

# Neuroimaging Case-Based Review

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Brain Summit 2024: Current Trends in Neurology

UTSouthwestern
O'Donnell Brain Institute

UTSouthwestern Medical Center

# **Aims and Outline**

#### Goals:

- review select neurological disorders with classic imaging manifestations.
- treatable conditions, impactful patient outcomes

#### **Format:**

- Cased-based discussion
- Clinical vignettes, selected images
- Interactive polling platform engagement and self assessment

## How to Join

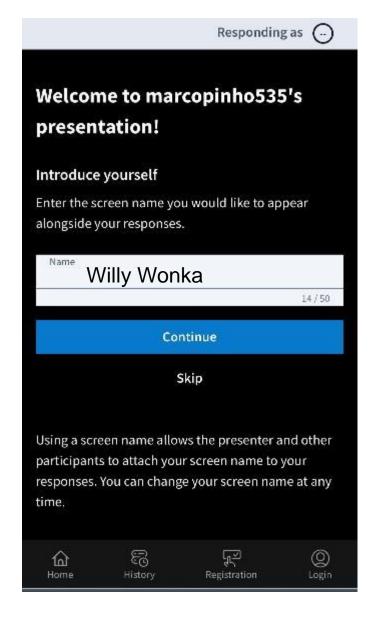
**QR** Code



### **Browser**



- 1 Go to PollEv.com
- 2 Enter MARCOPINHO535
- Respond to activity

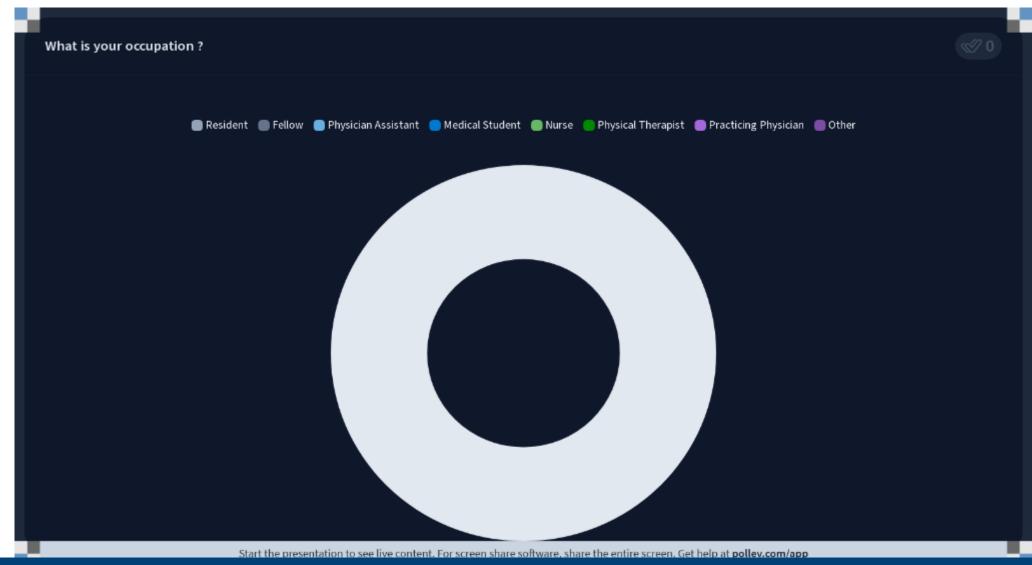


# Disclosures

- No financial relationships to disclose
- Polling tool just for engagement, don't take it too seriously

# Warm up!

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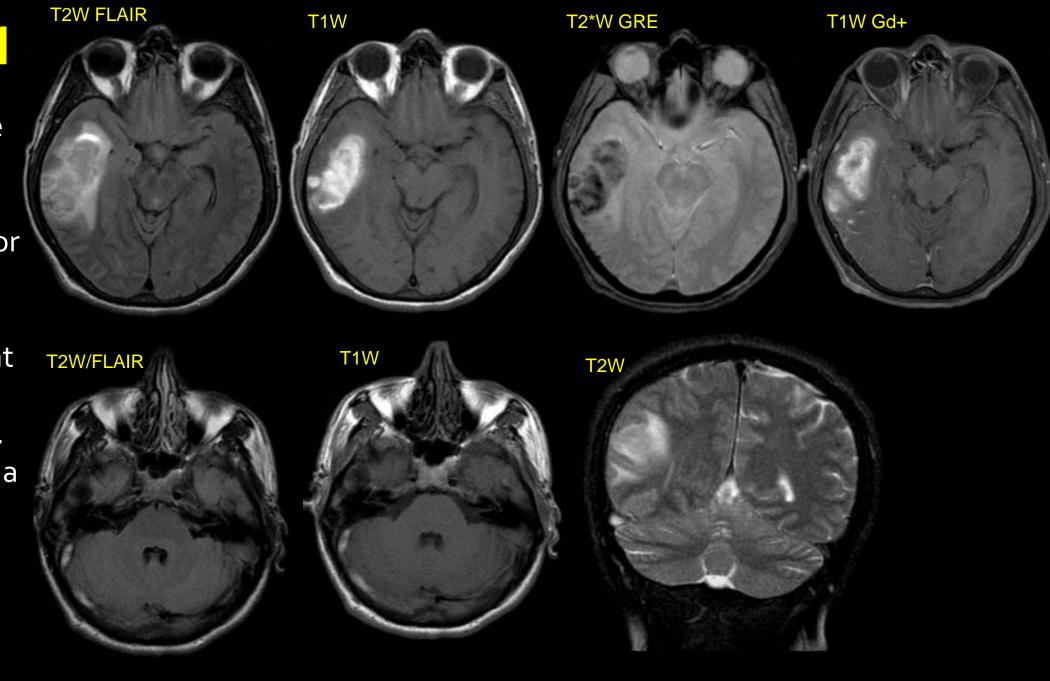


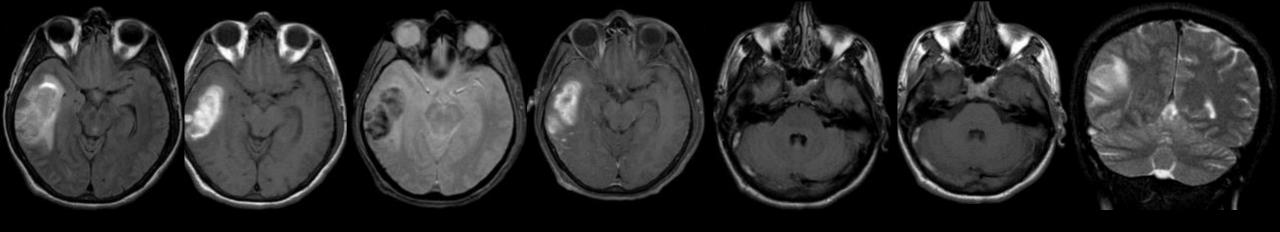
• 28 yo female

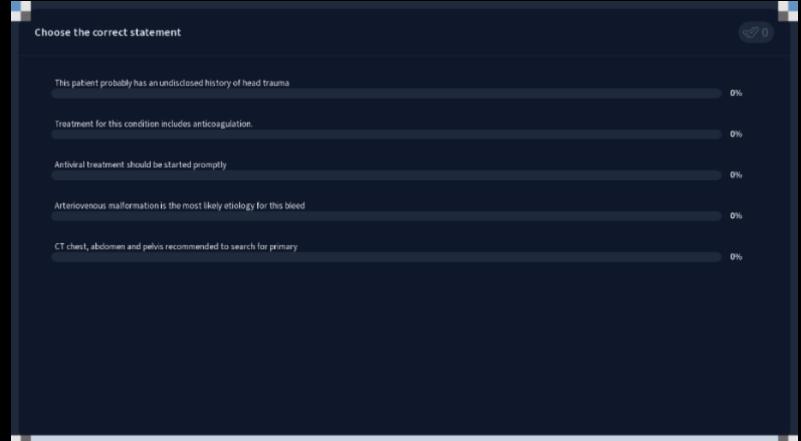
Worsening headaches for 10 days.

 No improvement with medications.

Negative CT a week ago.





