American Board of Surgery In-**Training Exam ABSITE - 2011** Matt Landman (With lots of material borrowed from Bailey and Forbes) January 7, 2011



"Never memorize what you can look up in books"

#### @ DESPAIR.COM



#### PROCRASTINATION

HARD WORK OFTEN PAYS OFF AFTER TIME, BUT LAZINESS ALWAYS PAYS OFF NOW.



#### WISHES

WHEN YOU WISH UPON A FALLING STAR, YOUR DREAMS CAN COME TRUE. UNLESS IT'S REALLY A METEOR HURTLING TO THE EARTH WHICH WILL DESTROY ALL LIFE. THEN YOU'RE PRETTY MUCH HOSED NO MATTER WHAT YOU WISH FOR. UNLESS IT'S DEATH BY METEORITE.



#### PRESSURE

IT CAN TURN A LUMP OF COAL INTO A FLAWLESS DIAMOND-OR AN AVERAGE PERSON INTO A PERFECT BASKETCASE.

#### ABSITE Review – January 2011

#### • January 7

- Endocrine surgery
- Surgical principles
- Hepatobiliary & Transplant surgery
- Surgical oncology
- January 14
  - Practice questions
- January 21
  - Thoracic surgery
  - Vascular Surgery
  - Surgical Critical Care and Trauma
  - Head & Neck
  - Peds Surgery
- January 28
  - High Yield ABSITE Topics
- January 29 ABSITE

#### How It's Broken Down?

- 220 questions
  - Junior level (PGY 1 & 2) Exam
    - 60% Basic Science
    - 40% Clinical Management
  - Senior Level (> PGY 3) Exam
    - 20% Basic Science
    - 80% Clinical Management

#### Break down . . .

<b>Content Category</b>	<b>Junior Level</b>	Senior Level
Body as a whole (General Surgery and Surgical Critical Care)	67%	25%
GI Tract	10%	25%
CV/Respiratory	7.8%	16.7%
GU, Head/Neck, Skin, MS, CNS	7.8%	16.7%
Endocrine, Spleen, Lymphoma, Breast	7.8%	16.7%

Endocrine surgery

- Asymptomatic Adrenal Mass
  - Rule out pheochromocytoma before biopsy
  - Surgery is indicated if:
    - > 4-6cm
    - Functioning
    - Enlarging
  - Breast cancer most common metastases to adrenal gland.

- Adrenal insufficiency
  - #1 cause is withdrawal of exogenous steroids
- Cushing syndrome
  - Diagnosis:
    - 1. 24-hour urine cortisol
    - 2. Low dose dexamethasone suppression test
    - 3. Measure ACTH
    - 4. If serum ACTH high—high dose dexamethasone suppression test
  - Etiology:
    - 1. Pituitary adenoma
    - 2. Ectopic ACTH (small cell lung CA)
    - 3. Adrenal adenoma

- Thyroid Hormones
  - Thyroxine (T4)
  - Triiodothyronine (T3)
  - Calcitonin
  - T4  $\rightarrow$  T3 peripherally by deiodinase
  - T3 10x more active
  - Calcitonin produced by parafollicular cells
    - Opposes PTH

- Hashimoto Thyroiditis
  - Autoimmune disease with lymphocytic fibrosis and painless goiter
  - Eventually degenerates into hypothyroidism
  - Also associated with thyroid lymphoma (usually B cell)
  - Rx:
    - Thyroid hormone replacement
    - Surgery for symptomatic goiters refractory to med management

- Asymptomatic thyroid nodule
  - Thyroid Function Test
  - Fine-needle aspiration (FNA)
- Papillary thyroid carcinoma
  - Most common thyroid CA (~70%)
  - Unfavorable prognosis
    - Age over 50 (women) or 40 (men)\*
    - Lymphatic invasion
    - Tall cell variant

- Hypoparathyroidism
  - s/p parathyroidectomy or thyroidectomy, radiation, autoimmune syndromes, defects in calcium-sensing receptor
  - Symptoms
    - Seizures
    - Personality changes
    - Chvostek sign (facial twitching with facial n. stim)
    - Trousseau sign (carpal spasm due to blood pressure cuff)
  - Primary disease low PTH and Ca
  - Secondary disease low PTH, high calcium
  - Pseudohypoparathyroidism
    - Due to genetic defects leading to low Ca, high phosphate, high PTH and insensitivity to PTH
    - Rare
  - Rx calcium and vitamin D

- Primary Hyperparathyroidism
  - Elevated PTH
  - Elevated serum Ca
  - Elevated urine Ca
- Secondary Hyperparathyroidism
  - Elevated PTH
  - Normal serum Ca
  - Elevated urine Ca
- Familial Hypercalcemic Hypocalciuria (defective PTH receptor)
  - Normal PTH (or slightly elevated)
  - Elevated serum Ca
  - Low urine Ca

