

CAIR Case of the Month

Case Courtesy of Drs. S. Neufeld and S. Sookhoo
University of Manitoba

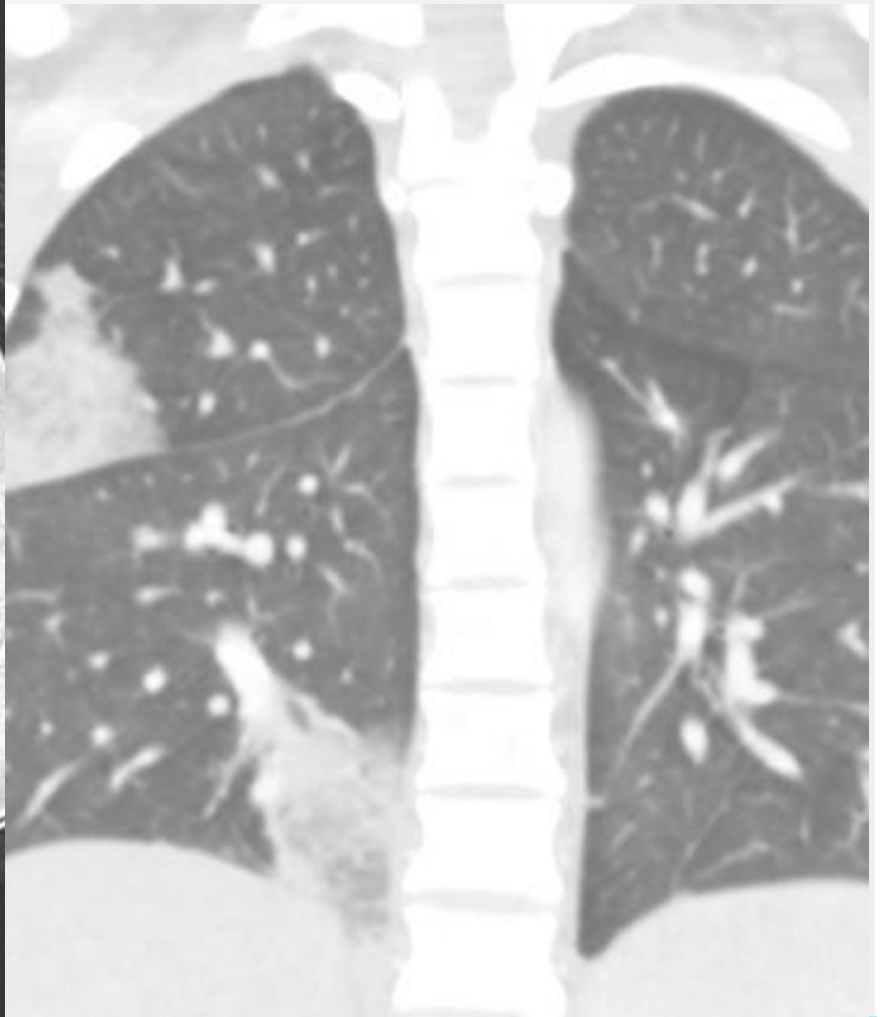
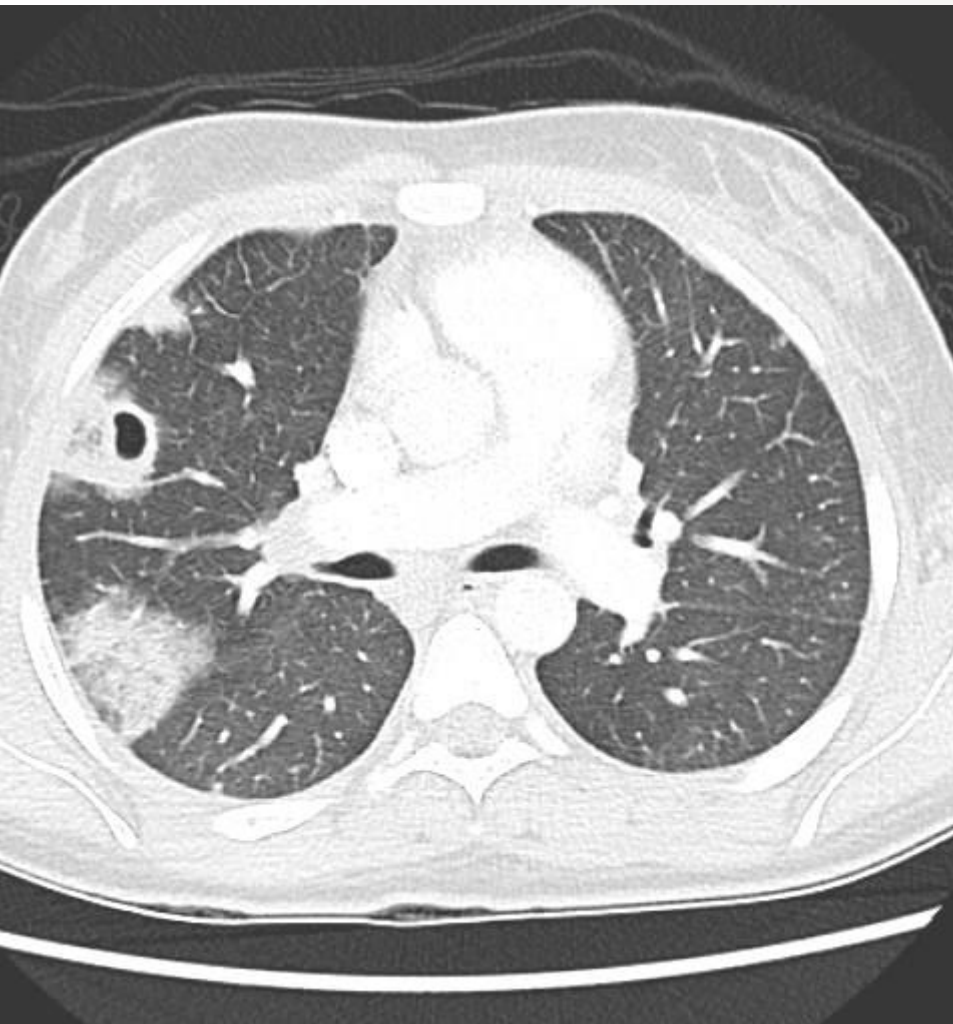