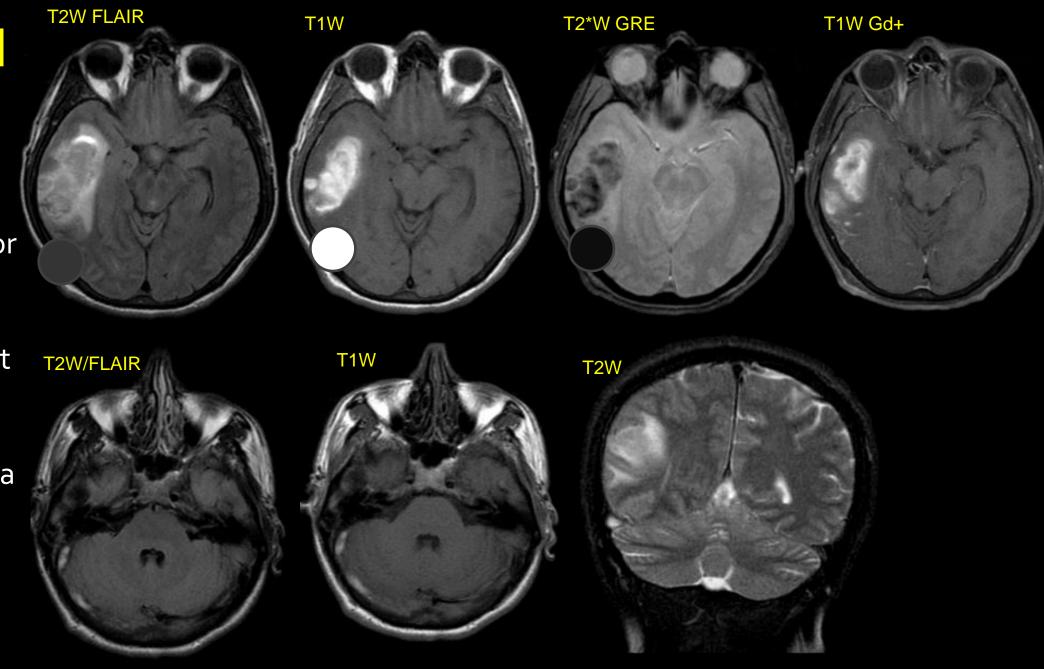


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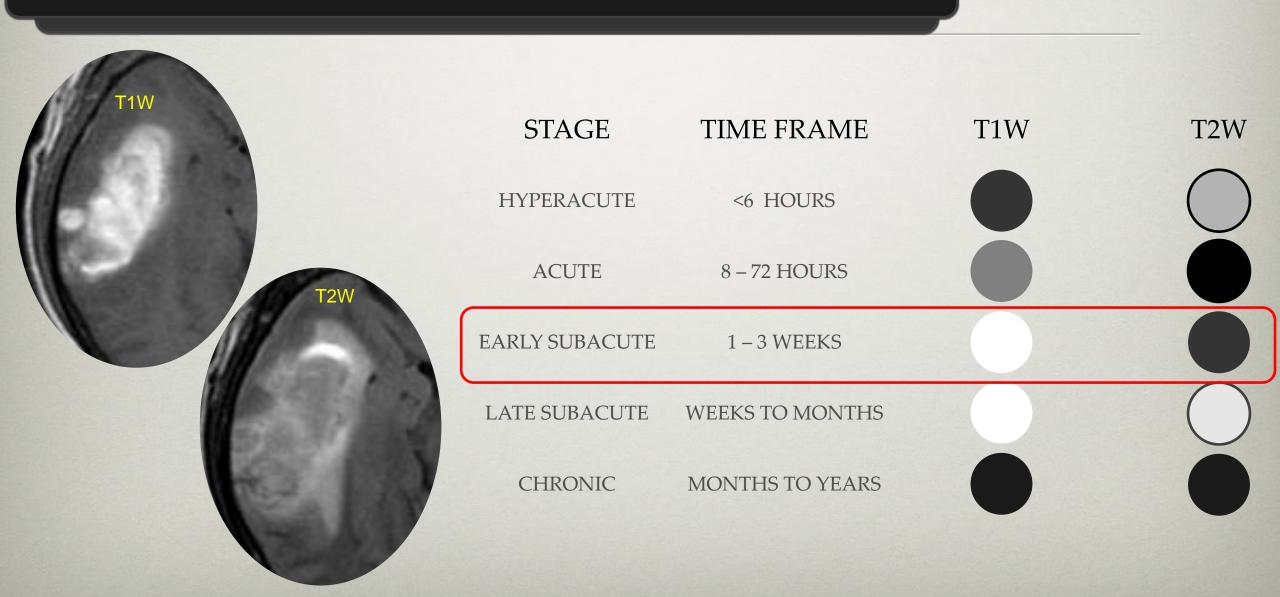
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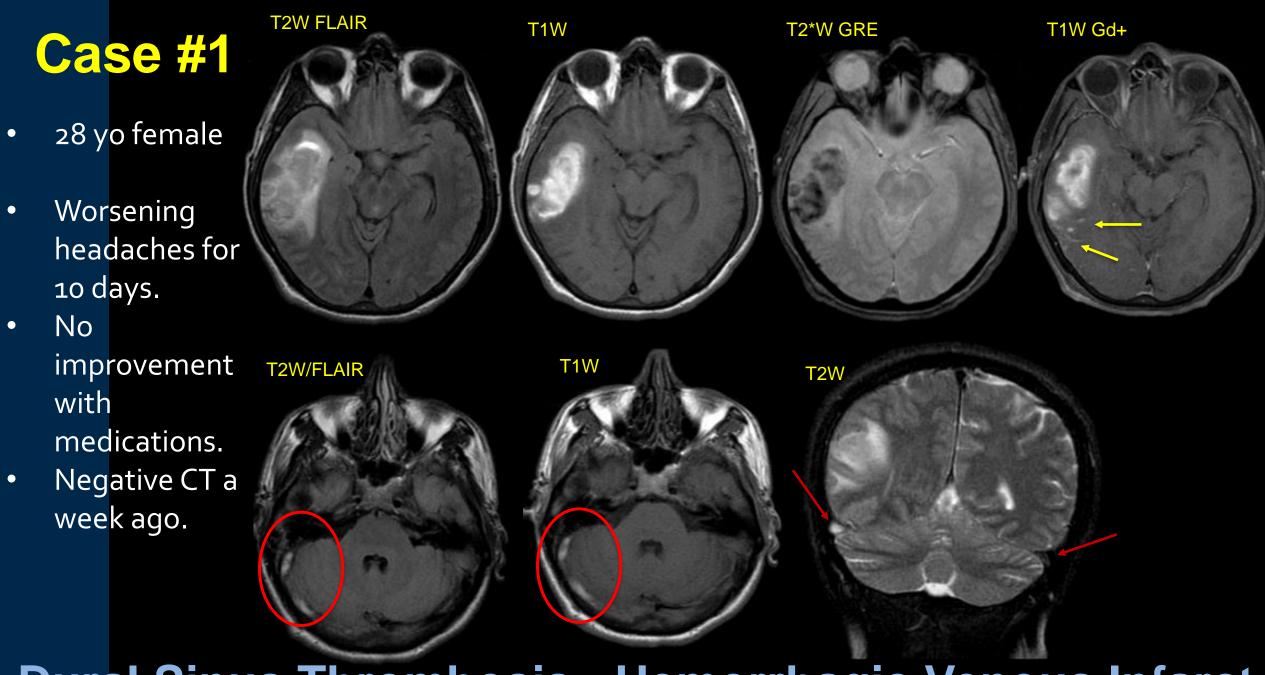
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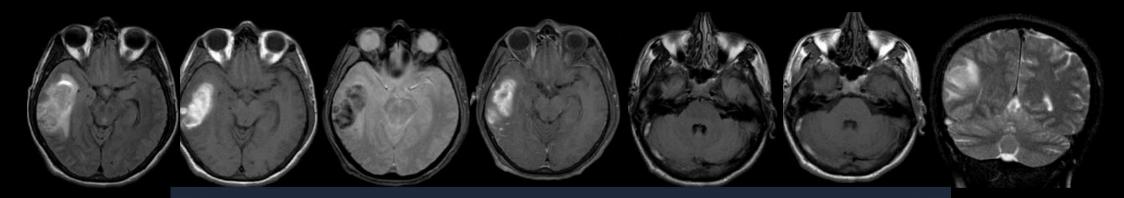


## Hemoglobin Degradation





**Dural Sinus Thrombosis - Hemorrhagic Venous Infarct** 



#### Choose the correct statement



This patient probably has an undisclosed history of head trauma



Treatment for this condition includes anticoagulation.