- Hypercalcemia
  - Most common presentation of general population and outpatients (cancer more common in inpatients)
  - Serum phosphate reduced
  - Symptoms
    - CNS changes (psychiatric overtones)
    - Nausea/abdominal pain (groans)
    - Nephrolithiasis (stones)
    - Bone pain (bones)
    - Bradycardia and shortened QT intervals
  - Other causes
    - Excess vitamins A & D, milk-alkali syndrome, thiazide diuretics

- MEN I
  - MENIN gene
  - 3 P's
    - <u>Parathyroid hyperplasia</u>
    - <u>Pancreatic islet cell tumors</u>
    - <u>P</u>ituitary adenoma

#### • MEN IIa

- Pheochromocytoma
- Medullary CA of thyroid
- Parathyroid hyperplasia
- MEN IIb
  - Pheochromocytoma
  - Medullary CA of thyroid
  - Mucosal neuromas
- Correct pheochyromocytoma 1st
- RET protooncogene

- Pituitary
  - Anterior and Posterior
    - Anterior adenohypophysis
      - Controlled by hypothalamic releasing factor/hormones
      - TSH, ACTH, LH, Growth Hormone, Prolactin, FSH
    - Posterior neurohypophysis
      - Innervated directly by the hypothalamus
      - Oxytocin (paraventricular nucleus)
      - Vasopressin/Antidiuretic hormone (supraoptic nucleus)
        - Secretion initiated by hypotension/increased osms

- Diabetes Insipidus
  - Central (decreased production of ADH) vs. nephrogenic (decreased renal response to ADH)
  - Central
    - Most common cause trauma
    - Loss of other anterior pituitary hormones
    - Inability to concentrate urine, dilute urine, hypernatremia
    - Compare urine to plasma osmolarity
    - Nephrogenic if no response to ADH admin
  - Rx
    - Central ADH
    - Nephrogenic HCTZ, amiloride or chlorthalidone

- Secretion of Inappropriate ADH (SIADH)
  - Ectopic secretion (ex lung ca)
  - CNS trauma
  - Medications
- Dilutional hyponatremia and high urine Na
- Urine Na above 20 mEq/L
- Renin-angiotensin system suppressed
- Rx:
  - Fluid restriction
  - Lasix for "water" diuresis
  - Saline infusion if Na falls below 110 mEq/L (rare)
  - If correct Na too fast = central pontine myelinolysis

- Endocrine pancreas
  - Insulin (beta cells)
    - Low C peptide levels exogenous insulin
  - Glucagon (alpha cells)
    - Leads to glucose release and provides energy during times of physiologic stress
    - Glycogenolysis
    - Present in early septic shock
  - Somatostatin (delta cells)
    - Inhibitory hormone
    - Octreotide is synthetic form

- Endocrine cancers of pancreas
  - Workup
    - Triple-phase CT of pancreas
    - Full biochemical workup
    - EUS location and tissue sample
    - Octreotide scan
    - If unable to locate on imaging, portal venous sampling

- Gastrinoma  $\rightarrow$  Zollinger Ellison syndrome
  - Elevated gastrin
    - GERD, abdominal pain, diarrhea, PUD
    - Serum gastrin >1,000 pg/mL diagnostic
    - Must be done off H2 blockers or PPI
    - Serum gastrin b/w 200-1000 require secretin stimulation test
    - Gastrinoma triangle
      - confluence of the cystic and CBD
      - junction of the 2<sup>nd</sup> & 3<sup>rd</sup> portions of duodenum,
      - junction of the neck and body of the pancreas
  - Rx: resection
    - Unresectable PPI possible gastrectomy



- Insulinoma
  - Hypoglycemia during fasting (glucose levels below 50 mg/dL)
  - Insulin-to-glucose ratio over 0.4
  - Elevated C peptide
  - 10% malignant
  - Whipple's triad
    - Symptoms of hypoglycemia
    - Low glucose measured at times of symptoms
    - Relief of symptoms with glucose
- Rx: enucleation vs. pancreatic resection depending on location
  - Unresectable disease octreotide vs. diazoxide

- Glucagonoma
  - Present with necrolytic migratory erythema, diabetes, anorexia, angular stomatitis
  - Diagnosis elevated glucagon during fasting
  - Most in body or tail of pancreas
  - Rx: resection
    - Unresectable disease octreotide

- VIPoma (Verner Morrison Syndrome)
  - WHDA syndrome
    - Watery diarrhea (profound and chronic)
    - Hypokalemia
    - Achlorhydria with acidosis
  - Most within body or tail of pancreas
- Rx: resection
  - Unresectable disease octreotide

You take a 35m to the OR for resection of a pheochromocytoma however after very careful inspection of both adrenal glands, you cannot find the tumor. The most likely location for the missing tumor is:

Lung
Liver
Aortic Bifurcation
Spleen



#### **Answer: Aortic Bifurcation**

- Most common location for an extra-adrenal pheochromocytoma is the Organ of Zuckerkandl which is located at the aortic bifurcation.
- 10% of pheochromocytomas are extraadrenal.
  - Most likely to be malignant compared to adrenal pheochromocytoma.

