Case Report

Giant prolactinoma case with side effects due to cabergoline

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Abstract

Dopamine agonists should usually be the first treatment for patients with prolactinomas of all sizes, because these drugs decrease serum prolactin concentrations and decrease the size of most lactotroph adenomas. Cabergoline is preferred first. When cabergoline develops side effects bromokriptin is switched. Cabergoline is more efficient than bromocriptine. Its side effect profile more favorable than bromocriptine. Cabergoline is an ergot dopamine agonist that is administered once or twice a week. Cabergoline can have rarely serious psychiatric adverse effects, including psychosis, impulse control disorders, dyskinesia, pulmonary fibrosis and valvular heart disease.

Prolactinomas are the most common pituitary tumors, 93%-95% of the cases are microadenomas. Macroprolactinomas larger than 40 mm, known as giant prolactinomas, are exceptionally rare, accounting for 0.5% - 4% of all prolactin-hypersecreting adenomas.

In our case, after the 7 x 6 cm giant macroprolactinoma operated we report the manic episode occurring during the second week of cabergoline treatment. In the treatment of patients with prolactinoma, cabergoline is a first choice drug because it has a better tolerance profile and is more effective, however, bromocriptine can be switched to when drug resistance or side effects develop to cabergoline.

Introduction

Prolactinomas are the most common pituitary tumors, often seen as microadenomas, and treatment is usually medical. Treatment of patients with lactotroph macroadenomas, no matter how large or how severe the neurologic symptoms, should also be initiated with a dopamine agonist, starting with cabergoline [1]. Patients whose macroadenomas are largely cystic should also be treated initially with a dopamine agonist since this treatment shrinks most of these [2]. Cabergoline, a dopamine agonist is generally used in the treatment of hyperprolactinemia, Parkinson’s disease, restless leg syndrome, and antipsychotic-induced prolactin elevation. It is generally well tolerated as compared to other dopamine agonist due to its more selective D2 receptor agonistic effect [3]. However, it can cause certain psychiatric side effects including psychosis, depression, impulse control disorders, and nightmares [4].

Prolactin (PRL)-secreting pituitary adenomas (PRLomas) are the most common pituitary secreting tumors, accounting for 32% - 45% of all pituitary tumors [5-7]. Prolactinomas are most often microadenomas (< 10 mm in diameter) and are diagnosed more frequently in women, who usually present with the hormonal sequelae of hyperprolactinemia including galactorrhea and oligomenorrhea. In men, macroprolactinomas (> 10 mm in diameter) are more commonly detected, and it has been hypothesized that the greater subtlety of male hormonal symptoms of hyperprolactinemia, gynecomastia, decreased libido, and impotence may account for the lower observed frequency and the larger size of clinically evident prolactinomas in men [8]. Giant prolactinomas are pituitary adenomas larger than 4 cm in size presenting very high serum prolactin levels (typically above 1000 ng/mL) and no concomitant growth hormone (GH) or adrenocorticotropic hormone (ACTH) hypersecretion. They represent less than 5% of all prolactinomas [9,10].

Here we present a case of a 33-year-old male who had 7x6 cm macroadenom. Occurred manic episode, personality changes and hallucinations, following the started cabergoline treatment after surgery, thereon bromocriptine therapy was started.

Case report

A 33-year-old male patient attended to ophthalmologist with a bilateral visual loss for four-five years.

He reported no history of galactorrhea, gynecomastia, but have headaches have and decreased libido. Then he
referred to endocrinologist. In laboratory examinations, hyperprolactinemia (> 200 ng/mL) and hypogonadotropic hypogonadism, were detected (Figure 1 and Table 1).

He was started on cabergoline 0.25 mg twice weekly, prednisone tablet 7.5 mg/daily and testosterone 250 mg IM injection every 3 weeks. Patient was hospitalized for treatment. After a week of cabergoline treatment patient had an epileptic attack due to apoplexi. The patient was urgently opereted. Postoperative patient was followed and discharged. Pathologic examination showed that; the tumor cells were positive for prolactin and luteinizing hormone (LH), follicle stimulating hormone (FSH), growth hormone (GH), thyroid stimulating hormone (TSH) were negative. Patient was brought to the emergency department by his relatives hallucinations, personality changes and manic episodes. These complaints seen in the patient were related to the side effect of the cabergolin. Therefore, bromocriptin treatment was started 1.25 mg/day and increased to 5 mg/day on this patient. There was no side effect with this treatment.

Discussion

Mesocortical and mesolimbic D2 and D3 receptors are thought to be involved in mood and behavior, and dopamine agonists can trigger psychosis or mania[11-13]. Impulse control disorders (ICDs), including problem gambling, hypersexuality, compulsive eating and compulsive shopping, may be mediated by D3 receptors in the mesolimbic system [12,13]. D2 receptor agonistic effect of cabergoline is responsible for manic episode. Initially high dose (0.5 mg per day) of cabergoline is associated with risk of psychosis [14]. However, no literature is available regarding relation between very low dose of cabergoline with giant macroprolactinoma.

In general, medically treated prolactinomas have a high remission rate of 90% [6]. When they are large and extend outside the sellar borders, the rate of prolactin normalization decreases to 70%-80% for macroprolactinomas [6] and 60%-68% for giant prolactinomas [9,10].

Transsphenoidal or transcral resection may be offered to the subset of patients with giant prolactinoma for whom even high-dose medical therapy fails to achieve tumor size and hormonal control [15], and patients with apoplexy or complications of pharmaceutical treatment (CSF leak). The reported rate of surgery following first-line dopamine agonist treatment is nearly 15% [9]. Severe chiasmal pressure might also lead to a decision to operate, although even severe optic damage and significant visual field defects can improve with medical treatment alone [9,10]. However, surgery is rarely curative. Most patients will not achieve hormonal remission postoperatively [6,9], will remain with a significant adenoma residue, and will require continuous medical treatment. Surgical resection may also be hampered by intratumoral fibrosis, which may develop following long-term dopamine agonist management [16,17]. Moreover, surgery in patients with gigantic tumors can be associated with complications such as diabetes insipidus, mental deterioration, CSF leak, meningitis, worsening vision, and even perioperative mortality [18].

Although cabergoline is highly effective in hyperprolactinemia treatment, it may induce hypomanic/manic episodes, pituitary apoplexi, personality changes and hallucinations such as in this case. Risk-benefit ratio should be analyzed with prospective controlled studies in future and cabergoline must be used carefully. Alternatively other drugs (e.g., bromocriptin, quinagolide) should be kept in mind. But there are more than other side effect profile (nausea, vomiting, dizzi-ness) and should be taken every day. Cabergoline is an effective drug but it should be kept in mind that cabergoline may develop side effects even at very low doses.

Giant prolactinomas are generally benign but they may be aggressive and invasive, disrupting nearby structures and higher serum prolactin levels. They show a fairly good response to medical treatment with dopamine agonists, combined with surgery in some patients and radiotherapy if required.
Giant prolactinoma case with side effects due to cabergoline

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