

# A CASE OF CUSHING'S SYNDROME DUE TO ADRENOCORTICAL ADENOMA WITH PUBARCHE AND OBESITY

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## BACKGROUND

Adrenocortical tumors in childhood represent very rare about 0.2% of all pediatric malignancies. Cushing's syndrome (CS) is characterized by clinical features caused by autonomous excessive glucocorticoid production from adrenal cortex. In ACTH-independent CS, the most common cause is unilateral cortisol-producing adrenocortical adenoma.



Figure 1. Patient; moon facies, buffalo hump, facial plethora with acne on her forehead and cheeks

## PATIENT

Fifteen months old girl was admitted with gradually gain weight, pubarche, and acne of the face since five months. There was no history of using drugs which contain steroid. Length was 74.4 cm (-1.31 SDS), weigh was 13.2 (2.16 SDS), moon facies, buffalo hump, facial plethora with acne on her forehead and cheeks. Her pubic stage was 3. In laboratory, glucose (fasting) 93 mg/dL, insulin (fasting) 24.99  $\mu$ IU/mL, total testosterone 76 ng/dl, cortisol (morning) 37.69 mg/dL, ACTH (morning) 5.04 pg/ml, cortisol (evening) 30.4  $\mu$ g/dL, ACTH (evening) 5.32 pg/ml, DHEA-SO<sub>4</sub> 238 pg/mL, sodium 138 mmol/L, potassium 4.53 mmol/L, WBC 20.8  $10^3$ /mL, RBC 5.05  $10^6$ /mL, hemoglobin 15.3 g/dL, and platelet 391  $10^3$ /mL. After evening dexamethasone implementation, cortisol (morning) was 29.36  $\mu$ g/dL. In magnetic resonance imaging identified pronounced thickening and nodularity in lateral crus of left adrenal gland.

## RESULTS

Patient was diagnosed with Cushing's syndrome due to adrenocortical tumor and administered laparoscopic left adrenalectomy. The removed mass was consistent with adrenal adenoma histologically. Postoperative fourth day control Cortisol (morning) 4.73 mg/dL, ACTH 18.7 pg/ml, and DHE-SO<sub>4</sub> <15.0 pg/mL.

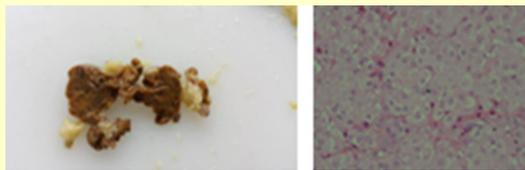


Figure 2. Adrenal adenoma size 5.5x3.5 cm, weight. 13 gr. The lesions consisting of large eosinophilic cytoplasm (H & E, x400)

## CONCLUSIONS

In adrenocortical tumors, distinguishing to adrenocortical cancer and adenoma is difficult. Herein we represent a rare case of CS because of adrenocortical adenoma with clinical features of excessive glucocorticoid production.

