

CASE REPORT

Transient normoprolactinemic galactorrhea induced by fluoxetine

Fluoksetine baęlı, geici normoprolaktinematik galaktore

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ABSTRACT

Hormonal side effects of antidepressants are infrequent. Galactorrhea is rarely reported among antidepressant related side effects. Antidepressants can directly stimulate postsynaptic 5- Hydroxytryptamine (5-HT) receptors in the hypothalamus or indirectly inhibit the tuberoinfundibular dopaminergic neurons through 5-HT, which may increase prolactin levels and later cause galactorrhea. However, galactorrhea may develop despite normal prolactin levels during antidepressant treatment. We present a case of normoprolactinemic galactorrhea in a woman, related with fluoxetine treatment. This report highlights the presence of unidentified mechanisms of selective serotonin reuptake inhibitor induced galactorrhea. *J Clin Exp Invest* 2013; 4 (1): 105-106

Key words: Fluoxetine, galactorrhea, side effect

INTRODUCTION

Several drugs can cause galactorrhea, and it needs to be differentiated from other local or neuroendocrinological causes. All conventional antipsychotic drugs block D2 receptors on lactotroph cells and thus remove the main inhibitory influence on prolactin secretion.¹ Tricyclic antidepressants and selective serotonin reuptake inhibitors (SSRI) are less frequent causes.^{2,3} There is evidence that serotonin may stimulate prolactin release directly via postsynaptic 5- Hydroxytryptamine (5-HT) receptors in the hypothalamus,⁴ or indirectly via 5-HT mediated inhibition of tuberoinfundibular dopaminergic neurons.⁵ However, galactorrhea due to antidepressants is not consistently associated with elevated prolactin levels, which may suggest still unexplained mechanisms of antidepressant-induced galactorrhea.^{6,7}

Herein, we report a case of euprolactinemic galactorrhea in a woman with generalized anxiety disorder while on treatment with fluoxetine.

ÖZET

Antidepresanlara baęlı hormonal yan etkiler sık görülmektedir. Galaktore, antidepresanlara baęlı yan etkiler arasında nadiren bildirilmiştir. Antidepresanlar, hipotalamustaki postsinaptik 5-Hidroksitriptamin (5-HT) reseptörlerini doğrudan uyararak veya tuberoinfundibuler dopaminergic nöronları 5-HT üzerinden dolaylı olarak inhibe ederek prolaktin seviyelerinde artışa ve sonuçta galaktoreye neden olabilirler. Ancak antidepresan tedavisi sırasında normal prolaktin seviyelerine rağmen galaktore gelişebilir. Bu çalışmada fluoksetin tedavisi ile ilişkili normoprolaktinematik galaktore gelişen bir kadın olgu sunulacaktır. Bu olgu sunumu ile selektif serotonin geri alım inhibitörlerine baęlı galaktorenin tanımlanmamış mekanizmalarının varlığı üzerine vurgu yapılacaktır.

Anahtar kelimeler: Fluoksetin, galaktore, yan etki

CASE

A 29-year-old woman with two children visited a psychiatric outpatient clinic with complaints of excessive and uncontrollable worry about minor life events, feeling restlessness, irritability, muscle tension, tiring easily, and poor sleep that started 8 months prior to admission. She had no history of endocrine or reproductive pathology or psychiatric problems. She was diagnosed as generalized anxiety disorder according to DSM-IV criteria and was started on fluoxetine 20 mg per day. After 4 weeks of medication, her complaints declined. However, she developed unilateral galactorrhea (the nonpuerperal discharge of milk-containing fluid from the breast). She had no history of galactorrhea, thyroid disease, and polycystic ovary disease. She first noticed the discharge on treatment day 21 and described it as white-creamy and from right nipple. She did not notice any bloody, greenish, or foul-smelling discharge nor report sexual dysfunction. She consulted her gynecologist, who recommend-

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ed mammogram and breast ultrasonography. The pregnancy test was negative. The results of these tests and breast examination were normal. Serum prolactin level on treatment day 28 was 18.18 ng/mL (reference range: 2.5-29 ng/mL). Because her galactorrhea developed after initiations of her medication with fluoxetine, her medication was discontinued. Bupirone 5 mg/day treatment was started and gradually raised to 20 mg/d. Eight days after stopping fluoxetine, the patient reported reduction and cessation of galactorrhea. During 3 months of follow-up, the patient maintained well on bupirone and there was no reemergence of galactorrhea.

DISCUSSION

Galactorrhea caused by the use of fluoxetine has been reported earlier and the commonly perceived cause is hyperprolactinemia.⁸ Fluoxetine has been shown to potentiate elevation of prolactin levels inducing other stimuli, including insulin, fenfluramine, and 5-HT.⁹ However, hyperprolactinemia is not the only mechanism responsible for the development of SSRI-induced galactorrhea.^{6,7} The exact mechanism of galactorrhea remains unknown in many cases.

Our patient developed galactorrhea without hyperprolactinemia after beginning fluoxetine therapy. The strict temporal relationship between the use of the drug and the onset of galactorrhea, as well as the resolution once treatment was discontinued, suggests a causal link between the two phenomena. Although unilateral galactorrhea induced by psychotropic medications is rare,¹⁰ our patient developed unilateral galactorrhea with normal mammogram and breast ultrasonography, indicating a complex relationship between antidepressant use and galactorrhea.

To the best of our knowledge, we are the first to report an association with fluoxetine use and galac-

torrhea without elevated prolactin level. Clinicians should consider fluoxetine as a possible cause of galactorrhea even with normal prolactin levels. Future research should investigate the exact mechanisms of antidepressant-induced normoprolactinemic galactorrhea.

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