Angioedema and Urticaria Associated with Fluoxetine in a Preadolescent Boy

Preadolesan Bir Erkek Çocukta Fluoksetinle İlişkili Anjiyoödem ve Ürtiker

Aslı Sürer Adanır1, Arif Önder2, Zehra Ece Soğucak3, Abdurrahman Erdem Başaran3, Ayşen Bingöl6, Esin Özatalay1

1Akdeniz University Faculty of Medicine, Department of Child and Adolescent Psychiatry, Antalya, Turkey
2Manisa Psychiatry Hospital, Clinic of Child and Adolescent Psychiatry, Manisa, Turkey
3Akdeniz University Faculty of Medicine, Department of Pediatrics, Antalya, Turkey
4Akdeniz University Faculty of Medicine, Department of Pediatric Allergy-Immunology, Antalya, Turkey

Abstract

Angioedema is defined as increased permeability and dilatation of the capillaries in the deep dermis or subcutaneous or submucosal tissues leading to localized swelling, often affecting the upper respiratory and gastrointestinal tracts. Angioedema may be mast cell (such as histamine)-mediated, bradykinin-mediated; or of unknown origin. Mast cell mediators are usually associated with urticaria and itch, whereas bradykinin has no association. Urticaria is defined as the vascular reaction of the upper dermis marked by itching and the transient appearance of raised patches that are redder or paler than the surrounding skin.1

Angiotensin-converting enzyme inhibitors (ACEIs), non-steroid anti-inflammatory drugs (NSAID), neuromuscular blockers and penicillin are the most common medications held responsible for angioedema,2 but it can also be observed as a rare but potentially life-threatening adverse effect of selective serotonin reuptake inhibitors (SSRIs). There are 4 reported cases of angioedema associated with SSRIs in the literature.3-6 Here, we report a 10-year-old boy who had urticaria associated with fluoxetine use, showed recovery after cessation of the drug, and manifested with urticaria and angioedema after re-prescription. In the literature, there are two previous case reports of angioedema with fluoxetine. One case of angioedema accompanied by urticaria and showed a flu-like presentation 2 days after ingestion of high-dose fluoxetine. The other case, reporting angioedema with 10 mg/day, was without urticaria and thought to be a pseudoallergic reaction. We assume that our case is unique in this respect for that angioedema developed with therapeutic doses and thought to be allergic.

Keywords: Adverse effect, angioedema, child, fluoxetine, urticaria

Introduction

Angioedema is defined as increased permeability and dilatation of the capillaries in the deep dermis or subcutaneous or submucosal tissues leading to localized swelling, often affecting the upper respiratory and gastrointestinal tracts. Increased vascular permeability may be mast cell (such as histamine)-mediated, bradykinin-mediated; or of unknown origin. Mast cell mediators are usually associated with urticaria and itch, whereas bradykinin has no association. Urticaria is defined as the vascular reaction of the upper dermis marked by itching and the transient appearance of raised patches that are redder or paler than the surrounding skin.1


Anahtar Kelimeler: Anjiyoödem, çocuk, fluoksetin, ürtiker, yan etki

Address for Correspondence/Yazılaşma Adresi: Aslı Sürer Adanır MD, Akdeniz University Faculty of Medicine, Department of Child and Adolescent Psychiatry, Antalya, Turkey
Phone: +90 242 249 67 97 E-mail: asladanir@hotmail.com ORCID ID: orcid.org/0000-0002-6223-756X
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fluoxetine use, showed recovery after the cessation of the drug, and manifested with urticaria and angioedema after the re-prescription.

Case
A 10-year-old boy presented to our pediatric emergency unit with diffuse itchy skin lesions (Figure 1) and swollen lips and tongue (Figure 2). His parents reported that he was on fluoxetine 10 mg treatment for 12 days for his anxiety symptoms, and he had mild urticarial lesions for 2 days. Within a few hours, the lesions had spread to all of his body and he started to have difficulty in talking and breathing because of swollen lips and tongue. Examinations and questioning of the patient and information obtained from the parents ruled out food or pollen allergy, insect bites, physical exercise, NSAID use or any other etiological agents. The parents did not report any novel stressors or infection symptoms prior to the onset of angioedema. He did not take any food supplements or medication except for fluoxetine. Hereditary angioedema (HAE) was ruled out by a negative family history and normal C4 and C1 inhibitor levels.

