

## Case Simulation

### High Blood Pressures

**Tim Nguyen\* and Harold Doerr**

Department of Anesthesiology, University of Texas Medical School, Houston, USA

**\*Corresponding author:** Tim Nguyen, Department of Anesthesiology, University of Texas Medical School, Houston, USA,  
E-mail: [Timothy.Nguyen@uth.tmc.edu](mailto:Timothy.Nguyen@uth.tmc.edu)

**Received Date: June 29, 2014**

**Accepted Date: Aug 06, 2014**

**Published Date: Aug 11, 2014**

**Citation:** Tim N., et al. High Blood Pressures. (2014) J Anesth Surg 1(1): 1-5.

#### Section 1: Demographics

**Patient Name:** Roy G Biv

**Scenario Name:** A day in the OR

**Simulation Developer(s):** Tim Nguyen and Harold Doerr

**Date(s) of Development:** 5/11/13 – 7/22/13

**Appropriate for following learning groups (circle all that apply)**

Faculty	CME						
Residents: (PGY)	1	2	3	4	5	6	7
Specialties:	Anesthesiology		Nurse Anesthesia			Surgery	
	Critical Care		Emergency Medicine			Obstetrics	
Medical Students (yr):	1	2	3	4			
Nurse Anesthesia Faculty: CEU							
Nursing Students (yr):	1	2					
Other:							

#### Section 2: Curricular Information

##### Educational Rationale

To learn to treat and diagnose pheochromocytoma in the perioperative setting

**Learning Objectives:** (ACGME Core Competencies: Medical knowledge (MK), Patient care (PC), Practice-based learning and improvement (PLI), Interpersonal and communication skills (CS), Professionalism (PR), Systems-based practice (SBP))

- Objective 1: Medical Knowledge
- Objective 2: Patient Care
- Objective 3: Practice-base learning
- Objective 4: Interpersonal and communication skills

##### Guided Study Questions

- Question 1: How to diagnose pheochromocytoma?
- Question 2: How to treat pheochromocytoma before surgery?
- Question 3: How to treat a patient with pheochromocytoma during surgery?

##### Didactics

##### Pheochromocytoma<sup>[1-3]</sup>

- Only important disease a/w adrenal medulla
- Most secrete both epi and norepi
- <0.2% hypertensive patients
- Surgery curative in >90%
- 90% are solitary tumors localized to single adrenal gland, usually on the right (non inherited)
- 95% located in abdomen

##### Inheritance

- 5% inherited as familial autosomal dominant trait
- MEN type IIA: medullary carcinoma of thyroid, parathyroid hyperplasia, and pheo
- MEN type IIB: medullary carcinoma of thyroid, pheo, and neuromas of oral mucoas
- a/w von Recklinghausen neurofibromatosis or von Hippel-Lindau disease (retinal and cerebellar angiomas)
- Rarely extra-adrenal or malignant
- 75% bilateral: bilateral adrenalectomy should be considered (familial type)

##### Clinical Presentation

- Most common in young to mid-adult life
- Tumors not innervated, catecholamine release independent of neurogenic control
- Sustained HTN or paroxysmal leading to risk of CVA, heart failure, dysrhythmias, and MI
- HA, palpitations, tremor, profuse sweating, pallor or flushing sometimes confused for malignant hyperthermia (Figure 1)

##### Diagnosis

- Free catecholamine concentration and metabolites in urine most common screening test (urinary vanillylmandelic acid and unconjugated NE and EPI levels measured in 24h urine)

collection)

- Less specific: ECG (LVH and nonspecific Twave abn), CXR (cardiomegaly), CBC (elevated Hct/hemoconcentration)
- US/CT/MRI for noninvasive localization of tumors
- Metaiodobenzylguanidine scintigraphy effective in localizing recurrent/extra-adrenal masses (Figure 2)

