disease, as there is no evidence of a decline in lung function following ABP, while reduced function has been observed following the use of other substances.4,5 The technique consists in the instillation through a chest tube of 1–2 mL/kg of blood previously extracted from the patient adhering to strict aseptic technique. After the blood is instilled, the cavity is rinsed with 10 mL of physiological saline solution and the chest tube is suspended 60 cm overhead for 30 min to prevent drainage of the instilled blood while allowing air to escape the chest.2,4,5 We recommend changing positions to facilitate a better distribution of the blood.

While the exact closure mechanism is not understood, it has been postulated that the blood acts as a patch that directly seals the defect, followed by an inflammatory response in the pleural cavity.1 In the case presented here, treatment with vinblastine and corticosteroids promoted the persistence of pneumothorax, making the patch-like effect all the more relevant.

The most frequent complications of ABP are fever, pneumothorax and empyema. Unlike other techniques, it is painless, does not require sedation or analgesia, and can be repeated. We did not observe any of the previously described complications in our patient.

Autologous blood patch pleurodesis is a safe, inexpensive and efficacious treatment for persistent air leak. Larger studies need to be conducted in the paediatric population prior to recommending its routine use.

Acknowledgments

We wish to thank the nursing staff of the PICU and the paediatric oncology ward.

Prolactinomas in a paediatric population

Prolactinomas en la población pediátrica

Dear Editor:

Case 1: female patient aged 15 years that sought care at her assigned health care centre for irregular menstruation. There was no personal or family history of interest, and the patient was not taking any medication. Her age of menarche was 11.5 years.

The patient was referred to the paediatric endocrinology unit after testing revealed an elevated prolactin level (198 ng/mL; normal value range, 8.5–26.5 ng/mL). During the structured interview, the patient reported one year of intermittent blurry vision and frontal headache of three months’ duration accompanied by light unilateral galactorrhea. The findings of the physical examination were: height, 169.5 cm (standard deviation [SD], +1.1); weight, 53.3 kg; body mass index (BMI), 19.29 kg/m² (SD, –0.79). The patient was at Tanner stage V of pubertal development with proportionate features. There was no galactorrhea, hirsutism or acne. The complete blood count and blood chemistry panel were normal, with the most salient finding being the serum prolactin level (205.6 ng/mL). Magnetic resonance imaging (MRI) revealed an enlarged pituitary gland with a superior border that was convex towards the midline and signal hyperintensity suggestive of subacute haemorrhage (Fig. 1). The pituitary gland extended into and obliterated the suprasellar cistern, exerting a mass effect on the optic chiasm. The ophthalmological examination evinced mild hyperopia, and the visual field test and eye fundus examination were normal. The patient was given a diagnosis of prolactinoma with subclinical apoplexy based on these findings, and started treatment with cabergoline at a dose of 0.25 mg the first week, followed by a maintenance dose of 0.50 mg a week. One month after treatment initiation, the follow-up MRI scan showed an adenoma measuring 10 mm × 8 mm (Fig. 2), values that may have been overestimated due to haemorrhagic features, and the serum prolactin level was 12.6 ng/mL. At present, the patient has regular menstrual periods, normal vision without headache and no galactorrhea.

Case 2: male patient aged 14 years that was being followed up for obesity and tall height with advanced bone age. The assessment of the patient’s headaches included performance of an MRI scan that revealed a 10 mm × 10 mm nodule in the left region of the pituitary gland, with remodelling of the angle of the sella turcica with no extension to the cavernous sinus compatible with adenoma (at the

References


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Patients in the literature, and most of the evidence consists in the description of isolated cases. Patients may remain asymptomatic or have very mild symptoms compared to those of pituitary apoplexy. There is no correlation between prolactin levels and the duration or severity of symptoms. There is also no evidence of a relationship between tumour size and the presence of haemorrhagic processes. In children, development of subclinical pituitary apoplexy is more frequent due to the decreased susceptibility to infarction of the tumour and a greater resistance to haemorrhage than adults, and is probably underdiagnosed.

As for treatment, cases of subclinical apoplexy, especially prolactin-producing adenomas, can be managed conservatively. Treatment with dopamine agonists can control prolactin levels and reduce tumour size significantly. In cases of hyperprolactinaemia secondary to prolactinoma, medication is recommended as the first-line treatment for both microadenomas and macroadenomas regardless of age. The primary goal of dopamine agonist therapy is to improve sex-gland and neurologic functioning. The followup requires close monitoring of the patient’s clinical manifestations and blood chemistry, and treatment may be discontinued after two years if prolactin levels are no longer elevated and there is no visible tumour remnant on MRI. If treatment is discontinued, serum prolactin should be measured every three months in the first year and annually thereafter, and MRI performed if serum prolactin levels increased past the normal range.

References


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