Resection of a Dopamine Secreting Paraganglioma Overlying the Left Main Coronary Artery

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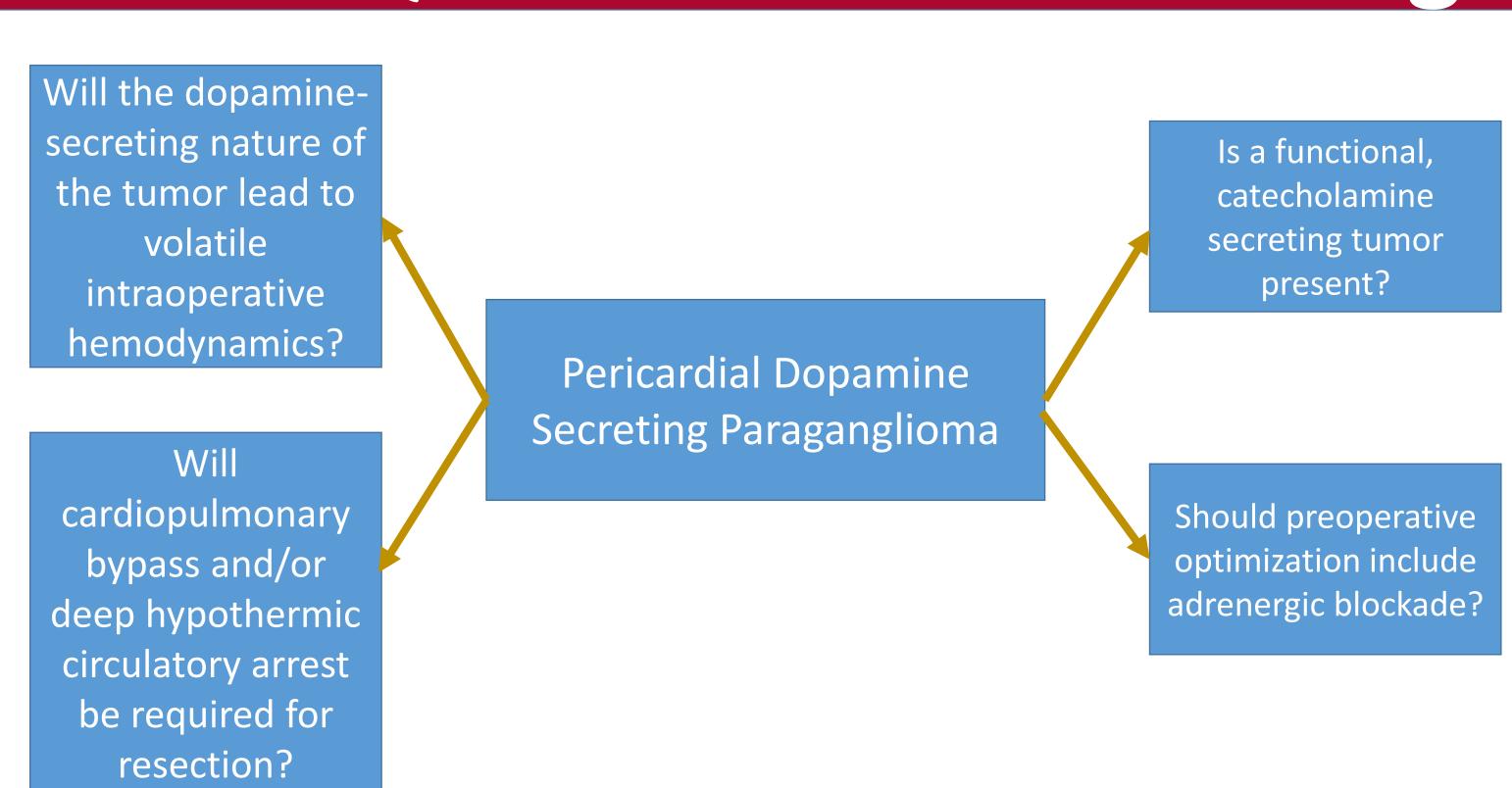




Case History

- A 66 year old male presented after a snowmobile accident resulting in right sided rib fracture
- Given the extent of trauma, a chest CT was preformed which suggested a left atrial appendage thrombus
- Follow up trans-esophageal echocardiogram, showed an extracardiac nodular echodensity adjacent to the left coronary cusp
- Cardiac MRI demonstrated an enhancing 4.2 cm soft tissue mass interdigitating with the left proximal coronary vessels
- Given the heterogeneity of the lesion, paraganglioma or hemangioma were considered
- Endocrinology studies performed were significant for 24 hour urine dopamine of 1,103 mcg/24 hours (nl 52-480)
- Cardiac catheterization evaluated the patency of the coronary vessels which identified no stenosis or apparent compression
- The patient denied any history of chest pain, palpitations, hot flashes, night sweats, or blood pressure swings
- He decided to pursue elective resection of the Paraganglioma

Clinical Questions and Challenges



Treatment Plan

Arterial Line

General

Anesthesia

Pulse Oximetry

Transesophageal Echocardiogram

Anticoagulation and

Antifibrinolytic

Therapy

Bladder emperatur

Continuous CVP

Tranexamic acid infusion for

the entirety of the case

Heparinization prior to

bypass followed by

Hemodynamic Management

perform preoperative alpha blockade

Nicardipine, Esmolol, Norepinephrine,

intermittently throughout the case,

otherwise no other vasopressors or

The anesthesia team elected to not

Prepared infusions: Nitroglycerin,

Epinephrine, Dopamine

Pt required Norepinpherine

Fentanyl, Midazolam, Rocuronium

• Induction: 2% Propofol,

• Maintenance: 1.5-2% Sevoflurane

Pathology



- Neoplastic cells are positive for chromogranin, synaptophysin, CD56, and S100 supporting diagnosis of pheochromocytoma.
- Grading of adrenal pheochromocytoma and Paraganglioma score of 2 equating to metastatic rate of 3.6%.