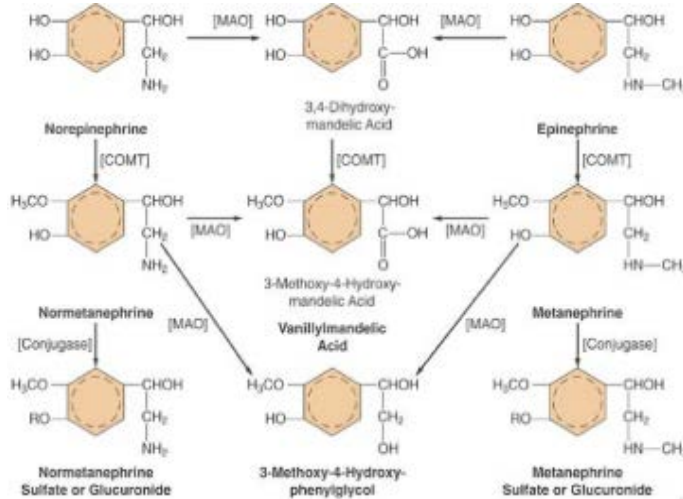


Figure1: Catabolism of norepinephrine and epinephrine

## CT abdomen

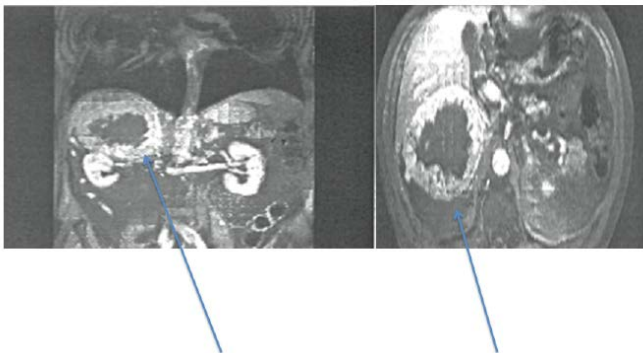


Figure2: CT abdomen

### Markers

- Vanillylmandelic acid excretion: 81% sensitive, 97% specific
- Catecholamine excretion: 82% sensitive, 95% specific
- Metanephrine excretion: 83% sensitive, 95% specific
- Abdominal CT: 92% sensitive, 80% specific
- Paroxysmal HTN, HA, sweating, tachycardia: 90% sensitive, 95% specific
- Ultrasound has largely been replaced by CT as it is limited because of overlying bowel gas

### Preop Preparation

- perioperative mortality was 45%
- decreased to 0-3% following  $\alpha$  antagonists for preop therapy
- $\alpha$ -adrenergic blockade is initiated once diagnosis of pheo is established

- Phenoxybenzamine, long-acting (24-48h) is noncompetitive presynaptic ( $\alpha_2$  and postsynaptic  $\alpha_1$  blocker dosed at 10mg q8h
- Doxazosin, terazosin, prazosin is a selective competitive  $\alpha_1$  blocker, dosed initially as 1mg PO qHS because of postural hypotension

### Preop cont

- Phenoxybenzamine or prazosin shown to be equally effective in controlling BP
- $\alpha$  blockade therapy recommended at least 10-14days prior to surgery until AM of surgery
- Allows for contracted intravascular volume/hematocrit to return to normal and stabilize BP
- $\beta$  adrenergic blockade occasionally added after  $\alpha$ -blockade in patients with persistent tachycardia/dysrhythmias
- Not be given until adequate  $\alpha$  blockade
- $\alpha$ -methyl tyrosine
- Inhibits enzyme tyrosine hydroxylase, rate-limiting step in catecholamine biosynthesis
- Reserved for those with metastatic disease or those which surgery is contraindicated and long-term medical therapy is required, i.e. patients with decompensated CHF, end stage COPD.
- When used in combo with  $\alpha$ -adrenergic blocking agents, there is significant reduction in catecholamine biosynthesis

### Pregnancy

- Unrecognized pheochromocytoma during pregnancy may be life-threatening to mom and baby
- Adrenergic blockers probably improve fetal survival, but safety has not been established and no well-designed controlled studies have been done to prove this.
- Perform surgery during first trimester or time of CS
- No reason to terminate early pregnancy, but should be aware of risk of spontaneous abortion from surgery

### Perioperative Management

- If can't start  $\alpha$ -blocker prior to surgery or <48h of treatment, can use nitroprusside
- If solitary tumor w/out mets, surgery involves laparoscopic retroperitoneal approach, and maybe later converted to open if needed
- During laparoscopy, pneumoperitoneum may cause release of catecholamines and hemodynamic changes can be controlled with vasodilator