A 30f recently diagnosed with MEN I syndrome develops confusion, somnolence, and a shortened QT on EKG. The next appropriate step is:

- 1. Alpha-blockade
- 2. Beta-blockade
- 3. Calcium
- 4. IVF and Lasix



#### Answer: IVF and lasix

- MEN I syndrome includes parathyroid hyperplasia, pancreatic tumors, and pituitary tumors.
- Symptoms of hypercalcemia:
  - Somnolence
  - Shortened QT on EKG
  - Confusion
- Treatment for hypercalcemia is IVF (NS) and lasix

After thyroidectomy, an opera singer has a loss of voice pitch. This is most likely due to injury of the:

- 1. Superior laryngeal nerve
- 2. Recurrent laryngeal nerve
- 3. Vagus nerve
- 4. Hypoglossal nerve



#### Answer: Superior laryngeal nerve

- Superior laryngeal nerve injury
  - Loss of projection and easy voice fatigability
- Recurrent laryngeal nerve injury
  - Reults in hoarsness
  - Bilateral injury can obstruct the airway

Which of the following is most consistent with secondary hyperparathyroidism?

- 1. Ca 11, PTH 200, Increased Ca
- Ca 8, PTH 200, Increased urine Ca, serum creatinine 7
- **3**. Ca 11, PTH 400, decreased urine Ca
- 4. Ca 8, PTH 45, nl urine Ca



# Answer: Ca 8, PTH 200, elevated urine Ca, serum creatnine 7

- Secondary hyperparathyroidism occurs primarily in dialysis patients who have chronic loss of Ca.
- Secondary hyperparathyroidism is associated with:
  - Elevated PTH
  - Normal serum Ca
- The most common indication for operation is bone pain as a result of Ca resorption.



- 1<sup>st</sup> st. deviation 68% of all values
- 2<sup>nd</sup> st. deviation 95.5% of all values
- 3<sup>rd</sup> st. deviation 99.7% of all values

- Prevalence
  - Measure of the number of cases of a particular disease or condition at a particular time point
  - Cross-section of a population
- Incidence
  - Measure of the rate of occurrence of new cases of a disease or condition in a specified period of time (usually one year)

- Sensitivity, Specificity, Positive predictive value and negative predictive value
- Sensitivity in patients with a disease the proportion of true positives to all with the disease (TP/(TP + FN)
  - High sensitivity good for screening tests b/c miss few who actually have the disease
- Specificity in patients without a disease, the proportion of who test negative (TN/(TN+FP)
- Trade off b/w sensitivity and specificity demonstrated in a receiver operator curve (ROC curve)

		Condition (as determined by "Gold standard")		
		Positive	Negative	
Test outcome	Positive	True Positive	False Positive (Type I error, P-value)	$\rightarrow$ Positive predictive value
	Negative	False Negative (Type II error)	True Negative	→ Negative predictive value
		↓ Sensitivity	↓ Specificity	

- Null hypothesis (H0) = treatment being evaluated will have no effect i.e. no difference b/w groups
- Type I error
  - Incorrectly rejecting H0
  - P-value
- Type II error
  - Incorrectly failing to reject H0 failing to see a treatment effect when one exists
  - Avoided by increasing the power of a study (done by increasing sample size)
  - Power = probability that a test will detect true differences

- T-test
  - Compares means of two groups (assumes normally distributed data)
- ANOVA
  - Similar to t-test but used for 3 groups
- Precision
  - Measure of reliability of a testing instrument the degree to which further measurements will show the same result
- Accuracy
  - Ability of a test to obtain a result that closely resembles the true result

- Primary prevention
  - Prevent the development of the disease
- Secondary prevention
  - Reduce the progression of the disease
  - Focuses on early disease detection
- Tertiary prevention
  - Efforts to reduce the negative impact an existing disease has on a patient's life
    - Ex: treatment of heart disease to reduce symptoms of dyspnea

- Breast Disease important nerves relative to mastectomy
  - Long thoracic nerve
    - Winged scapula
  - Thoracodorsal nerve
    - Weak arm pullups and adduction
  - Intercostobrachial nerve
    - Provides sensation to medial arm and axilla