Case

- 24 year old female with history of granulomatosis with polyangiitis (GPA – formerly Wegener’s granulomatosis)
 - **Renal biopsy confirmed**
- Worsening shortness of breath, fevers, hemoptysis
- CT chest ordered in Emergency Department to reassess disease and rule out infection



Prior (2016)

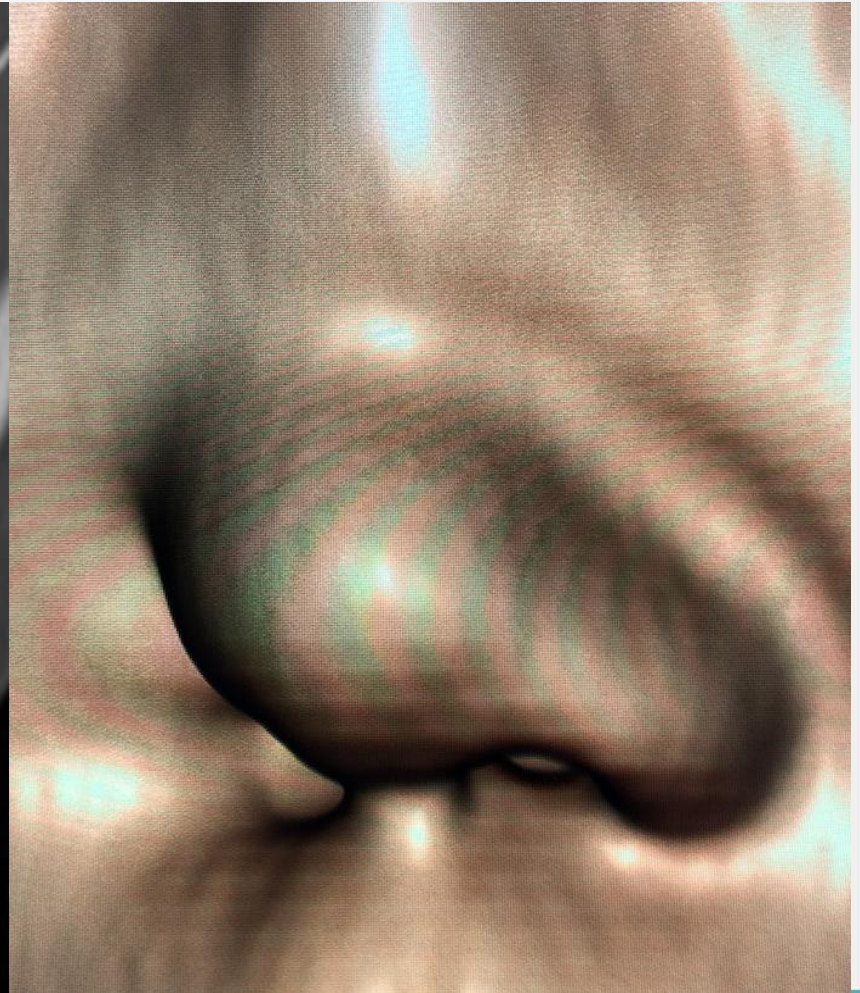


Prior Chest CT Findings

- Multiple ill defined nodules
 - Cavitation
 - Ground glass halos
- Impression
 - Granulomatosis with polyangiitis



Priors (2014)