### Periop cont

- Manipulation of tumor may produce marked elevations in BP
- Acute hypertensive crises treated with IV nitroprusside or phentolamine
- Phentolamine is a short acting  $\alpha$  adrenergic antagonist that's given as 2-5mg IV bolus or continuous infusion as the half-life is 19mins
- Tachydysrhythmia controlled with IV propranolol (1mg in-

cremets) or continuous infusion of ultrashort-acting selective  $\beta_1$  adrenergic antagonist esmolol

### Periop cont

- Long-acting beta blockers disadvantages: persistence of bradycardia/hypotension after tumor removed
- Magnesium sulfate infusion with intermittent boluses has been successful to control BP after maximizing  $\alpha$  adrenergic antagonists
- Nicardipine, nitroglycerin, diltiazem, fenoldopam, and prostaglandin E1 have all been used anecdotally

### Periop cont- hypotension

- Reduction of BP after ligation of tumor's venous supply can be dangerously abrupt, and close communication required
- Initial therapy: Restore intravascular fluid deficit
- If remains hypotensive: phenylephrine
- After surgery: catecholamines return to normal over several days
- 75% patients are normotensive within 10 days

### Assessment Instruments

- Mac3 or Miller
- Endotracheal tube
- Arterial line set up

### Section 3: Preparation

#### Monitors Required

1	Non-Invasive BP Cuff			
1	Arterial Line			
	CVP			
	PA Catheter			
1	5 lead EKG			
1	Temperature Probe			
1	Pulse Oximeter			
1	Capnograph			
	BIS			

#### Other equipment required

1	Anesthesia Machine	1	ETT		
1	Pumps		LMA		
	Brochoscope	1	Laryngoscope		
	Defibrillator				
	Hotline				
	Nerve Stimulator				
	Echo Machine and Probe				

#### Supporting Files (cxr, ekg echo, assessment, handouts, etc)

1. Chest xray – no pulmonary infiltrates pleural effusions, or pneumothorax
2. Electrocardiogram – sinus tachycardia and inverted T wave on leads V5 and V6
3. Trans-Thoracic Echocardiogram – Ejection Fraction of 55%

with no regional wall motion abnormalities

Abdominal Computerized Tomography: possible acute appendicitis.

#### Time Duration

Set-up	20 mins
Preparation	4 hours
Simulation	20 mins
Debrief	20 mins
Programming/scripting	8 hours

#### Case Stem

**Case Stem (one to two paragraphs on pertinent patient and scenario information-this should be the stem for the learner and should include location, physician/help availability, family present, etc.)**

60 year old male with no past medical or surgical history presented to the Emergency Room with sudden onset dyspnea and palpitations<sup>[1,2]</sup>. For the past 2 weeks, he had been experiencing include intermittent headaches, palpitations, anxiety, and occasional shortness of breath on exertion, which is what brought him to the hospital. This morning, his wife noted increased anxiety leading to “he can’t catch his breath.” Work up includes an Electrocardiogram, cardiac enzymes, chest x-ray, stress test, and transthoracic echocardiography, all of which were normal. Incidentally, a abdominal Computerized tomography (CT) scan was performed, and an acute appendicitis could not be ruled out, so he was sent to the Operating Room to do a diagnostic laparoscopy and appendectomy.

#### Background and briefing information for Facilitator/coordinator's eyes only

In the ER, his blood pressure ranged from noninvasive systolic 140-280 millimeters mercury over diastolic 30-150 millimeters mercury, so a CT (computerized tomography) abdomen showed a 6-centimeter round mass retroperitoneally on the right adrenal gland suggesting pheochromocytoma<sup>[4,5]</sup>, but this was missed initially as it only showed a possible acute appendicitis. Serum epinephrine was 18.6 nanogram/milliliter (normal: 0.01-0.18 nanogram/milliliter), norepinephrine 24.2 nanogram/milliliter (normal: 0.06-0.45 nanogram/milliliter), and dopamine 4.8 nanogram/milliliter (normal 0-0.09 nanogram/milliliter). No metaiodobenzylguanidine (MIGB) scan performed because of acute renal failure.

