Macroprolactinoma appearing after resolution of hydrocephalus

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Summary

The role of mechanical forces influencing the growth of a pituitary adenoma is poorly understood. In this paper we report the case of a young man with hyperprolactinaemia and an empty sella secondary to hydrocephalia, who developed a macroprolactinoma following the relief of high intraventricular pressure.

Learning points:

• The volume of a pituitary tumour may be influenced not only by molecular but also by local mechanical factors.
• Intratumoural pressure, resistance of the sellar diaphragm and intracranial liquid pressure may play a role in the final size of a pituitary adenoma.
• The presence of hydrocephalus may hide a pituitary macroadenoma.

Background

Factors controlling pituitary adenoma growth and aggressiveness are still poorly understood. The significance of molecular alterations acting through changes in the regulatory mechanisms of the cell cycle and subsequently in the growth pattern of pituitary adenomas is currently being investigated (1). However, to the best of our knowledge, the role of physical forces as conditioning factors in tumour growth has not yet been considered. In this study, we report a clinical case suggesting the possibility of such a role in a patient with a pituitary prolactinoma.

Case presentation

A 24-year-old man first consulted because of a progressive loss of libido and erectile dysfunction that had started at the age of 18. He had also had slow mentation and ill-defined psychomotor troubles since the age of 15. Physical examination was normal with the exception of scarce facial beard which he shaved sporadically; his weight was 79 kg, height, 172 cm and blood pressure, 120/70 mmHg.

The ophthalmologic evaluation showed normal visual acuity (10/10 in both eyes) but a bilateral papilledema.

Investigations

Adrenal and thyroid function tests were within the range of normalcy. Testosterone level was 3.4 ng/mL (normal value (nv): 3.5–8.0). The main finding was a serum prolactin (PRL) concentration of 880 and 1000 ng/mL, on two separate occasions suggesting the diagnosis of a prolactinoma. However, instead of a pituitary adenoma, MRI revealed a very important ventricular dilatation and a partially empty sella with herniation of the suprasellar cisternae, the third ventricle and the optic chiasm into the pituitary fossae; idiopathic aqueductal stenosis was diagnosed (Fig. 1).

The patient was hospitalized and 24-h monitoring of intracranial pressure (IP) showed a mean figure of 16 mmHg (normal values <13 mmHg) (2). His clinical condition deteriorated with the loss of communication, disorientation and prostration.
Treatment

After two unsuccessful attempts to place shunting devices (a medium pressure valve and a flow-dependent valve respectively), a ventricular-peritoneal shunt using a mean-pressure valve was performed. A progressive improvement began on the sixth postoperative day with recovery of oral language, and continued until the total normalization of all neuropsychological functions. At that time, IP was 7 mmHg, MRI showed a ventricular dilation reduction and a 13 mm image compatible with pituitary macroadenoma was now evident (Fig. 2); serum PRL was 953 ng/mL. After three months, an additional reduction of the ventriculomegaly was observed (Fig. 3) but serum PRL remained high at 400–500 ng/mL, despite treatment with increasing doses of bromocryptine (up to 25 mg/day). This treatment had to be discontinued because of the development of psychotic signs which subsided after a few days. A trial with cabergoline was indicated and then the patient was operated on by the transsphenoidal approach to resect a prolactinoma confirmed by the pathologist with immunomarcation showing strong positivity for PRL.

Outcome and follow-up

In the following months, serum PRL fell to between 90 and 200 ng/mL. After discharge, he was treated for panhypopituitarism with levothyroxine 100 µg/day, hydrocortisone 20 mg/day, testosterone gel 1 application/day. He was lost to monitoring for a very long period but reappeared at the beginning of 2016, almost 20 years after his first consultation. He was clinically stable under treatment and a physical exam conducted on him was normal. MRI showed no pituitary tumour but a moderate dilation of the ventricular system (Fig. 4).
the ventricular system (Fig. 4). Hormonal lab showed undetectable serum LH, FSH, GH and cortisol; prolactin level was 61 ng/mL. The long-term patient’s evolution has been satisfactory and he lives and works normally.

Discussion

Around 80% of prolactinomas are microadenomas, though macroadenomas are frequently observed in young male patients (3). Normal mean intrasellar pressure is not known but it is unlikely to exceed normal intracranial pressure (ICP). An ICP value below 13 mmHg characterizes normal CSF circulation; above 18 mmHg the CSF circulation is clearly disturbed; between 13 mmHg and 18 mmHg there is a grey zone (2). On the other hand, it is known that pituitary intratumoral pressure can be variable, depending on the size of the tumour. In a two-centre study, intrasellar pituitary tissue pressure (ISP) was measured during transsphenoidal surgery in 107 patients with pituitary adenomas or intrasellar cysts; ‘normal’ ISP in patients with microadenomas less than 5 mm diameter or a partial empty sella was $12 \pm 3$ mmHg. Raised ISP of more than 15 mmHg was found in 75% of the cases (4). On the other hand, little is known about the possible influence of variable resistance of the anatomical surrounding structures on the growth of pituitary tumours. The sellar diaphragm is composed of two layers of dura mater forming the upper limit of the sella turcica, but its resistance to pressure on both sides is difficult to define; it could be because anatomically weakened resistance is involved in the pathogenesis of primary empty sella (5). In the same way, individual variation in the resistance of the sellar diaphragm can partially explain the greater or smaller suprasellar expansion of pituitary adenomas. There is also a remarkable variation in the morphology of the diaphragm opening which could explain the growth of pituitary tumours towards the cavernous sinus or towards the suprasellar region (6).

Our patient presented a mean ICP of around 16 mmHg accompanied by a clear clinical intracranial hypertensive syndrome and imaging of a huge hydrocephalus which prompted neurological treatment. Interestingly, in the present case higher than normal IP could have impeded tumour expansion, without influencing PRL secretion. We assumed that the pituitary adenoma was not the cause of hydrocephalus because of its moderate size. Besides, the tumour expansion took place shortly after intracranial pressure decrease, even before normalization of the ventricular dilation, leading to the clear visualization of a typical macroadenoma image. The endocrine clinical findings in this case are not different from those shown in the usual cases of prolactinoma, but the main interest of the observation is the unusual form of presentation which allowed us to speculate on the role of mechanical factors influencing the size of a pituitary adenoma.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Patient consent

Written informed consent has been obtained from the patient for the publication of the case report.

Author contribution statement

Dr O D Bruno is a consultant endocrinologist and professor of medicine who was in charge of the patient together with Dr G Isaac. Dr Fernandez Pisani and Prof. A Basso were the neurosurgeons responsible for the surgical procedures. All the authors reviewed and revised the present manuscript.

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