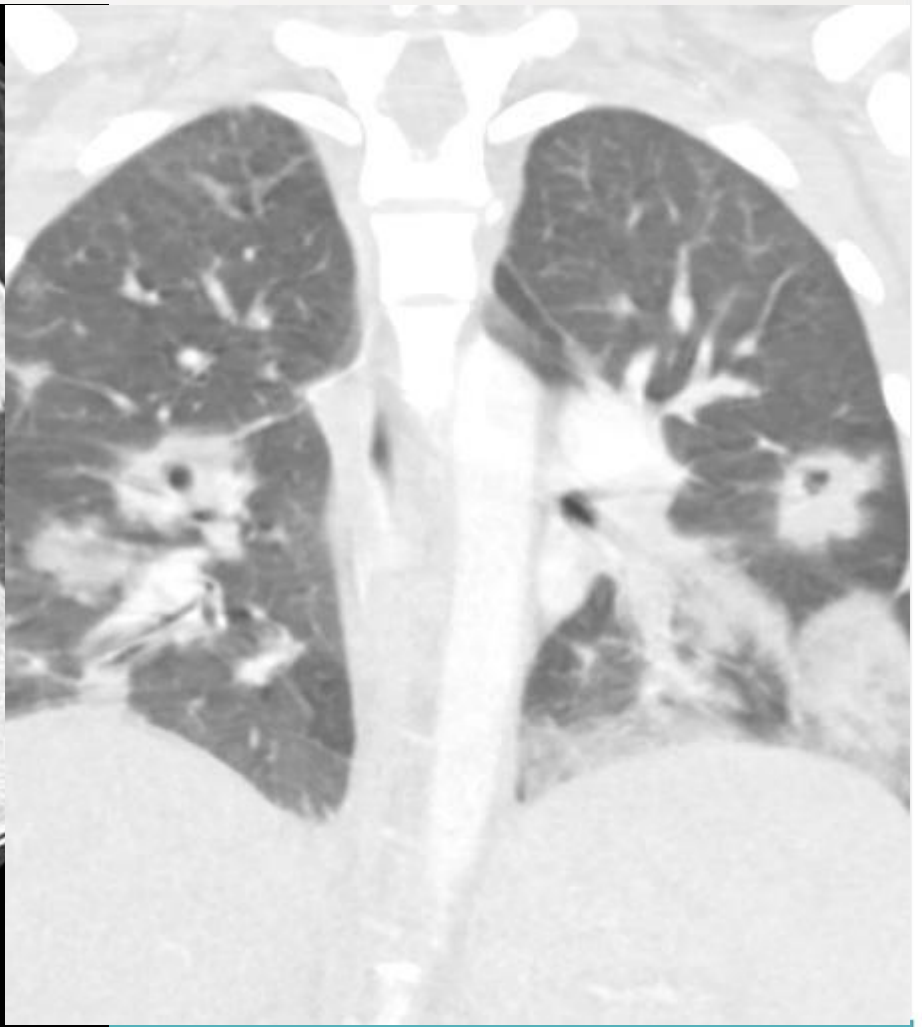
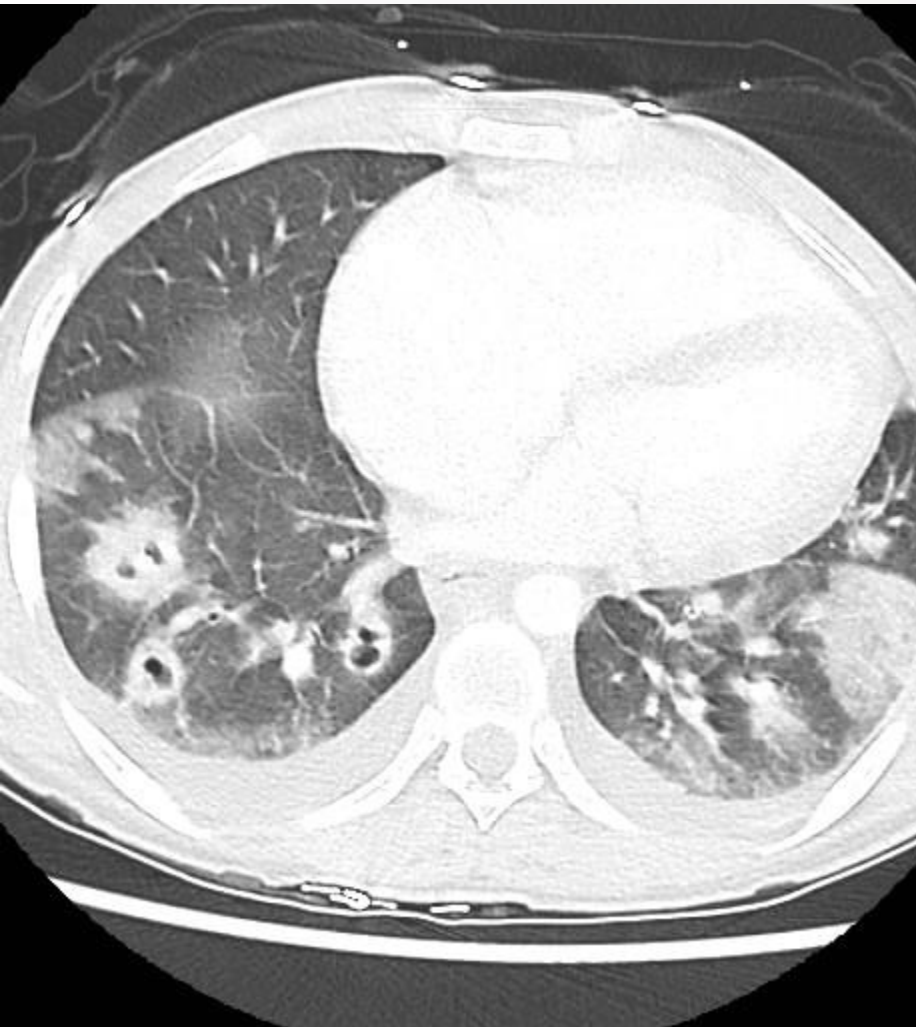