#### Patient Data Background and Baseline State

#### Patient History (should follow standard H and P format)

#### Review of Systems:

- Central Nervous System: + headache, no strokes or seizures or focal deficits
- Cardiovascular: + palpitations and chest pain, no murmurs/arrhythmias
- Pulmonary: + dyspnea with exertion and now at rest, no wheezing or asthma or cough
- Renal / Hepatic: + decreased urine output, acutely, no cirrhosis

or liver problems

- Endocrine: no diabetes or excessive thirst<sup>[2]</sup>
- Hematology/Coagulation: no easy bruising or bleeding

**Current Medications and Allergies:** No current meds, No known drug allergies

**Physical Examination:**

General: No acute distress, well developed, well nourished

Weight, Height: 77kg, 170 centimeters, Vital Signs: noninvasive blood pressure 180/100 mmHg (millimeters of mercury) Heart Rate 101 beats per minute Temperature 99.9 Fahrenheit, Respiratory rate 20 breaths per minute.

Oxygen saturation: 95% on Room Air, Airway: Mallampati I, normal mouth opening, full range of motion of the neck, Lungs: bilateral equal breath sounds, no wheezes, rales, rhonchi, Heart: sinus tachycardic, no murmurs, gallops, rubs

**Laboratory, Radiology, and other relevant studies:**

Hematocrit: 45%, Chest XRay: no pulmonary infiltrates, pleural effusions, or pneumothorax, Electrocardiogram: sinus tachycardia and inverted T wave on leads V5 and V6

**Baseline Simulator State: What underlying alterations in physiology would this patient have when compared to “perfect” 70 kg man or woman? Include target numbers. This will comprise your baseline state**

- Vitals: Blood Pressure 180/100 mmHg (millimeters Mercury) Heart Rate 141 beats per minute Temperature 99.9 Fahrenheit Respiratory Rate 20 breaths per minute Oxygen saturation:97% on Room air
- Neurologic: alert and oriented to person, place, time, and no focal deficits
- Respiratory: bilateral equal breath sounds
- Cardiovascular: sinus tachycardia
- Gastrointestinal: positive bowel sounds, diffusely mildly tender to palpation
- Genitourinary: decreased urine output
- Metabolic: normal
- Environmental: normal OR setting

State	Patient Status	Student learning outcomes or actions desired and trigger to move to next state	
1. BASE-LINE	Awake, alert and oriented to person, place, time, responsive, noninvasive BP (Blood Pressure) 120s-180s/60-90s mmHg. HR (heart rate) 70-150s BPM (beats per minute)	Learner Actions: o IV (intravenous) phentolamine (i.e. 5 milligrams IV x 1) o Preop PO alpha antagonist i.e. phenoxybenzamine (i.e. 10milligrams by mouth once daily) or doxazosin	Operator: o Decrease systolic BP by 10-20 points if alpha antagonist started o Proceed to MILD if started on alpha antagonists o Proceed to Moderate if not Teaching Points: o Starting alpha 1

		(i.e. 1 milligram by mouth once daily), terazosin (i.e. 1milligram by mouth once daily), prazosin (i.e. 1milligram by mouth once daily)	blockade before beta blockers o Control HTN Trigger: alpha blockers- proceed with MILD, if given beta blockers without alpha blockers, proceed to SEVERE, if student fails to start alpha blockers at all, proceed to SEVERE and after 10mins DEATH
2. MILD	Awake, alert and oriented to person, place, time, responsive before induction. Invasive BP 120s-180s/60-90s mmHg. HR 70-150s BPM, then converted over to general anesthesia with same vitals.	Learner Actions: o General anesthesia, regular induction o Avoid Desflurane o Propofol 50 microgram/kilogram/hour + Sufenta 0.2 microgram/kilogram/hour with ½ MAC of Sevoflurane or Isoflurane o Continue IV phentolamine 5milligram IV every 5 mins	Operator: o Proceed to general anesthesia state Teaching Points: o Careful attention to vital signs o Avoid Desflurane because of tachycardia and non-neurogenic catecholamine release o Extremely important to achieve adequate depth of anesthesia before proceeding with Direct laryngoscopy to minimize sympathetic nervous system response Trigger: 5 minutes, proceed to MODERATE, unless student used desflurane, proceed to SEVERE; if student did not adequately anesthetize patient before direct laryngoscopy, proceed to SEVERE for 2 minutes and return to moderate
3. MODERATE	Unresponsive, under general anesthesia, very labile blood pressures (BP 70-200s/40-110s mmHg). (beginning to middle of the surgery)	Learner Actions: o Continuous infusion of IV phentolamine 5 milligrams IV every 5 minutes o IV magnesium 2grams IV over 30 minutes o IV beta blocker i.e. esmolol infusion 50 micrograms/kilogram/minute or propranolol 1 milligram IV x once o IV nitroprusside 0.3-4 microgram/kilogram/minute o IV nicardipine 3-5 milligram/hour	Operator: o Labile blood pressures, but not as bad as SEVERE Teaching Points: o Blood pressure will rapidly decrease after resection of the tumor Trigger: 5 minutes, or if student does not use anything besides alpha antagonist, proceed with SEVERE, if student uses IV beta blockers, nitroprusside, or any other antihypertensive agent, proceed with MILD again and then RESOLUTION afterwards in 5 minutes

