

LAPAROSCOPIC BILATERAL ADRENALECTOMY

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Anatomy of adrenal gland

- Weight = 4-6 gm. (each)
- Size = 4-5 cm. x 2-3 cm. x 0.5-1 cm. (each) Shape = pyramidal (right), crescent (left)
- Color = golden orange, mustard
- Consistency = fine granular texture , friable Location = T11, subdiphragmatic







Laparoscopic Gold standard Procedures

Cholecystectomy
 Adrenalectomy

History Backgroud

1992 Joseph Petelin (France) Michael Gagner et al (Canada) Higashara E. et al (Japan)

Laparoscopic Approachs

- **1. Transperitoneal anterior approach**
- 2. Posterior retroperitoneal approach
- 3. Transperitoneal lateral (flank) approach



Indications of Laparoscopic Adrenarectomy

1. Aldosteronoma

- 2. Cushing syndrome
 2.1 Cortisol-producing adenoma
 2.2 Primary adrenal hyperplasia
 2.3 Failed treatment of ACTH
 Dependent Cushing's
- 3. Pheochromocytoma
- 4. Nonfunctioning cortical adenoma (incidentaloma) > 4 cm.
- 5. Small vinilizing adenoma
- 6. Adrenal metastasis
- 7. Miscellaneous

(myelolipoma, adrenal cyst, ganglioneuroma)

My own experience in Laparoscopic Adrenalectomy Four left adrenalectomy (three aldosteronoma, one Cushing) One right adrenalectomy (aldosteronoma) One bilateral adrenalectomy (Cushing)

Cushing's Syndrome

- Obesity, Peculiar fat deposition
- Amenorrhea, Impotence
- Purple striae
- Hypertension, Diabetes
- Other features that constitute the syndrome :-

Moon facies, Hirsutism, and Acne





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The causes of Cushing's syndrome Exogenous (most common cause)

- Administration of steroids
- Endogenous (rare, 5-10 per million)
- ACTH-hypersecreting pituitary adenoma (75%)
- Primary adrenal Cushing's syndrome (15%)
- Ectopic ACTH syndrome :neuroendocrine tumors or bronchogenic malignancies (<10%)



The surgical treatment Adrenalectomy is more than 90% effective in the treatment of primary adrenal Cushing's syndrome

Bilateral adrenalectomy is indicated in

- Patients with macroadenoma or hypophysial hyperplasia in which medical treatment and transsphenoid surgery have failed
- Bilateral benign tumors, metastatic neoplasia
- Ectopic ACTH syndrome due to occult or disseminated tumors, tumor cannot be localized

A CASE REPORT

A 29 year-old male who has two years histories of Cushing's disease and transsphenoidal pituitary tumor resection was admitted to the hospital because of recurrent Cushing's syndrome for two weeks.

Symptoms and Signs Truncal obesity, Purple striae, Moon facies Extreme muscle weakness, Hypertension Laboratory tests Severe hypokalemia Cortisol level 53.50 microgm./dl (normal 6-17) ACTH 37.9 pg/ml (normal 0-21) Low-dose dexa. Suppresion test - negative **High-dose dexa.** Suppresion test - negative MRI brain - small pittuitary gland, no mass **CT chest & abdomen - negative for neoplasms CT abdomen - bilat. adrenal hyperplasia**

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Results **Operative time was 305 minutes,** including repositioning. There were no intraoperative and postoperative complications. The patient resumed a regular diet on the first postoperative day. Inpatient postoperative hospital length of stay was 7 days, mainly for steroid-replacement and medical management.

CONCLUSIONS

Laparoscopic bilateral adrenalectomy for Cushing's disease refractory to medical management can be performed with low morbidity. Symptoms and signs of hypercortisolism rapidly improve postoperatively.