Prior Sinus CT Findings

- Saddle nose deformity
 - inferior/middle turbinates not identified
 - Septal perforation
- Impression:
Thought to be granulomatosis with polyangiitis but biopsy showed no GPA
Differential includes chronic Percocet inhalation



Admission



Current Chest CT Findings

- Multiple ill-defined nodules with cavitation
- Ground glass ‘halos’ and ‘reverse halos’
- Impression
Granulomatosis with polyangiitis

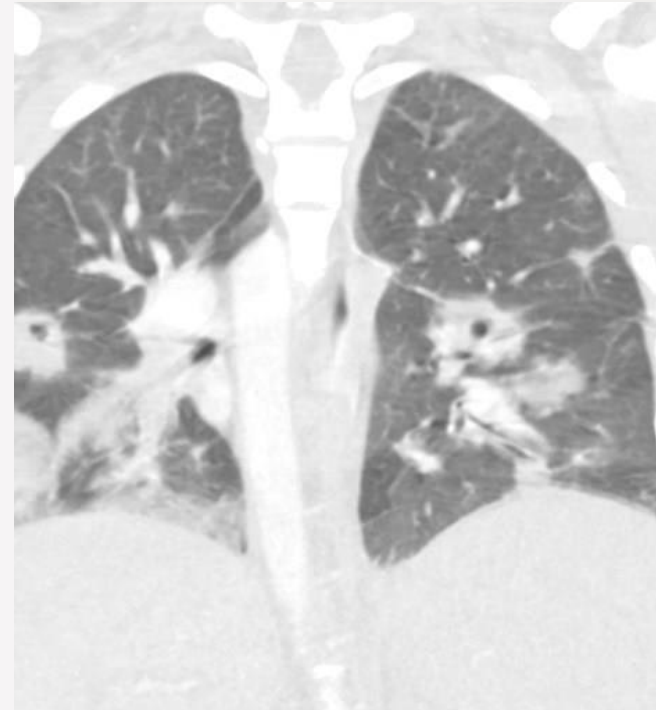
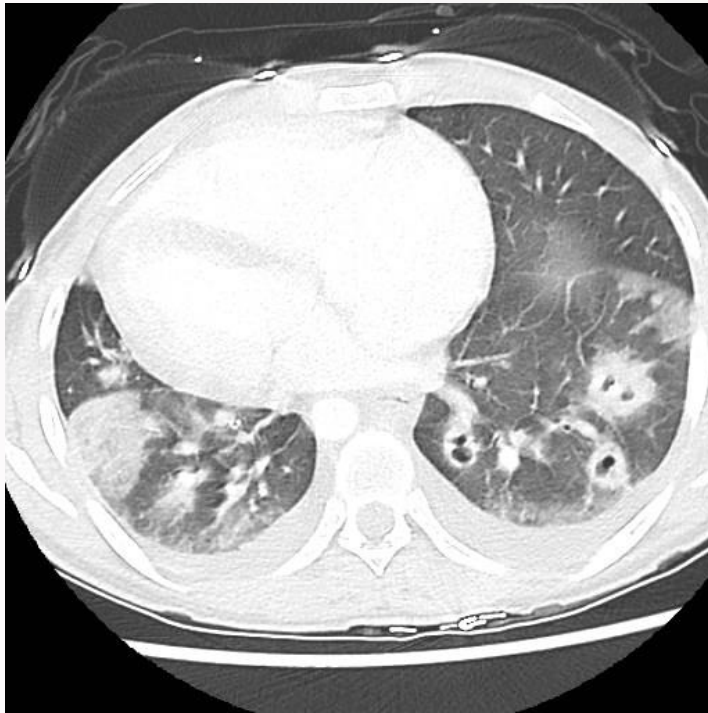


Course in Hospital

- Blood Cultures:
 - Streptococcus agalctiae
 - Enterococcus faecalis
 - Candida albicans
 - Acinetobacter
 - Lactobacillus
- Treatment:
 - Ceftriaxone
 - Ampicillin
 - Micafungin
- Echocardiogram:
 - Tricuspid regurgitation
 - 10 mm x 40 mm tricuspid valve vegetation
 - Severe right ventricular enlargement
 - Increased right sided pressure