Postoperative Course

The patient recovered uneventfully

Conduit Harvesting and Cardiac Bypass Onset

inotropes were used

- Distal Aorta and PA Reconstruction
- Saphenous Vein harvesting for potential conduit
- Heparinization
- Distal Aorta, SVC, and IVC cannulated
- Cardioplegia cannula in coronary sinus via right atrium
- Transition to bypass
- Aortic crossclamping

 Tumor mobilized from PA and roof of right atrium

Tumor Resection, CABG,

- For complete tumor mobilization, the PA and aorta distal to the sinotubular junction were transected
- In order to remove the tumor, the left main coronary artery was sacrificed
- Reconstruction of PA and
- CABG performed to LAD and left circumflex artery

 After rewarming and reperfusion, pt was

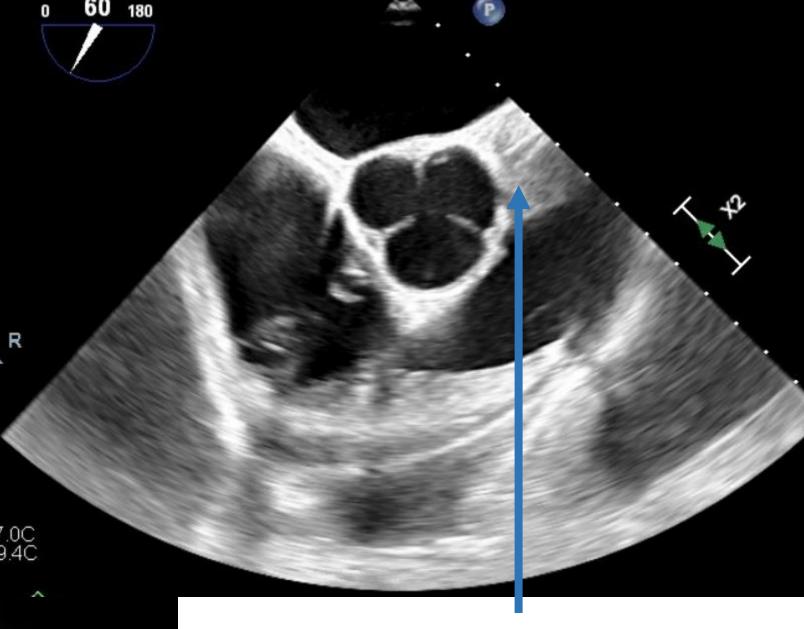
weaned gradually off

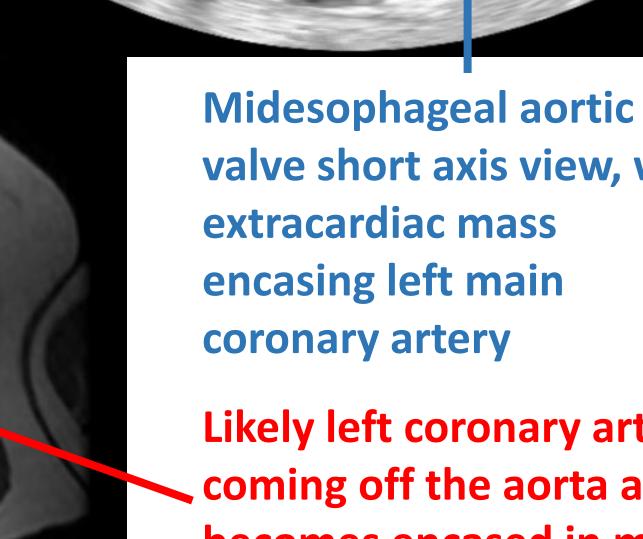
Transition off Bypass

- TEE confirmed return of biventricular function and filling status
- Heparin was reversed with protamine & coagulation function assessed with TEG
- Sternotomy was closed
- Patient remained intubated and transferred to cardiac ICU for recovery

Cardiac MRI and Echocardiogram

Heterogeneously hyperintense on T1. **Voiding throughout** mass could be intratumoral vessels or encasement of nearby vessels.





valve short axis view, with extracardiac mass encasing left main coronary artery

Likely left coronary artery coming off the aorta as it becomes encased in mass

Conclusion

- Paragangliomas have a frequency of 500-1600 cases in the US per year and of those cases, purely dopamine secreting lesions are exceedingly rare. [1]
- Given the unique pericardial location of this lesion, the following were required: aortic root reconstruction, pulmonary artery reconstruction, and coronary artery bypass grafting.
- Due to the catecholamine secreting nature of these lesions, anesthesia and surgical teams should anticipate labile hemodynamics and consider preoperative alpha blockade.

[1]: Chen, H., Sippel, R. S., O'Dorisio, M. S., Vinik, A. I., Lloyd, R. V., Pacak, K., & North American Neuroendocrine Tumor Society (NANETS) (2010). The North American Neuroendocrine Tumor Society consensus guideline for the diagnosis and management of neuroendocrine tumors: pheochromocytoma, paraganglioma, and medullary thyroid cancer. Pancreas, 39(6), 775–783. https://doi.org/10.1097/MPA.0b013e3181ebb4f0