Antiviral treatment should be started promptly

Arteriovenous malformation is the most likely etiology for this bleed

CT chest, abdomen and pelvis recommended to search for primary



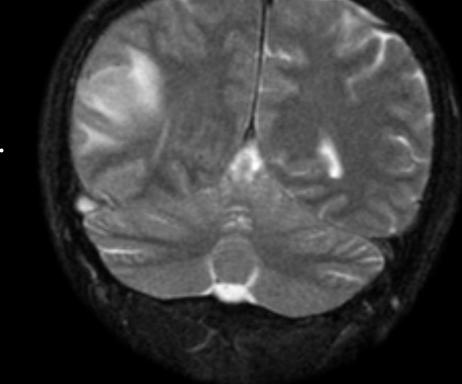






## Venous Sinus Thrombosis

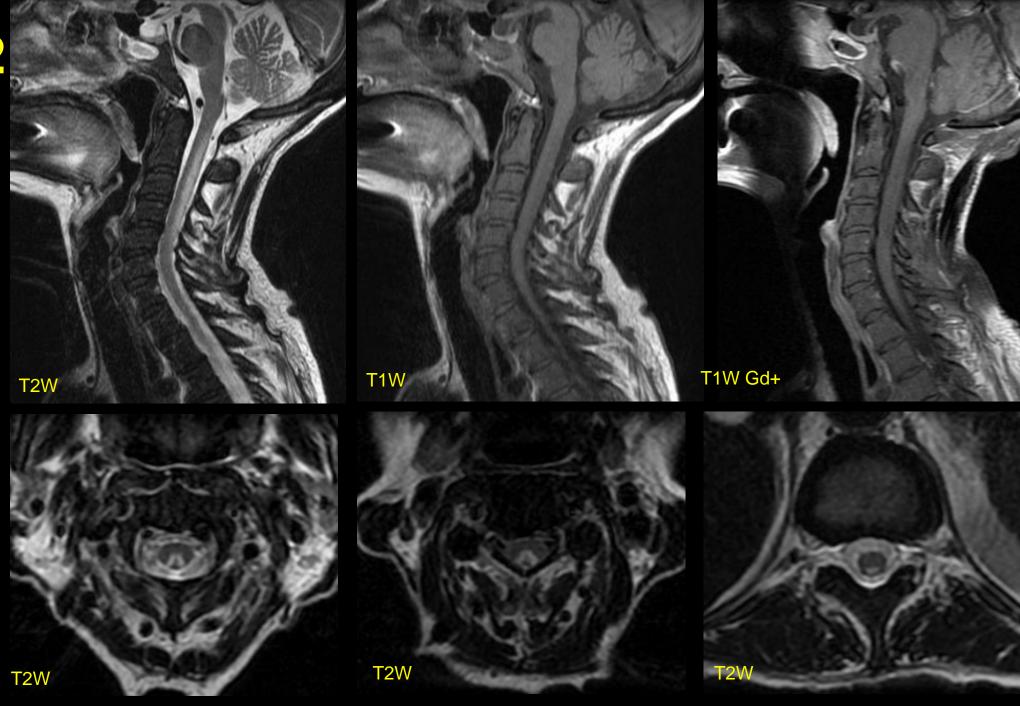
- Always search for venous disease:
  - unexplained parenchymal hemorrhage
  - infarcts in a non arterial distribution.
- Noncontrast MRI will often point to the correct diagnosis.
- MR Venography: confirmation and staging.



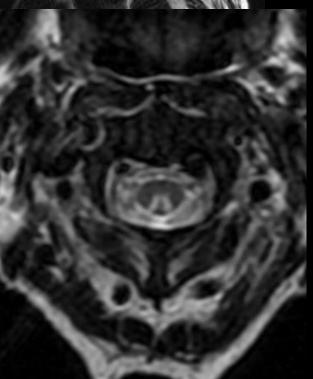
- Tx: heparin even with hemorrhagic venous infarction!
- Rarely catheter directed thrombolysis

40 yo M.
Progressive
paresthesia of the
hands and feet,
ataxia.

Hx of longstanding inflammatory bowel disease and prior surgeries.







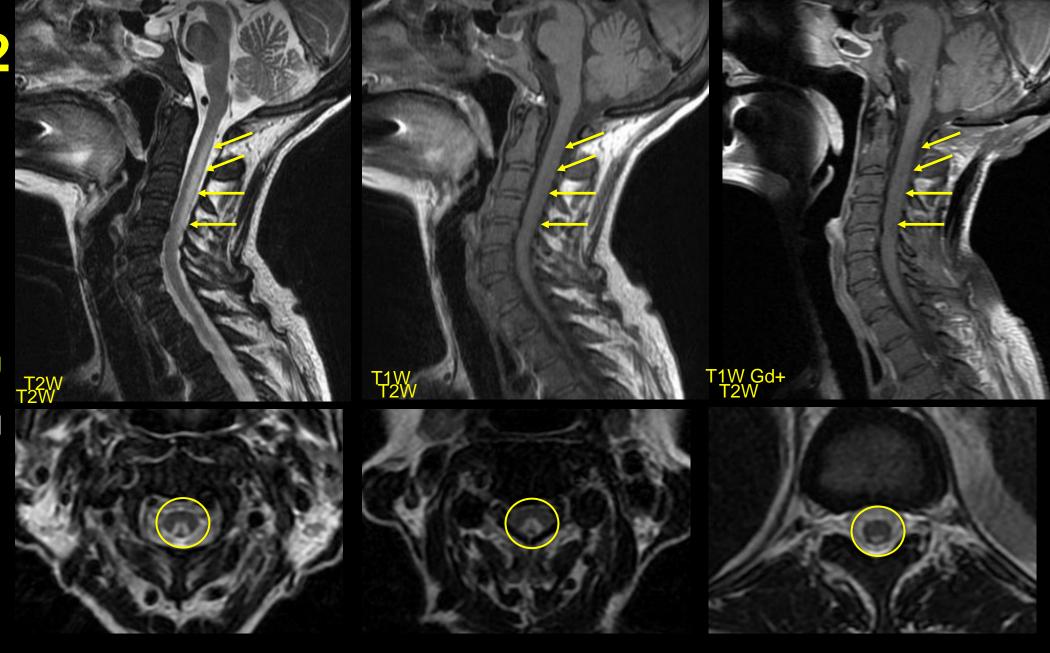
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## **Subacute Combined Degeneration – B12 Deficiency**

# Subacute Combined Degeneration

#### Pathophysiology

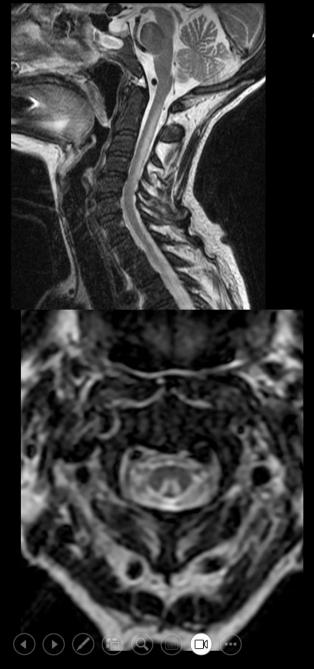
- B12 deficiency
- Risk factors: pernicious anemia, IBD, strict vegetarians.
- •Degradation posterior and lateral columns spinal cord
- Symmetrical myelopathy (distal proprioception and vibration with progression to all modalities, ataxia and distal weakness)

#### Imaging Findings

- Myelopathy posterior and lateral columns
- Inverted "V" sign, long segmental
- Reversible

#### Differential Diagnoses

- Copper or vit E deficiency: may look identical
- HIV and methotrexate induced myelopathy: may look identical
- Infectious: tabes dorsalis, herpes and HIV myelopathy
- Transverse Myelitis
- MS, Neuromyelitis Optica
- Cord Infarct



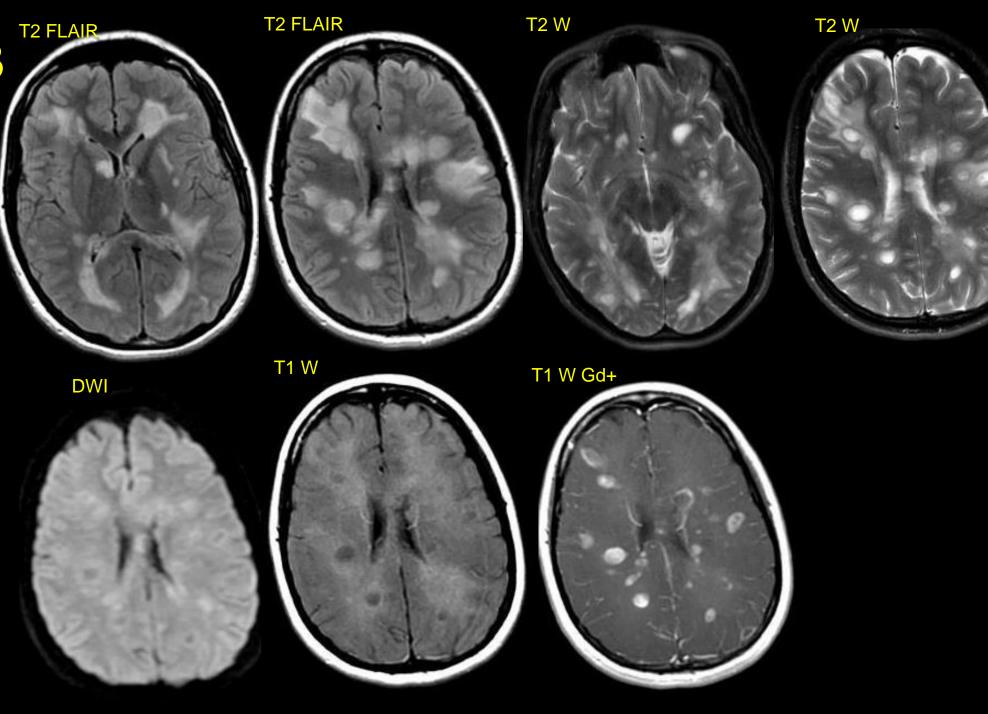
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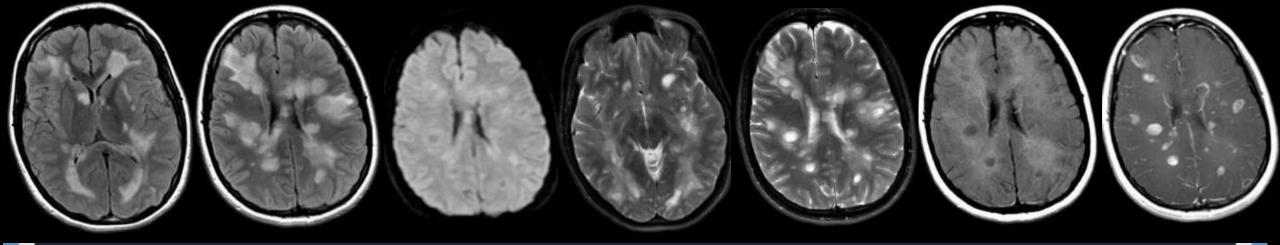
Hx of longstanding inflammatory bowel disease and prior surgeries.