Workup of breast mass

- Evaluation is influenced by the patient's age, the physical findings, and the perceived level of breast cancer risk.
  - •<30 years old:
    - Diagnostic US
  - •30-39:
    - Bilateral diagnostic mammogram and/or targeted US
  - •<u>></u>40:
    - Bilateral diagnostic mammogram with targeted views + US

**Breast Cancer treatment options** 

•Simple mastectomy

- Not indicated for Breast CA treatment
- Used for DCIS and LCIS
- Lumpectomy and SLNB (or ALND), postop XRT
- Modified radial mastectomy
  - Removes all breast tissue including the nipple areolar complex and level I axillary LN
- Radical mastectomy
  - Rarely performed

- BRCA 1 and 2
- Tumor suppressor genes; DNA repair
- Increased risk of breast cancer (40-85%)
- BRCA I
  - Ovarian cancer 40% lifetime risk
  - Male breast cancer 1% lifetime risk
  - Associated with endometrial cancer
- • BRCA II
  - Ovarian cancer 10% lifetime risk
  - Male breast cancer 10% lifetime risk
- • Considerations for prophylactic mastectomy
  - Family history + BRCA gene
  - LCIS

- Indications for XRT:
  - Skin or chest wall involvement
  - + margins
  - Tumor > 5 cm
  - Inflammatory CA
  - Advanced nodal disease
- Chemotherapy indicated if + nodes or tumor > 1cm
- Tamoxifen indicated for PR or ER + tumors

#### • LCIS

- No calcifications, no palpable mass
- Marker for the development of breast CA (not premalignant)
- 5% risk of having a synchronous breast CA at the time of diagnosis

- Phyllodes tumor
  - Fibroepithelial breast tumor
  - Majority are benign
  - 1cm margins
  - Some malignant and behave like sarcomas in that they spread hematogenously (i.e. no SLNBx/ALND required)

- Pseudoangiomatous Hyperplasia (PASH)
  - Benign, enlarging breast mass
  - Extensive empty spaces on histology (surrounded by stromal cells)
  - Endothelial markers are negative (as opposed to angiosarcoma which has markers for VEGF)
  - Rx: typically observation

- Familial adenomatous polyposis (FAP) is autosomal dominant and all have cancer by 40.
- APC gene on chromosome 5.
- With FAP, need total colectomy prophylactically by age 20.
- With FAP, check duodenum for polyps or CA q 2years.
- Most common cause of death is periampullary tumors.
- In Gardner's syndrome, patients get colon CA (associated with APC gene) and desmoid tumors/osteomas.

- Lynch syndrome (hereditary nonpolyposis colon cancer) are autosomal dominant and associated with DNA mismatch repair gene.
- HNPCC has predilection for R-sided and multiple CAs.
- Lynch I is just colon CA risk and Lynch II patients also have an increased risk of ovarian, endometrial, bladder, and stomach CA.
- With HNPCC, need surveillance colonoscopy starting at age 25 or 10 years before primary relative got CA.

- The quinolone antibiotics (ciprofloxacin) work by inhibiting DNA production (DNA gyrase).
- Tamoxifen works by binding to and inhibiting the estrogen receptor.
- Taxol (Paclitaxel) is a tubulin inhibitor, thus it inhibits microtubules.
- Proscar (Finasteride) works by inhibiting steroid 5alpha reductase.
- Plasmids are the most common mechanism of acquiring antibiotic resistance.

**Chemotherapy Toxicities** 

- Bleomycin pulmonary fibrosis
- Cyclophosphamide hemorrhagic cystitis
- Doxorubicin cardiomyopathy
- Vincristine peripheral neuropathy
- Cyclosporine renal toxicity

Transplant Surgery

- Type 1 Immediate Hypersensitivity
  - IgE-mediated leads to activation of mast cells and granulocytes
  - Ex: allergic reactions = hives, wheals, urticaria, itching
  - anaphylaxis

- Type II antibody mediated hypersensitivity
  - IgG and IgM against cell surface or extracellular proteins
  - Tissue injury via cell opsonization and phagocytosis and/or complement activation of leukocytes
  - Ex: autoimmune hemolytic anemia, Graves, Bullous pemphigoid

- Type III complex mediated hypersensitivity
  - Immune complexes of circulating IgG and IgM deposit in tissues
  - Tissue damage via complement and Fc receptor recruitment and activation of leukocytes
  - Ex: serum sickness, polyarteritis nodosum, rheumatoid arthritis

- Type IV T Cell mediated hypersensitivity
  - Activation of CD4+ T cells (delayed hypersensitivity) and CD8+ T cells
  - Damage via cytokine-mediated inflammation, macrophage activation and direct cell lysis
  - Ex: TB skin test, contact dermatitis