Second Look

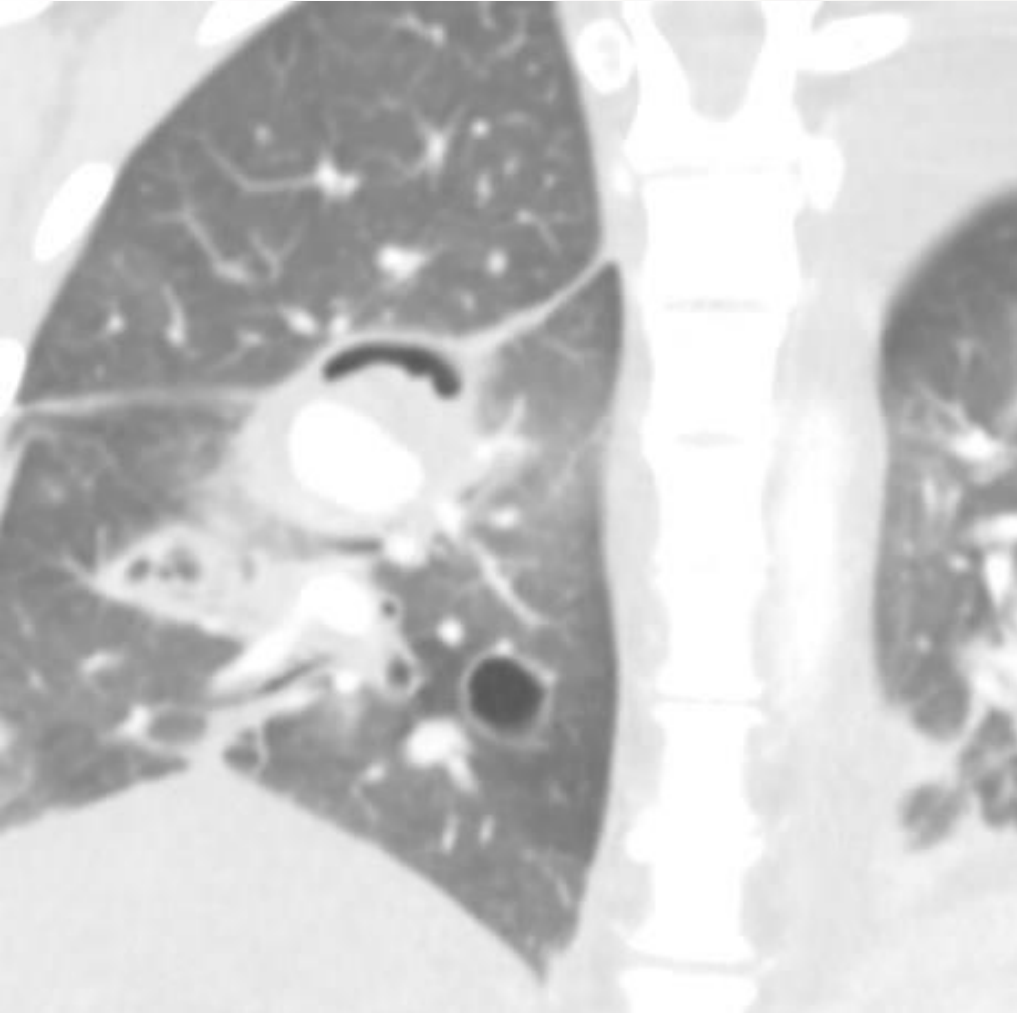


With the additional information the diagnosis is less clear

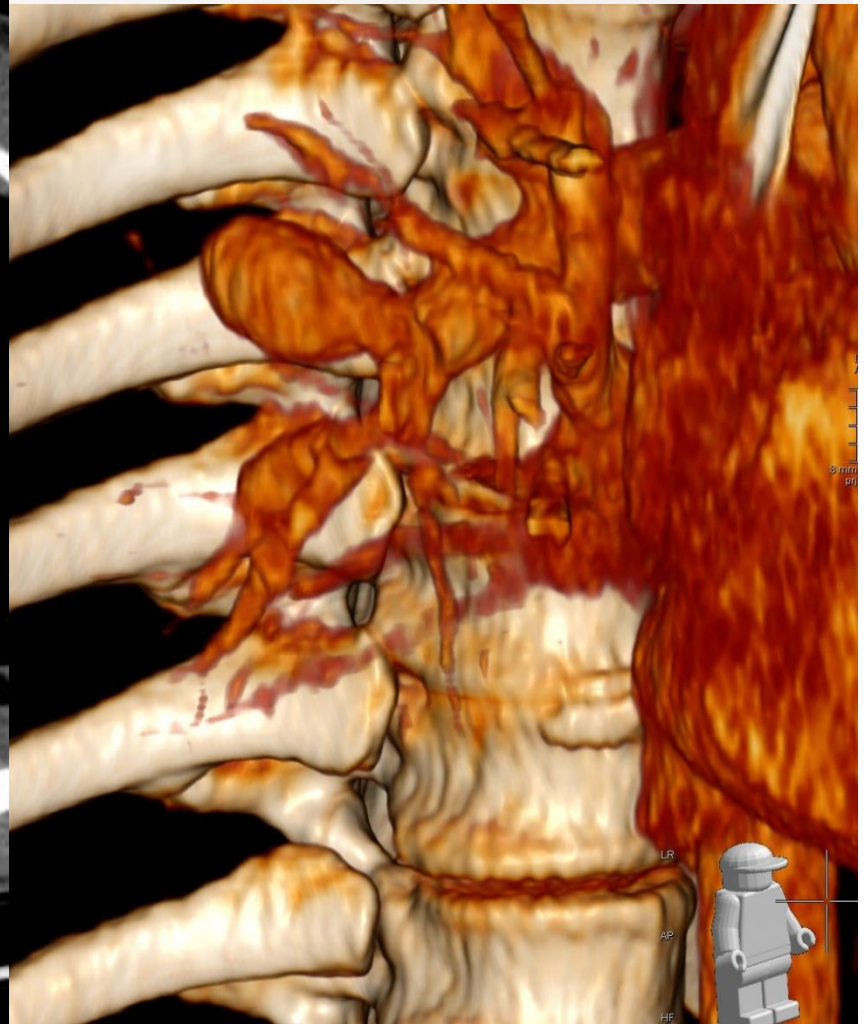
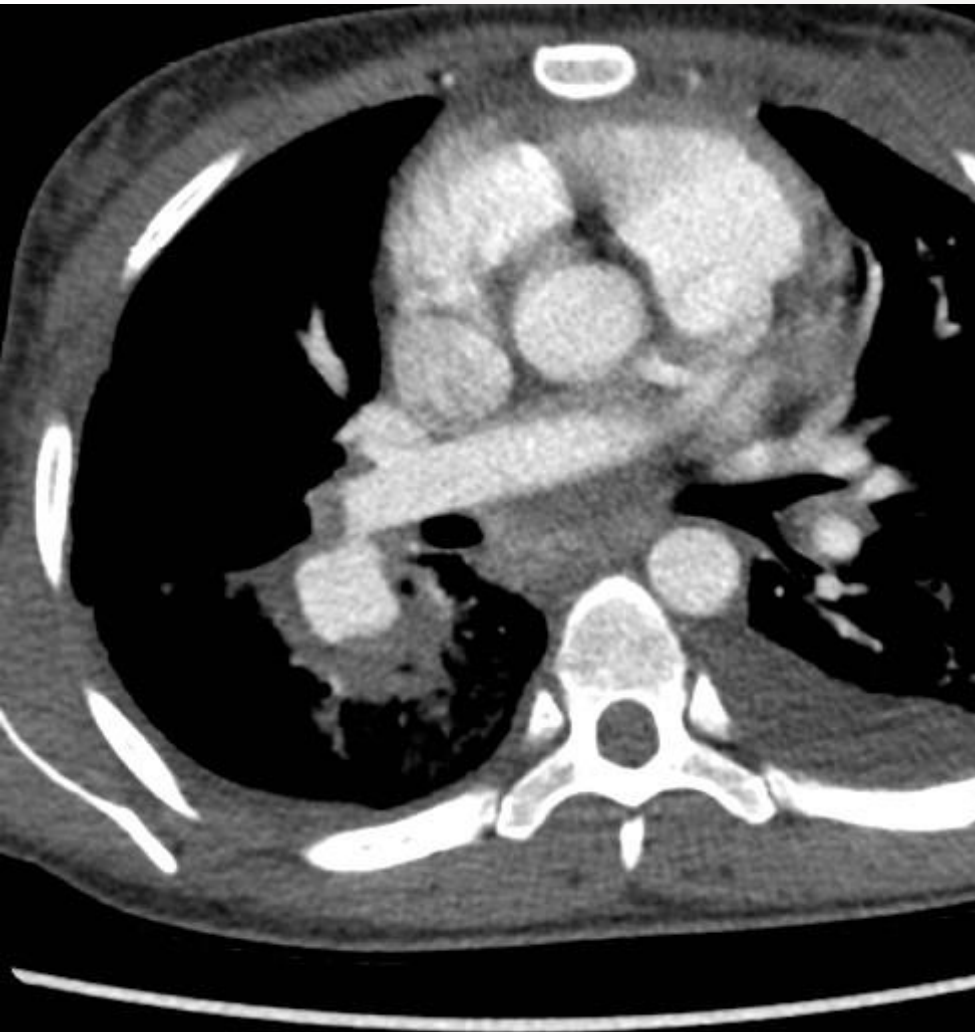
- **Granulomatosis with Polyangiitis**
- **Septic Emboli**



Repeat CT (1/2)



Repeat CT (2/2)



Repeat CT Findings

- 2.4 cm pseudoaneurysm adjacent to worsening cavitation
- Bilateral cavitory nodules
 - **Increased slightly in size**
 - **No new lesions**
- Progressive consolidation
- Multiple pulmonary emboli
- Impression:
New pseudoaneurysm



Pulmonary Artery Pseudoaneurysms

- Rare
- Diagnosed incidentally or following rupture
 - Massive hemoptysis with high mortality
- Management
 - Typically endovascular
 - Surgical management rarely indicated
- Etiology:
 - Post-Traumatic
 - Infectious
 - Vasculitis (Bechet's)
 - Congenital



Pathogenesis of Mycotic Pseudoaneurysm

- Contiguous involvement from adjacent sepsis
 - Rasmussen (TB)
- Hematogenous spread
 - Micro-emboli into vasa-vasorum
- Infection of pre-existing intimal defect
- Direct inoculation at time of vascular trauma



Treatment

- Ongoing antibiotic treatment
- Embolization of mycotic pseudoaneurysm
 - Urgent given the rapid expansion over only 10 days
 - Cardiac anesthesia present during procedure
- Tricuspid valve replacement
 - lobectomy to be performed during cardiac OR if embolization unsuccessful