 7 yo male, recovering from GI infection

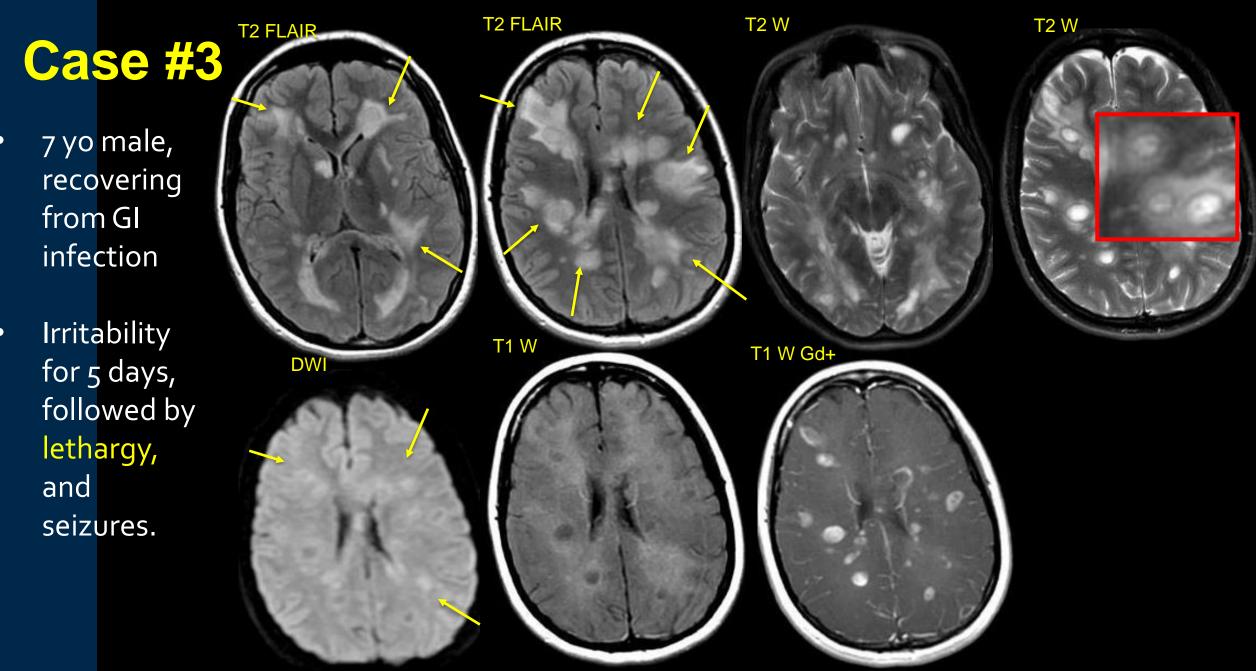
Irritability
 for 5 days,
 followed by
 lethargy,
 and
 seizures.



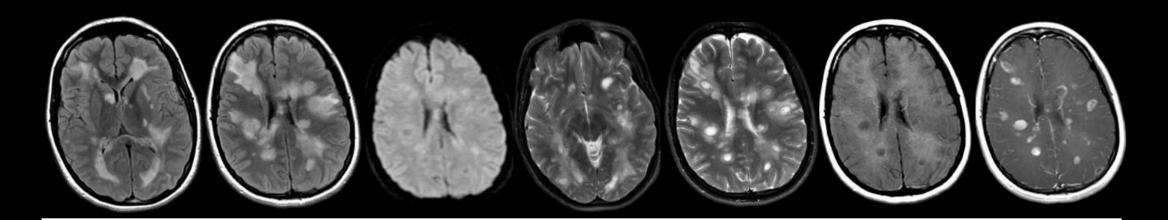


#### What statement is true about the presumed diagnosis for this case?

Antibiotic coverage should be started immediately, even before LP results	
	0%
Clinical and imaging findings favor de possibility of septic emboli	
	0%
This patient is at risk of hydrocephalus	
	0%
Given the combination of solid and ring enhancing lesions, metastatic disease is the top diagnostic consideration	
	0%
The prognosis for this patient is favorable with out without treatment, with expectation of good neurological recovery	
	0%



Acute Demyelinating Encephalopathy Syndrome



#### What statement is true about the presumed diagnosis for this case?



(E)

Antibiotic coverage should be started immediately, even before LP results

Clinical and imaging findings favor de possibility of septic emboli

This patient is at risk of hydrocephalus

Given the combination of solid and ring enhancing lesions, metastatic disease is the top diagnostic consideration

(D)



The prognosis for this patient is favorable with out without treatment, with expectation of good neurological recovery

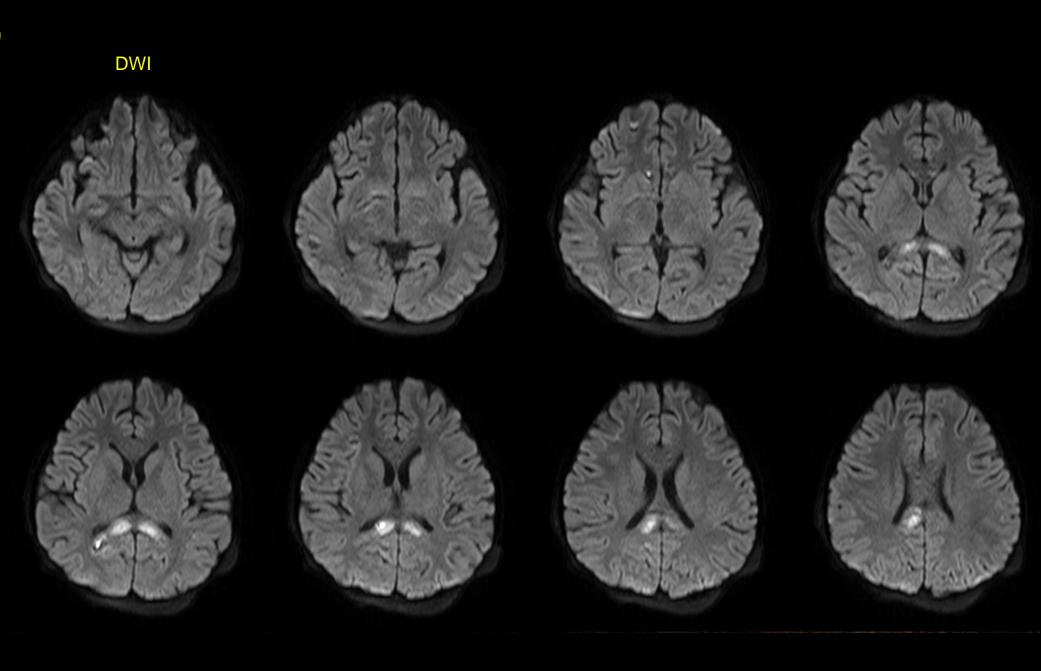
# Acute Disseminated Encephalopathy Syndrome

• Monophasic acute inflammation and demyelination associated with multifocal neurological symptoms.

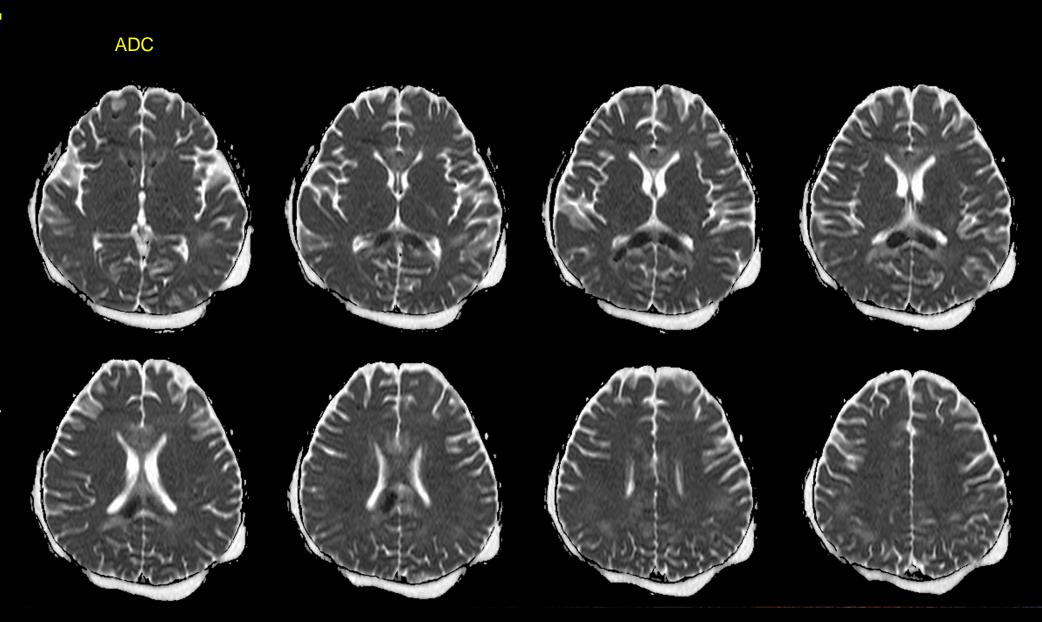
- Debated pathogenesis
  - autoimmune mechanism
  - environmental stimulus + genetically susceptible
- Recovery phase (~1 month) nonspecific systemic infection (75%)
  - Postinfections Encephalomyelitis
  - postvaccinal (< 5%)</li>
  - Common presentation of MOGAD in pediatric patients
- Clinical characteristics: acute/subacute presentation, multifocal neurological symptoms, encephalopathy
- Imaging: multifocal tumefactive WM lesions, enhancement, basal ganglia, little or no mass effect, no dissemination in time.



