks, but this was stopped because her insomnia worsened. Imipramine (20 mg/day) was then prescribed for 3 weeks, but her urinary symptoms worsened. Meanwhile, she became hopeless about her OAB and stopped visiting her urologist. She was then prescribed duloxetine (30 mg/day), and the dosage was escalated to 60 mg/day after 5 weeks of titration. Six weeks after duloxetine initiation, she described her urgency and frequency as 50% better. About a month later, she no longer had nocturia, and her micturition frequency has decreased to every 2 hours during the day, allowing her to concentrate during class. While she was taking duloxetine (60 mg/day), her depressive symptoms including insomnia and agitation also improved along with her alleviation of OAB symptoms over the next 6 months. Her urination frequency decreased to eight or fewer times a day, and she no longer had trouble with daily living or with riding the bus. She stated that she experienced no urinary urgency while taking the College Scholastic Ability Test, which lasted for more than 6 hours. Complaints of urgency and frequency did not recur and remained absent throughout treatment with duloxetine (60 mg/day). At the time of submission of this manuscript, she planned to attend college.

**DISCUSSION**

To the best of our knowledge, this is the first study reporting a case of a female adolescent having OAB and depression which was successfully treated with duloxetine. A previous study reported a case of detrusor overactivity successfully treated with duloxetine. The chief complaint of a 47-year-old female in the report was stress incontinence, which is a current indication for duloxetine. In contrast, the case reported here affected an adolescent with purely irritative symptoms without stress or urge incontinence. The fact that our patient’s OAB was refractory to three anticholinergic medications and three antidepressants, but resolved after duloxetine administration, is also noteworthy.

Studies have suggested that duloxetine improves urinary incontinence by inhibiting reuptake of serotonin and norepinephrine at the pre-synaptic neuron in Onuf's nucleus in the sacral spinal cord. Both alpha-receptors and serotonin 5-HT2 receptors in Onuf’s nucleus facilitate the storage reflex by contracting the urethral external sphincter muscle. Furthermore, results of an animal study suggested that duloxetine could increase vesical maximal capacity five-fold. Duloxetine’s ability to increase bladder capacity may be an important contributor to the improvement of urgency and frequency in our case.

We did not perform a urodynamic test, which could be an important limitation because it could have provided a more objective measure of severity of the OAB. We did not perform this test because it is usually indicated in patients with diagnosed or suspected neurological disease, and our patient had no evidence of such a condition. Another potential issue is that the safety and effectiveness of duloxetine in pediatric patients have not been established. Nonetheless, its use in children or adolescents is not contraindicated, and recommendations for duloxetine in such patients are to balance potential risks with clinical need. Finally, our patient’s urinary urgency and frequency may have been a symptom of anxiety, in which case, relief of urinary symptoms was simply a result of the drug’s anxiolytic effect. We believe reaching such a conclusion could overlook a patient’s disabling urinary symptoms. In line with our belief, studies not only showed association between OAB with anxiety, but also illustrated that people are inclined to consider anxiety symptoms more serious than OAB. Moreover, constant urinary urges led to anxiety in our patient, so we believed that the anxiety symptoms were secondary to her urgency and frequency.

Regardless of the exact mechanism of successful treatment, our report suggests that duloxetine could be a treatment option for OAB. It also highlights the need for further studies of duloxetine’s efficacy in the treatment OAB, especially in those with purely irritative symptoms.

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