Procedure

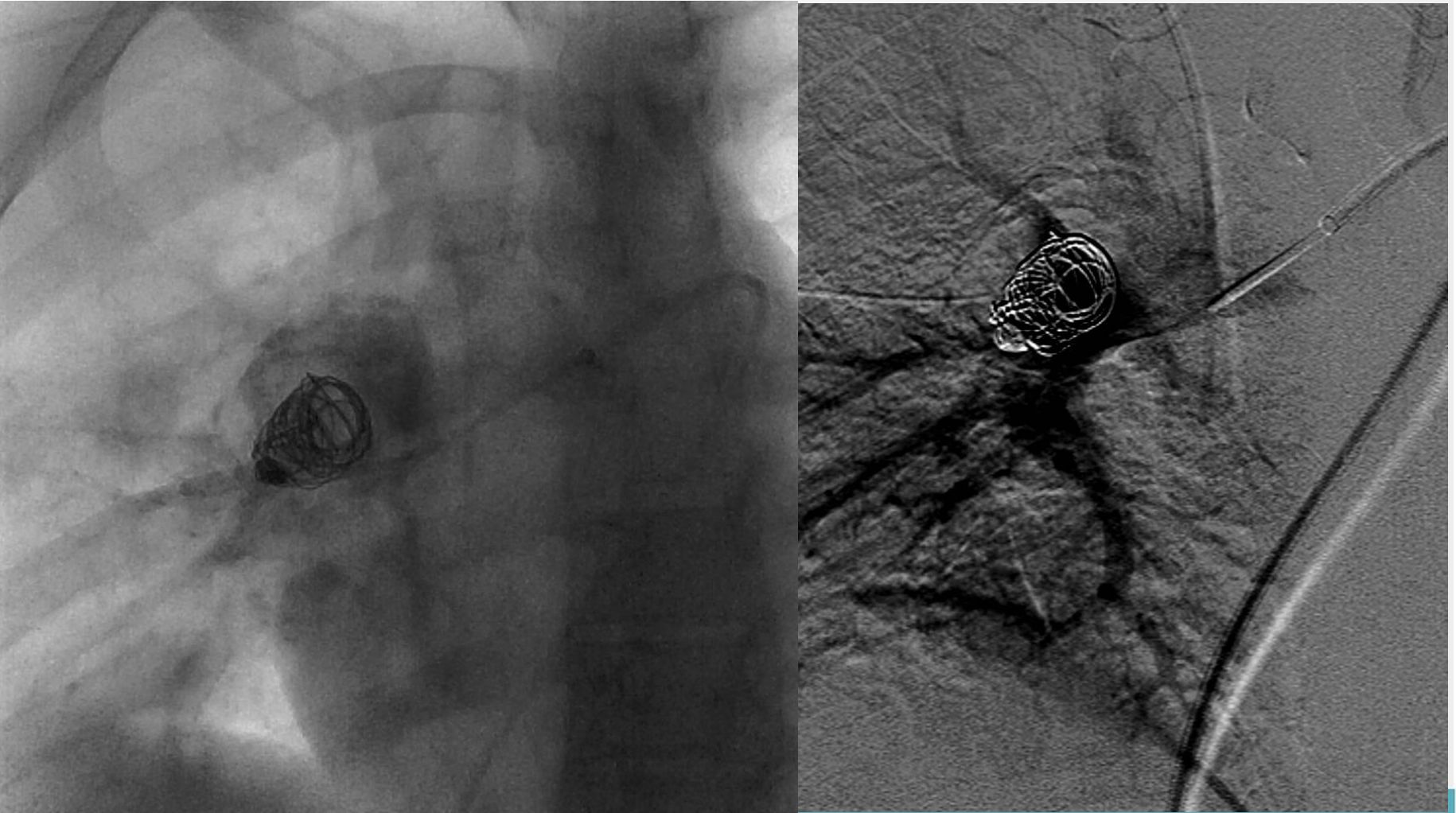
- Right femoral access
- APC catheter advanced into main PA
- Pulmonary angiogram performed
- APC catheter advanced
- 7-French sheath advanced into right main PA



Embolization



Completion Angiogram



Procedural Details

- Microcatheter advanced into aneurysm sac
- 5 micro coils were successfully deployed
- The 6th coil plugged in the catheter, necessitating catheter removal
- Access to the aneurysm sac could not be re-established
- Completion angiogram demonstrated aneurysm exclusion