- 11 yo female, transfer from outside hospital
- Admitted due to MCC, hit while riding a bike
- Multiple rib and limb fractures
- Normal Head CT admission
- Acute
   neurological
   deterioration 5
   days post
   admission



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11 yo female, transfer from outside hospital

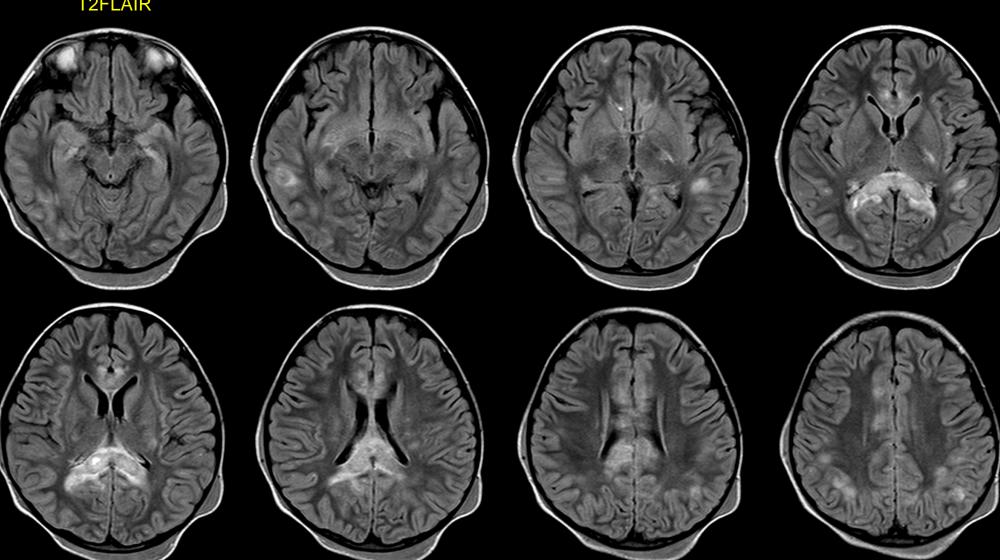
Admitted due to MCC, hit while riding a bike

Multiple rib and limb fractures

Normal Head CT admission

Acute neurological deterioration 5 days post admission

T2FLAIR

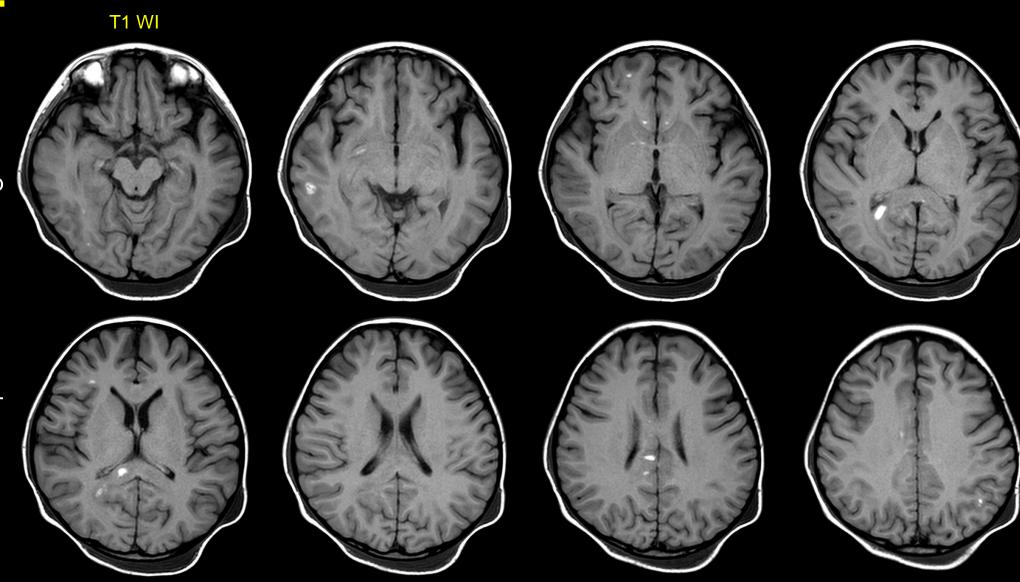


 11 yo female, transfer from outside hospital

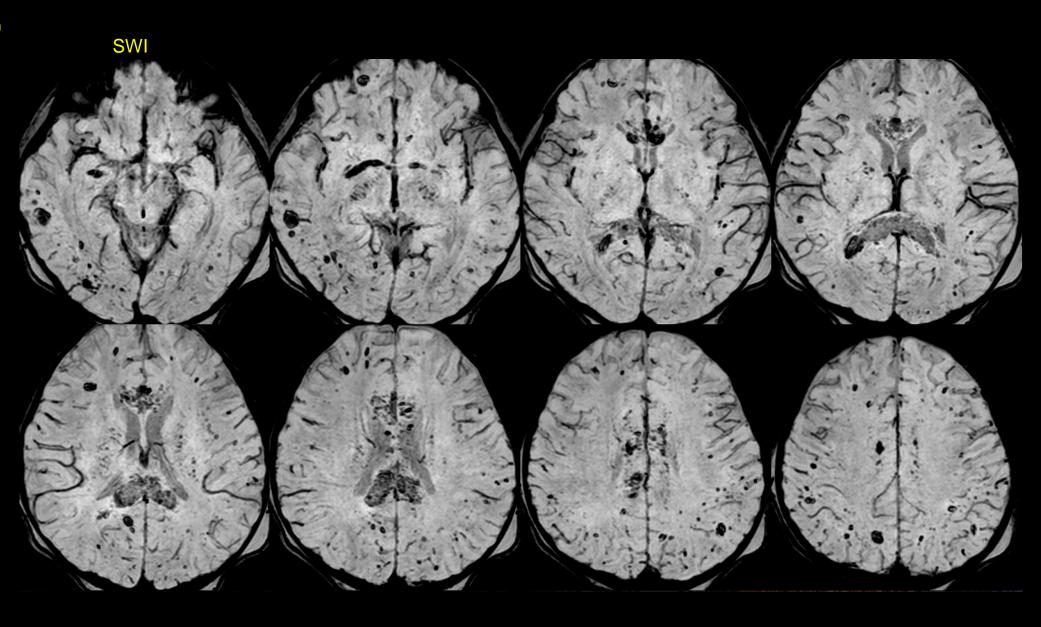
 Admitted due to MCC, hit while riding a bike

 Multiple rib and limb fractures

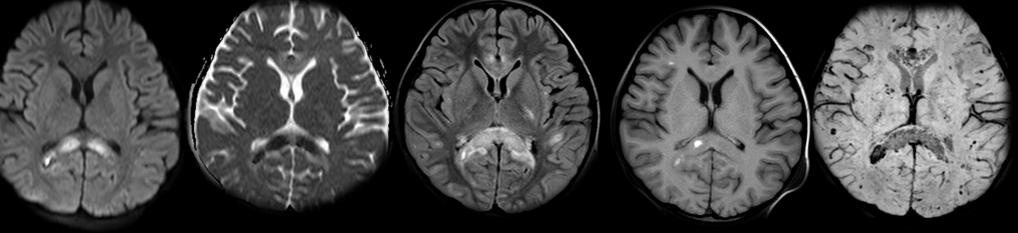
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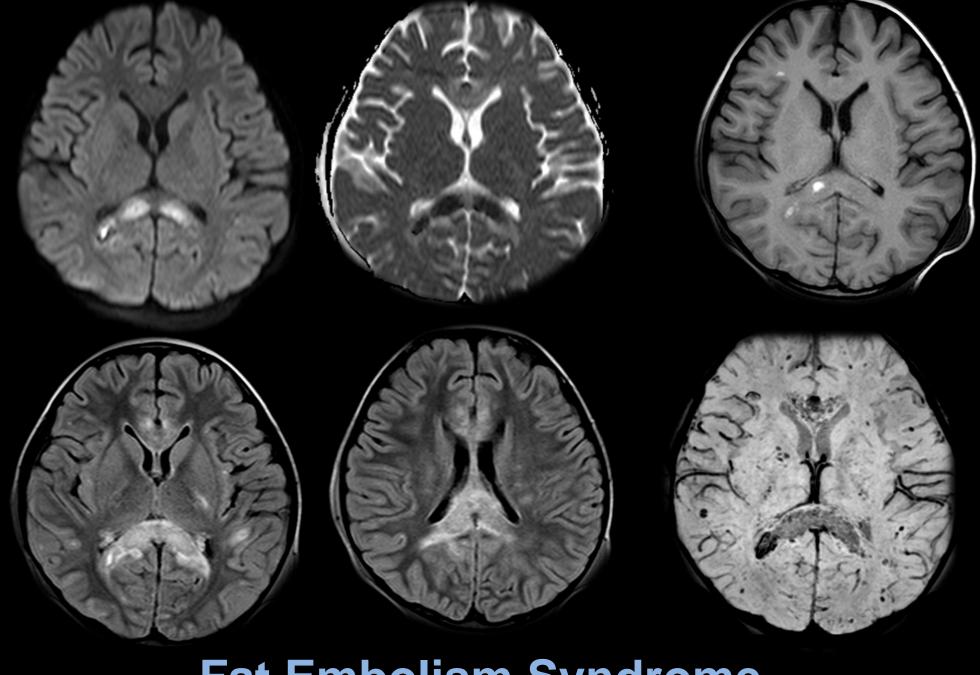