Upon questioning and medical reports, it was learned that he was brought to the emergency department once more with diffuse urticaria two and a half years ago, while he had been taking fluoxetine 10 mg for fifteen days. Three days before presenting to the emergency department, the family had taken him to their primary care physician, and oral pheniramine had been prescribed, but he did not benefit from the treatment. His lesions had been attributed to fluoxetine treatment and prednisolone and pheniramine were started, and cessation of fluoxetine was recommended. The lesions healed up in 2 days after then.

His lesions were attributed to fluoxetine treatment once more and intravenous prednisolone and pheniramine were given and fluoxetine was stopped. Angioedema regressed quickly after medication and complete recovery of the lesions was observed within 2 days. The patient is on sertraline treatment without any side effects now.

The patient’s mother provided written informed consent for publication of this case report and accompanying images.

Discussion
Fluoxetine is usually a well-tolerated drug in children and adolescents. The most frequent adverse events are sleep problems, gastrointestinal symptoms and headache. Urticaria, skin eruptions and rarely angioedema are also reported by the manufacturer. There are also 4 case reports of angioedema associated with SSRIs in the literature. There are three main forms of angioedema: extrinsic factor-induced angioedema, angioedema with C1-INH deficiency and idiopathic. The first form includes angioedema associated with allergic and non-allergic reactions due to various antigens, such as drugs, pathogens, foods, animals, venom, latex, etc. The second form is associated with C1-INH deficiency and divided into two subtypes, as HAE and acquired angioedema. The causes of the last form, idiopathic angioedema, are unknown. Drug-induced angioedema

Figure 1. Urticarial lesions on the front of the leg and foot

Figure 2. Swollen lips, indicating angioedema
is classified in the extrinsic factor-induced angioedema and divided into three main categories depending on the mechanism. The first group is IgE-mediated immediate hypersensitivity reactions, especially associated with iodinated contrast media, beta-lactam antibiotics, neuromuscular blocking agents, quinolones and pyrazolones. The second includes the adverse effects of NSAIDs due to inhibition of cyclooxygenase resulting in major alterations in arachidonic acid metabolism and is generally non-allergic. Thirdly, ACEI-induced angioedema, a reaction, in which the inhibition of the degradation of bradykinin is observed, resulting in angioedema, but not urticaria. To differentiate a kinin-dependent angioedema, from the IgE-mediated one and NSAID intolerance, it is useful to determine as if angioedema is accompanied by urticaria or not. Here, we report a case of angioedema in a preadolescent, thought to be associated with fluoxetine. The temporal correlation of the occurrence of urticaria with the use of fluoxetine in the absence of comorbid illness or concurrent medications followed by its disappearance with discontinuation and re-occurrence (this time with angioedema) with the repeated use of it strongly suggested that the case was associated with fluoxetine. An evaluation with the Naranjo algorithm revealed a score of 8, denoting a probable adverse effect. Due to the lack of patient and family history for similar reactions and normal C4 and C1 inhibitor levels, the diagnosis of HAE was excluded. We considered it to be an allergic reaction, as it was accompanied by urticaria. We think that he did not have angioedema at the first time he had used fluoxetine because of oral pheniramine he used for three days.

Fluoxetine is rarely associated with hypersensitivity reactions. In one case, it was reported to be related with a severe systemic hypersensitivity reaction including drug rash, eosinophilia, and systemic symptoms in a 4-year-old girl. There is only one other case of angioedema accompanying urticaria and flu-like symptoms developing 2 days after ingestion of high dose fluoxetine. The other case of angioedema attributed to treatment fluoxetine with 10 mg/day was without urticaria and thought to be a pseudoallergic reaction. Our case is unique in this respect for that angioedema developed with therapeutic doses and thought to be an allergic reaction. Regardless of the cause, clinicians should be aware of such rare but potentially life-threatening adverse effects of SSRIs and monitor patients closely. Parent education is also imperative.

Ethics

Ethics Committee Approval: None, as it is a case report.

Informed Consent: The parents of patient signed written informed consent form.

Peer-review: Externally peer-reviewed.

Authorship Contributions


Conflict of Interest: No conflict of interest was declared by the authors.

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