4. SE-VERE	Unresponsive, under general anesthesia, very labile blood pressures (BP 150-200s/40-110s mmHg). Heart Rate is 120-180s BPM (surgeon is manipulating the tumor or no alpha antagonist started)	<b>Learner Actions:</b> <ul style="list-style-type: none"> <li>o Continuous infusion of IV phentolamine, may need to increase dose.</li> <li>o Need to add beta blockade like propranolol 1 milligram IV x once or esmolol 50 micrograms/kilogram/minute or metoprolol IV 1-2 milligram x once</li> </ul>	<b>Operator:</b> <ul style="list-style-type: none"> <li>o Tell student that surgeon also sees an adrenal mass that he would like to take out as well</li> <li>o Tell the examinees that the surgeon has just resected the tumor before proceeding with the RESOLUTION state</li> <li>o Very labile blood pressures as surgeon is manipulating tumor releasing a lot of catecholamines</li> </ul> <b>Teaching Points:</b> <ul style="list-style-type: none"> <li>o Blood pressure will rapidly decrease after resection of the tumor</li> </ul> <b>Trigger:</b> after starting beta blocker proceed to MILD for 2 minutes before proceeding to RESOLUTION or 5 minutes proceed to RESOLUTION if student fails to add anything in addition to alpha blockers; DEATH if student did not add any antihypertensives at all since the beginning, especially alpha antagonists
5. RESOLUTION	Blood pressure low 60s/40s mmHg as surgeon just resected tumor. HR 50s (beats per minute) if learner used long acting beta blocker or still on Esmolol drip	<b>Learner Actions</b> <ul style="list-style-type: none"> <li>o Start a pressor i.e. norepinephrine to maintain blood pressure, and preferably a drip i.e. Norepinephrine 0.2 micrograms/kilogram/minute</li> <li>o IV (Intravascular) fluids: 1-2Liters Lactated Ringers or Normal Saline</li> </ul>	<b>Operator :</b> <ul style="list-style-type: none"> <li>o Increased the Blood Pressure (BP) to normal values 120s/80s mmHg if the examinees titrate in a pressor</li> </ul> <b>Teaching Points:</b> <ul style="list-style-type: none"> <li>o Ligation of tumor's venous supply can be cause reduction in BP abruptly</li> <li>o May need to start a pressor after tumor is resected</li> <li>o Long acting beta blocker i.e. propranolol may lead to persistence of bradycardia/hypotension after tumor removal</li> <li>o Need to restore intravascular fluid deficit</li> </ul> <b>Trigger:</b> student starts a pressor like Norepi or phenylephrine, return to normal vital signs 120s/80s mmHg Heart Rate 80s beats per minute. If student fails to start a pressor, IV fluids, go to hypotension 60s/40s mmHg and DEATH in 5 mins

## References

1. Roizen, M.F. Anesthetic Implications of Comorbid Diseases. (2010) Miller's Anesthesia. Philadelphia: Churchill Livingstone/Elsevier.
2. Barash., Paul, G. Endocrine Function. (2009) Clinical Anesthesia. Philadelphia: Wolters Kluwer/Lippincott Williams & Wilkins.
3. Uchida, N., Ishiguro, K., Suda, T., et al. Pheochromocytoma Multisystem Crisis Successfully Treated by Emergency Surgery: Report of a Case. (2010) Surg Today 40(10): 990-996.
4. Domi, R., Laho, H. Management of Pheochromocytoma: Old Ideas and New Drugs. (2012) Niger J Clin Pract 15(3): 253-257.
5. Myklejord, D.J. Undiagnosed Pheochromocytoma. (2004) Clin Med Res 2(1): 59-62.