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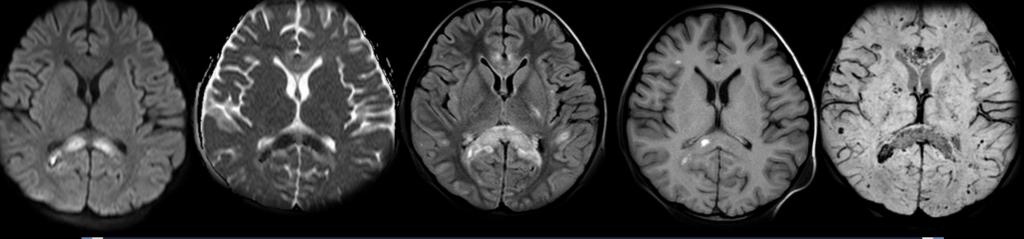
## Select the correct statement for this condition: The primary mechanism for this imaging abnormality is a coagulopathy 0% Pulmonary and cutaneous manifestations are not usually present in this clinical scenario 0% Vascular imaging is recommended and will likely disclose a central source of embolism 0% CSF tap highly recommended to identify the causative infectious agent 0% Treatment for this condition is mainly supportive medical care 0%

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**Fat Embolism Syndrome** 

- 11 yo female, transfer from outside hospital
- Admitted due to MCC, hit while riding a bike
- Multiple rib and limb fractures
- Normal Head CT admission
- Acute
   neurological
   deterioration 5
   days post
   admission





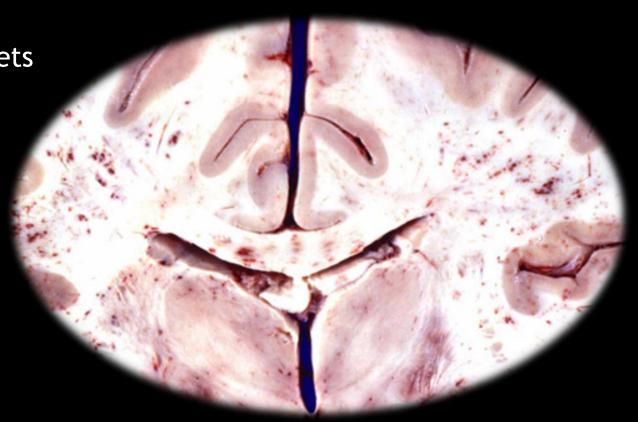
# Fat Embolism Syndrome

Occlusion of small CNS vessels by fat droplets

- Traumatic injury long bones and pelvis
- Subclinical fat embolism: common
- Clinical manifestations: 3-10%

### Classical Triad

- Pulmonary dyspnea
- Cutaneous petechial rash
- CNS symptoms



# Fat Embolism Syndrome

- Occlusion of small CNS vessels by fat droplets
- Traumatic injury long bones and pelvis
- Subclinical fat embolism: common
- Clinical manifestations: 3-10%

- Classical Triad
  - Pulmonary dyspnea
  - Cutaneous petechial rash
  - CNS symptoms

# Diagnostic Criteria (2 major or 1 major and 4 minor)

Major criteria	Petechial rash Respiratory insufficiency Cerebral involvement
Minor criteria	Tachycardia
	Fever
	Retinal changes
	Jaundice
	Renal signs
	Thrombocytopenia
	Anaemia
	High ESR
	Fat macroglobinemia

# Fat Embolism Syndrome

## Computed Tomography

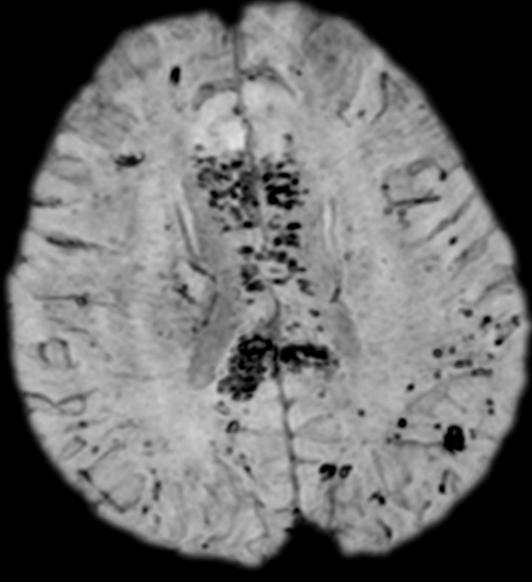
- Usually normal
- Patchy hypodensity, petechial Hemorrhages

#### MRI

- T2WI: patchy areas of nonspehyperintensity
- SWI: widespread microhemorrhages
- DWI: multifocal punctate pattern

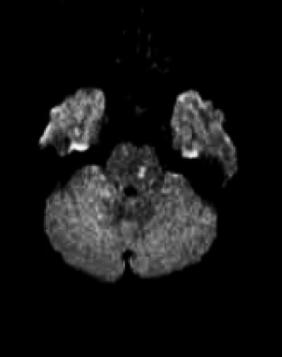
#### Dd

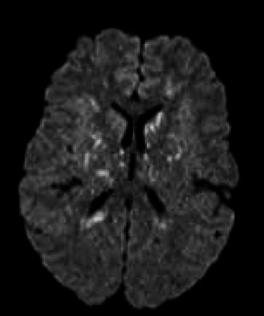
- Diffuse Axonal Injury
- Cardiogenic/septic cerebral emboli
- Vasculitis
- Thrombotic thrombocytopenic purpura