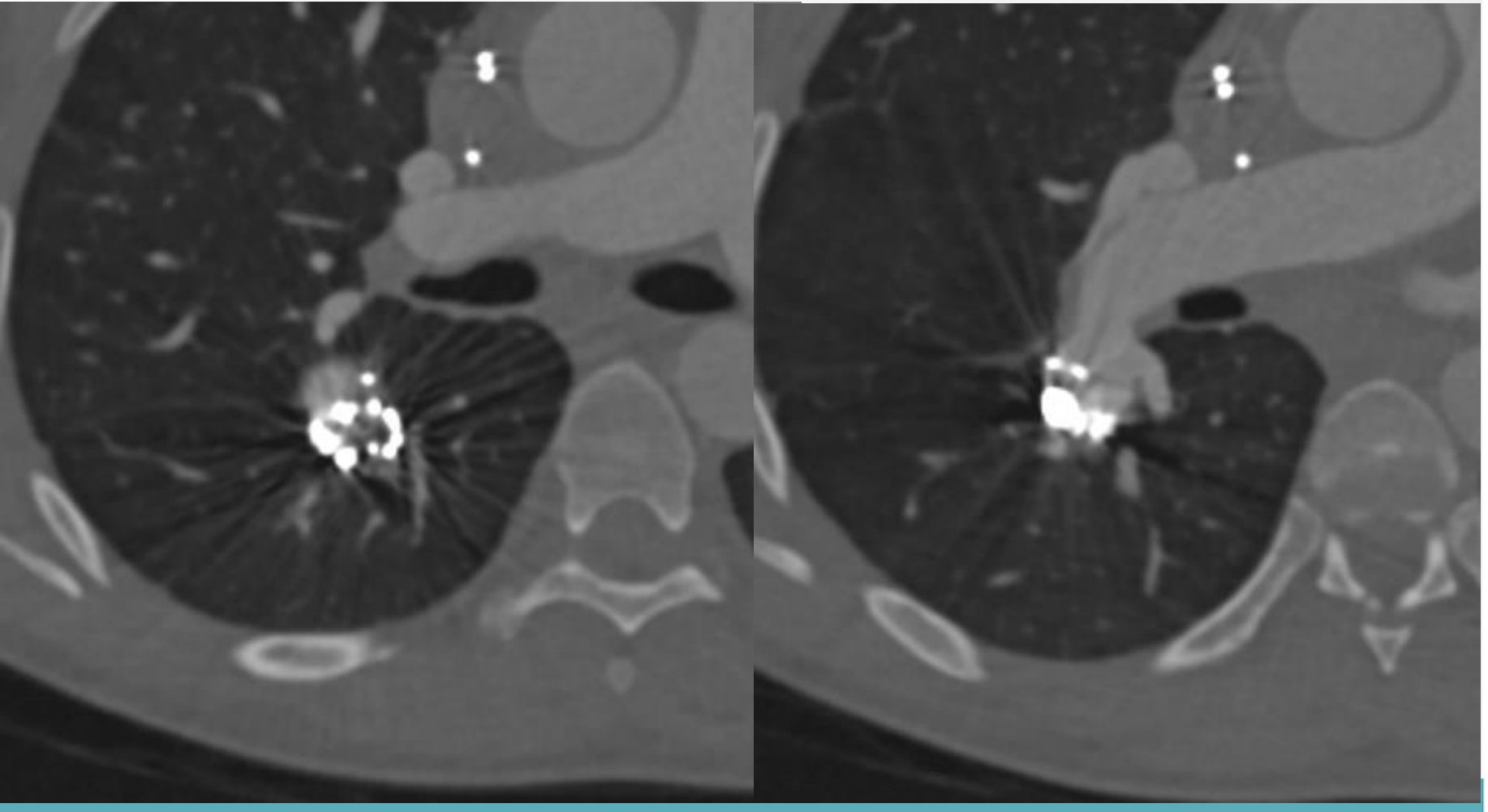


Course in Hospital

- Successful tricuspid valve replacement the following day
 - Follow up echo showed no vegetation or complication
- Ongoing antibiotics
- Left against medical advice approximately 10 days post-operatively



Follow-up CT



Follow-up CT Findings

- Small amount of filling in residual aneurysm neck
- Stable size
- Assessment somewhat limited due to steak artifact



Summary:

- Granulomatosis with polyangiitis
 - Diagnostic challenge
 - Multi-systemic disease of medium-sized vessels
 - Upper respiratory tract / pulmonary / renal
- Pulmonary artery pseudoaneurysms
 - Rare
 - Infectious / traumatic / vasculitic / congenital
- Endovascular management is mainstay



References:

- Ananthakrishnan, L., Sharma, N., & Kanne, J. P. (2009). Wegener's Granulomatosis in the Chest: High-Resolution CT Findings. *American Journal of Roentgenology*, 192(3), 676–682. <http://doi.org/10.2214/AJR.08.1837>
- Guillaume, B., Vendrell, A., Stefanovic, X., Thony, F., & Ferretti, G. R. (2017). Acquired pulmonary artery pseudoaneurysms: a pictorial review. *The British Journal of Radiology*, 90(1074), 20160783. <http://doi.org/10.1259/bjr.20160783>
- Lee, W.-K., Mossop, P. J., Little, A. F., Fitt, G. J., Vrazas, J. I., Hoang, J. K., & Hennessy, O. F. (2008). Infected (Mycotic) Aneurysms: Spectrum of Imaging Appearances and Management. *RadioGraphics*, 28(7), 1853–1868. <http://doi.org/10.1148/rg.287085054>
- Pakalniskis, M. G., Berg, A. D., Policeni, B. A., Gentry, L. R., Sato, Y., Moritani, T., & Smoker, W. R. K. (2015). The Many Faces of Granulomatosis With Polyangiitis: A Review of the Head and Neck Imaging Manifestations. *American Journal of Roentgenology*, 205(6), W619–W629. <http://doi.org/10.2214/AJR.14.13864>

