

Evaluation of Clinical Features and Treatment Response of Cases with Hyperprolactinemia

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Objective

In this study, we planned to evaluate the patients with hyperprolactinemia etiology, clinical features and treatment responses

Material and Method

We evaluated retrospectively the features, clinical follow-up data and treatment responses of the patients with hyperprolactinemia in our pediatric endocrinology clinic between 01.01.2012-31.12.2018.

Results

- Thirty-one patients with hyperprolactinemia underwent follow-up in a seven-year period.
- The mean age of these cases was 15.7 ± 1.5 years.
- 27 cases (87.1%) were female.
- The most common complaints in girls were menstrual irregularity / amenorrhea (n = 14) and galactorrhea (n = 8). **Table 1**
- In terms of etiology, microadenomas were observed in 9, macroadenomas in 4, idiopathic hyperprolactinemia in 12, and drug-induced hyperprolactinemia in 4 cases.
- The mean prolactin level in the whole group was 104.7 ± 145.4 (27.8-813) ng/ml.
- According to the etiology, the mean prolactin level was 43,6 ng/ml in drug-induced hyperprolactinemia, 137,8 ng/ml in microprolactinoma group, 285 ng / ml in macroprolactinoma group, idiopathic hyperprolactinemia group was 49.6 ng/ml.
- **All cases with prolactin level above 100 ng/ml were diagnosed as adenoma** (3 macroadenoma and 5 microadenoma).
- Medical treatment was performed in 24 patients (23 cases with cabergoline, 1 case with bromocriptine) and surgery was performed in 3 cases (1 patient with Rathke cleft cyst, 2 cases with macroadenoma).
- Seven patients were followed without treatment.
- The mean cabergoline dose was 0.52 ± 0.21 (0.25-1) mg/week.
- **The mean time to normalization of prolactin levels was 2.6 ± 3 (1-12) months.**
- **There were no side effects related to cabergoline.**
- Postoperative prolactin levels did not return to normal in 2 patients with hyperprolactinemia due to macroprolactinoma and Rathke cleft cyst, and the need for medical treatment continued.
- Multiple pituitary hormone deficiency (gonadotropin, TSH, ACTH) was diagnosed in one patient with macroprolactinoma. The patient's cabergoline treatment (1 mg / week) and prolactin levels returned to normal in the first month. At the eighth month the adenoma shrank almost completely.

Table 1. complaints in girls

Complaints	n=27 (%)
Menstrual irregularity / amenorrhea	14(51,8)
Galactorrhea	8 (29)
Short stature	1 (3,7)
Delayed puberty	1 (3,7)
Hirsutism	1 (3,7)
Anti-dopaminergic drug	2 (7,4)

Table 2. complaints in boys

Complaints	n=4 (%)
Gynaecomastia	1 (25)
Galactorrhea	1 (25)
Short stature	1 (25)
Headache	1 (25)

Table 3. etiology, characteristics and treatment of hyperprolactinemia cases

	n	Sex	Age (year)	Prolactin (ng/ml)	Treatment
Microadenoma	9	6/3	$15,4 \pm 1,5$	137 ± 8 (43-308)	Medical 9
Macroadenoma	4	4/0	$16,5 \pm 1,3$	285 ± 353 (75-813)	Medical 4 Surgery 2
anti-dopaminergic drug	4	4/0	$15,6 \pm 0,9$ (15-17)	$43 \pm 5,4$ (35,7-48)	Medical 1 Untreated 3
Rathke cleft cyst	1	1/0	13	67	Medical+ Surgery
Macroprolactinemia	1	1/0	17,5	29	Untreated
Idiopathic hyperprolactinemia	12	11/1	$15,7 \pm 1,6$ (12-18)	49 ± 19 (28-88)	Medical 9 Untreated 3
Total	31	27/4	$15,7 \pm 1,5$ (12-18)	104 ± 145 (28-813)	Medical 24 Surgery 3 Untreated 7

Conclusion

The cases diagnosed as hyperprolactinemia were mostly adolescent girls. The most common cause was the pituitary adenoma (42%) and 64% of these adenomas were microadenomas. All cases with prolactin level > 100 ng / ml were diagnosed as adenomas. In patients with hyperprolactinoma, a good response to medical treatment including those due to adenoma, and a small number of cases required surgical treatment.