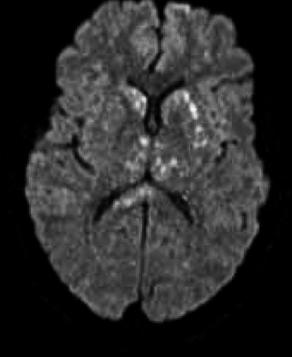


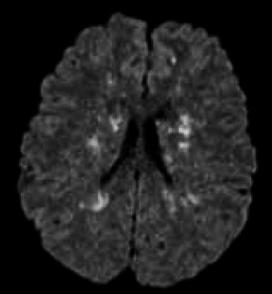
# Starfield Pattern





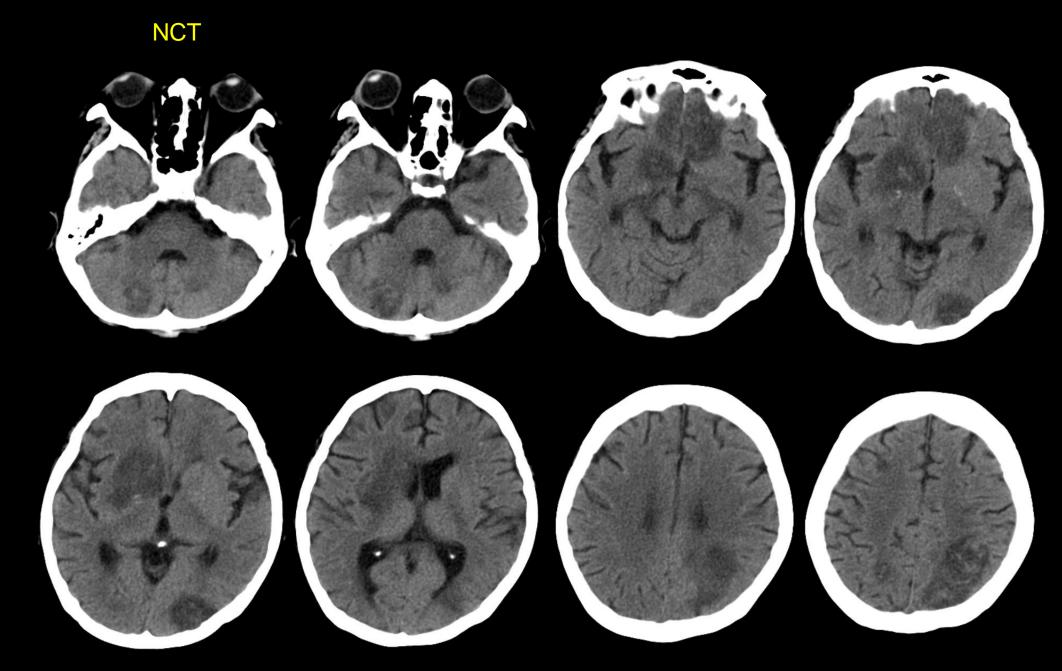


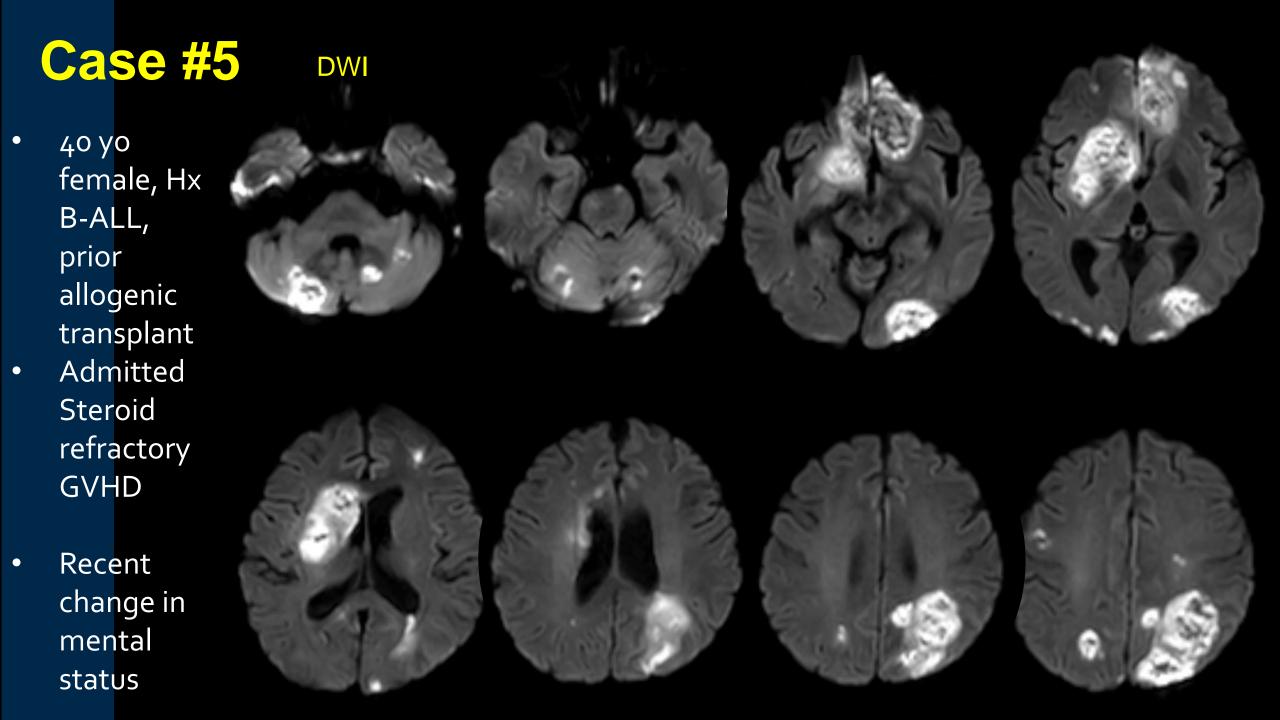






- 40 yo
   female, Hx
   B-ALL,
   prior
   allogenic
   transplant
- Steroid refractoryGVHD
- Recent change in mental status



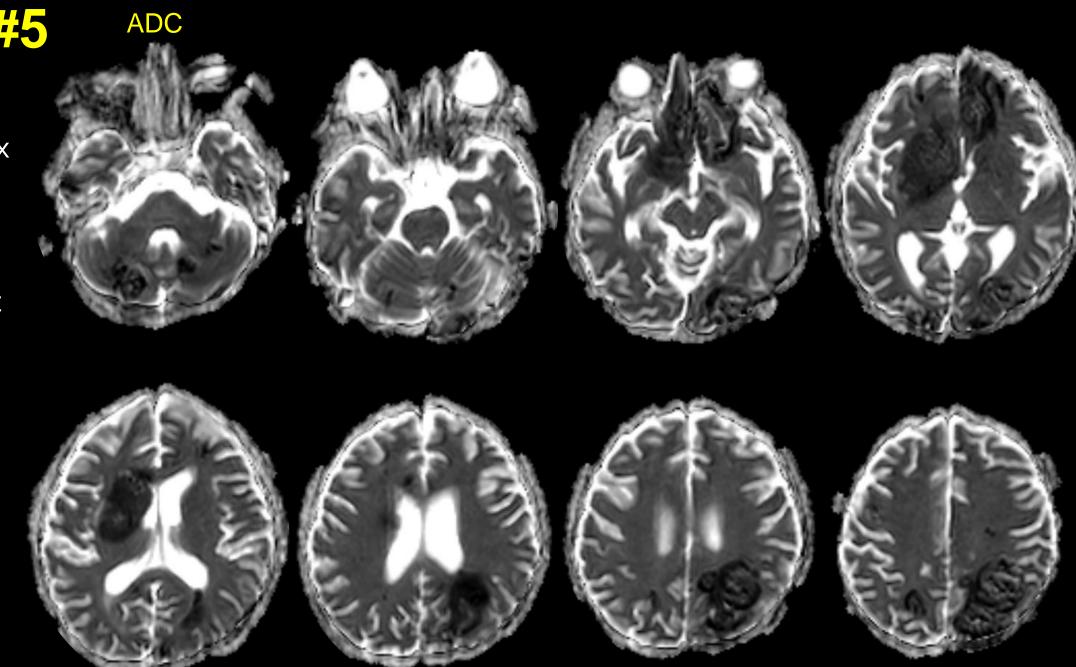


Case #5

40 yo
female, Hx

female, Hx B-ALL, prior allogenic transplant

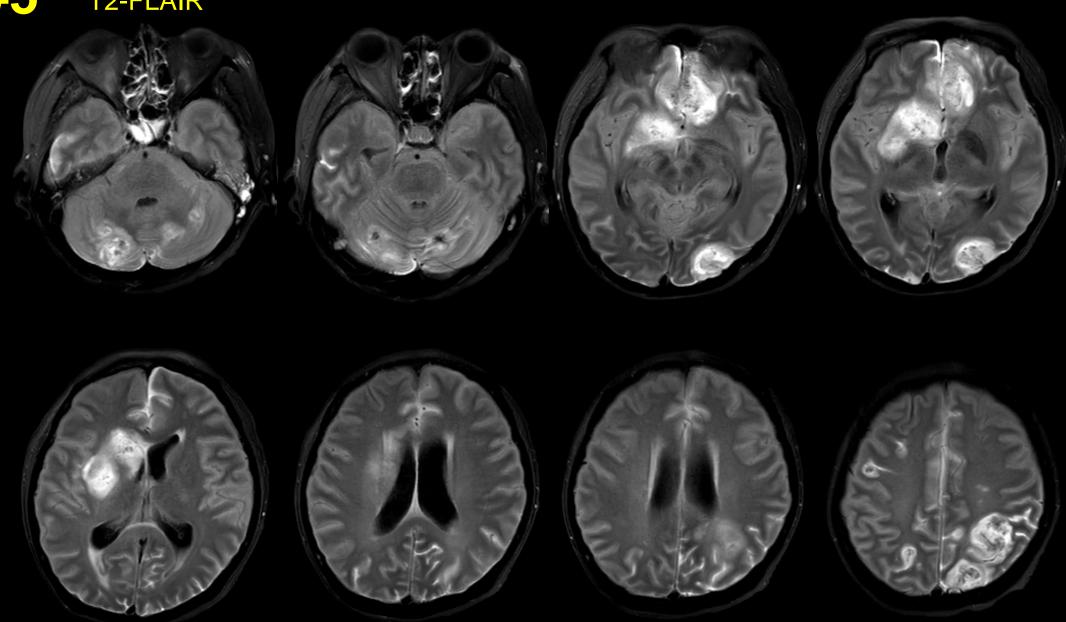
Admitted
 Steroid
 refractory
 GVHD



## Case #5 T2-FLAIR

40 yo
female, Hx
B-ALL,
prior
allogenic
transplant

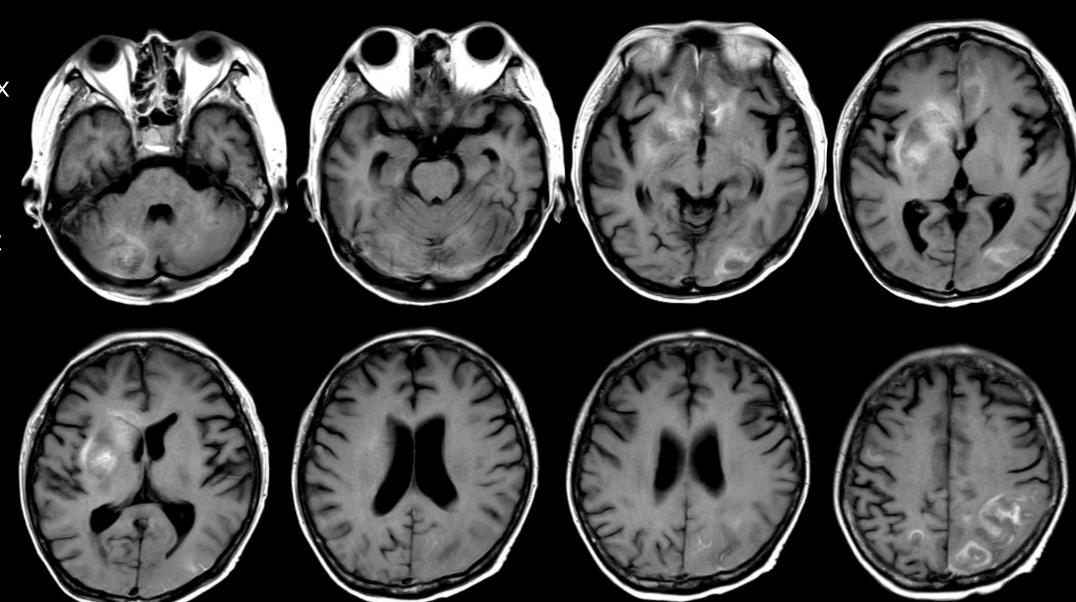
Admitted
 Steroid
 refractory
 GVHD



## Case #5 TI-WI

• 40 yo
female, Hx
B-ALL,
prior
allogenic
transplant

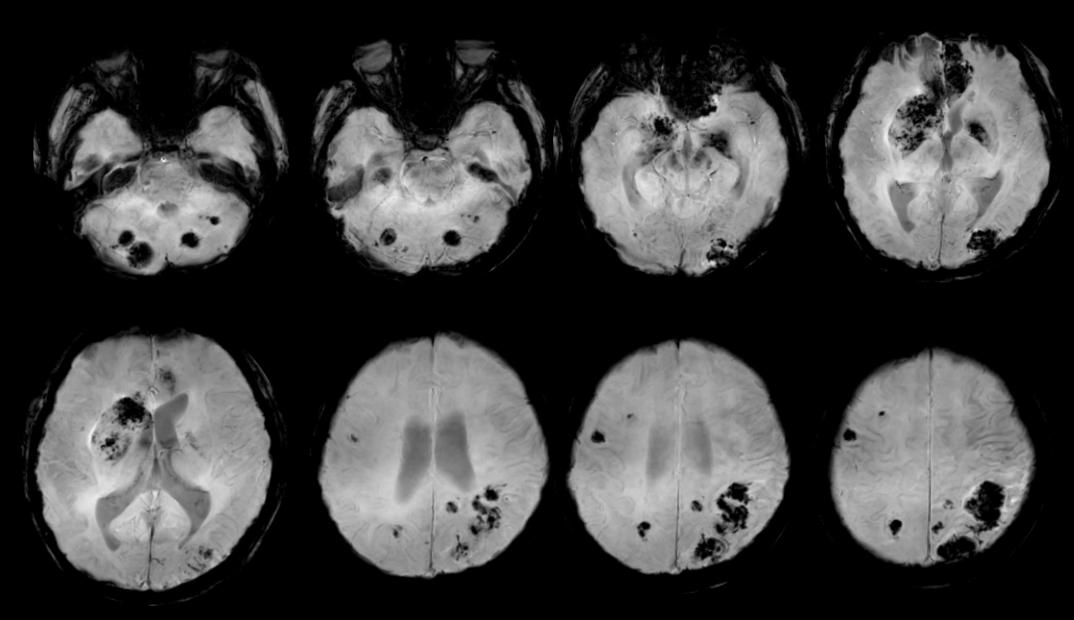
Admitted
 Steroid
 refractory
 GVHD



SWI

40 yo
 female, Hx
 B-ALL,
 prior
 allogenic
 transplant

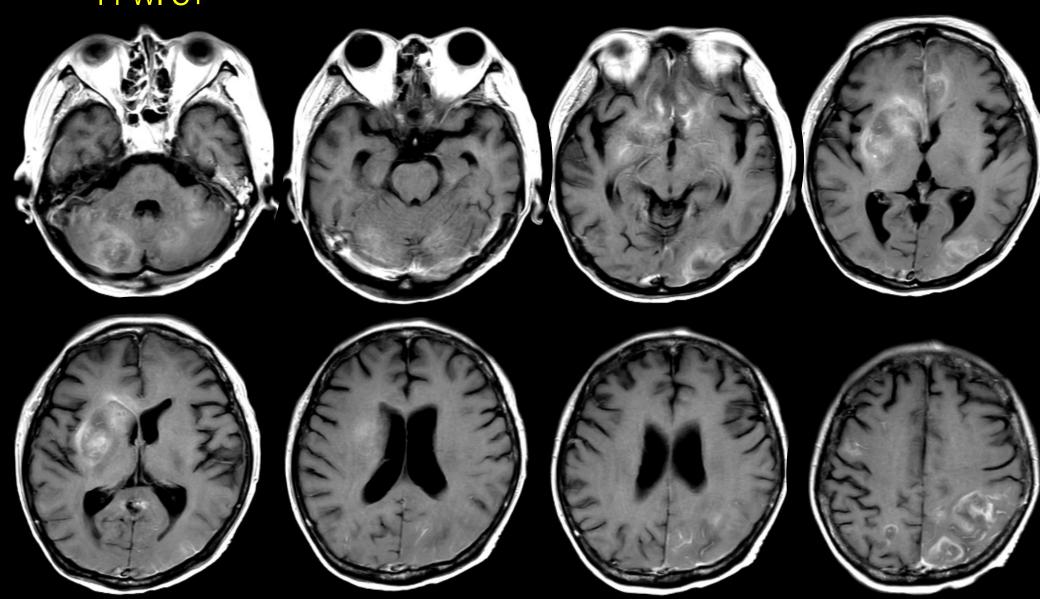
Admitted
 Steroid
 refractory
 GVHD



T1 WI C+

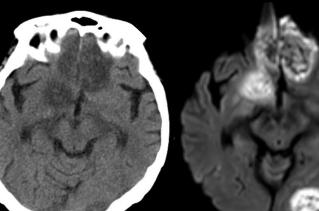
40 yo
 female, Hx
 B-ALL,
 prior
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 transplant

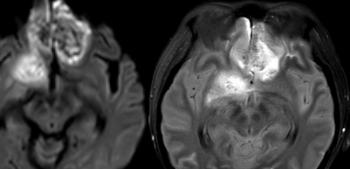
AdmittedSteroidrefractoryGVHD

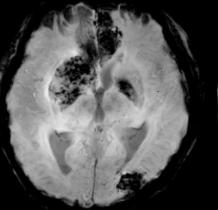


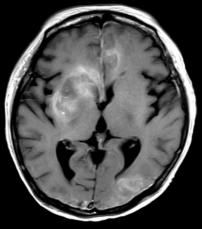
T2-FLAIR Case #5 NCT ADC DWI 40 yo female, Hx B-ALL, prior allogenic transplant Admitted T1WI Subtrac T1WI C+ T1WI SWI Steroid refractory **GVHD** Recent change in mental status

- 40 yo
  female, Hx
  B-ALL,
  prior
  allogenic
  transplant
- Admitted Steroid refractory GVHD
- Recent change in mental status









#### Please select the incorrect statement:

This is a treatable condition, however, the mortality rates are very high.	
	0%
Hemorrhagic components are very typical for this entity, as seen in this case	0%
	0 70
Methotrexate based intrathecal chemotherapy should be strongly considered	
	0%
Vascular Imaging is likely of no benefit in this case	0%
	0.70
Chest CT may offer clues to the correct diagnosis	
	0%

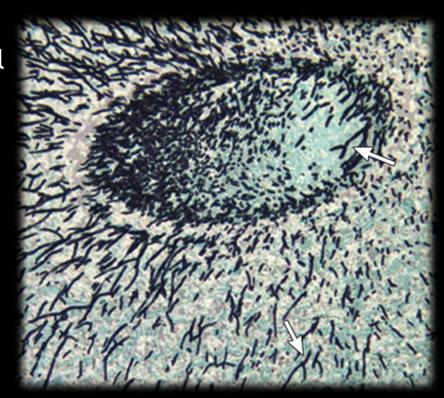
T2-FLAIR Case #5 **NCT** DWI **ADC** 40 yo female, Hx B-ALL, prior allogenic transplant Admitted T1WI Subtrac T1WI C+ T1WI **SWI** Steroid refractory **GVHD** Recent change in mental Angioinvasive Fungal Cerebritis status

## Angioinvasive Fungal Cerebritis

- Life threatening CNS infection, usually Aspergillus or Mucor
- Immunocompromised, hematogenous versus direct extension
- Poor prognosis with ~ very high mortality for cerebral aspergillosis
- Pulmonary aspergillosis treatable

#### Mechanism

- Invasion walls of both small and large blood vessels
- Thrombosis, infarction, hemorrhage, necrosis
- Abscess if immune response partially preserved

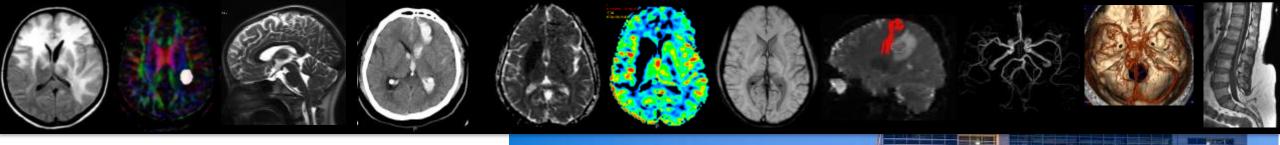


Tempkin et al 2006

40 yo
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Admitted Steroid refractory GVHD





# Thank you for participating!

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