

Learning Objectives

- 1) Present an unique case of spinal myoclonus
- 2) To increase awareness of prevalence, risk factors, severity, and adverse effect to myelograms
- 3) To review literature and treatment plans for adverse effects to myelograms including ascending tonic-clonic seizure syndrome

Patient Presentation

48 year old female with a pmhx of hypothyroidism, diabetes mellitus, and chronic back pain presented to the emergency department two days after receiving a lumbar CT myelogram for lower extremity muscle spasms and cramps. On evaluation, the patient was very agitated with acute onset of muscle spasms screaming that she was in a lot of pain. Denied any fever, chills, chest pain, shortness of breath, nausea, vomiting, and diarrhea.

Physical Exam

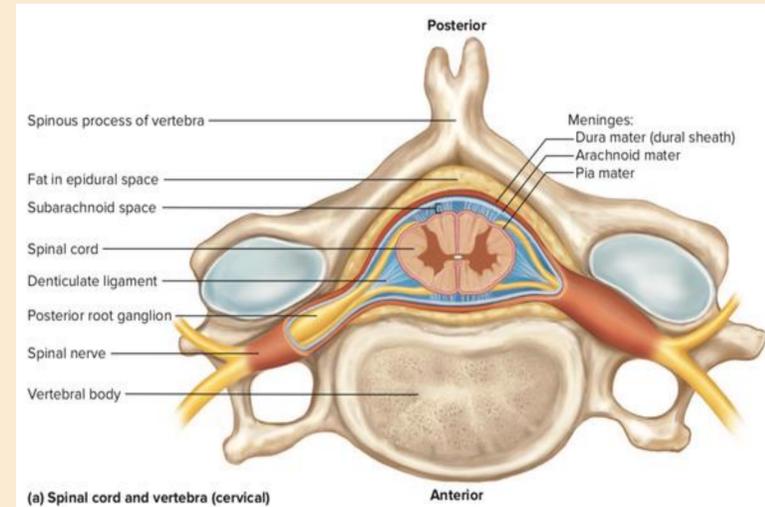
Vitals: Temp: 98.2, **BP: 193/82**, **HR: 155**, **RR: 28**, O₂: 95% on 2L NC
 General: In moderate distress, having active lower extremity muscle spasms
 HEENT: Normocephalic, atraumatic, PERRLA, EOMI
 Cardio: **Tachycardic**, distinct S₁ + S₂, No murmurs, rubs, or gallops, +1 lower extremity edema bilaterally. Pulses equal and intact.
 Respiratory: **Wheezing bilaterally**. Increased work of breathing, currently on 2L NC
 GI: Soft, non-tender to palpation, +bowel sounds, no guarding or rebound tenderness
 MSK: **Actively having muscles spasms, able to mover her upper extremities without problem, lower extremities rigid and difficult to bend**



Lab Values

142	113	27	13.5	CPK: 1504
4.4	14	0.91	311	BNP: 160
			40.3	D-dimer: 724
				Lactic Acid: 1.6

Myelogram and Spinal Cord Anatomy



Inadvertent Intrathecal Use of Contrast Media for Myelography

- There have been over 14 cases of adverse effects from contrast media following a myelography
- Normally seen with ionic contrast media
- The clinical presentation widely varies from the spread, location, and severity but there are several case studies highlighting tonic-clonic movements
- Reaction occurs with 30 minutes to 72 hours after contrast initiation with painful spasms of the lower extremities associated with spasms initially and then contraction
- Frequently elicited by merely touching the affected extremity
- Ascending paralysis that can lead to a generalized seizure and loss of consciousness
- Autopsy and histopathologic examination following a case report showed localized, cellular reaction of the brain and spinal cord with brain edema
- Theory that the contrast may change or destabilize the membrane potential of affected neurons resulting in abnormal function

Hospital Course and Follow Up

- Patient was intubated for airway protection and given multiple doses of benzodiazepines to help control the muscle spasms and prevent any ascending tonic-clonic seizures from occurring
- Given continual supportive care with IV fluids, benzodiazepines, and frequent neuro checks
- EEG negative for any seizure activity
- Patient was able to be extubated 48 hours after the event and able to move all extremities at that time without any muscle spasms

Ascending Tonic-Clonic Seizure (ACTS) Syndrome

- 1) The patient has undergone a myelographic examination during the preceding 6 hours
- 2) Painful tonic-clonic muscular spasm are present
- 3) The spasms occur in paroxysms lasting approximately 15 – 30 minutes
- 4) The spasms may be precipitated by touching or handling the affected limbs
- 5) Following a lumbar myelogram, the spasms spread from the lower limbs in an ascending pattern to involve also the trunk and upper limbs
- 6) The patient is at first fully conscious, but becomes stuporous or comatous as the symptoms ascend in a cephalad direction



Treatment

- Should be aimed at controlling muscle spasms and seizures to avoid pain, fractures, hyperthermia, acidosis, and muscular injury. Early intubation has been successful throughout the case studies.
- Intravenous benzodiazepines should be used first and if this does not relieve symptoms, use neuromuscular blocking agents
- Recommend continual treatment for 48 hours prior to stopping any medications
- The procedure will mask external signs of seizure, an EEG may be helpful to distinguish between ATCS syndrome from a seizure
- Steroids might have a role in preventing cerebral edema but there is no scientific evidence
- Place patient in the “head and trunk up” position as the contrast is heavier than CSF
- In refractory cases, anesthesia can perform a careful CSF exchange-washout

Take Home Points

- There are serious life threatening adverse reactions to contrast media myelograms that health care professionals need to be aware of and learn how to treat
- The ionic contrast can affect neurons leading to abnormal function causing muscle spasms, ascending tonic-clonic seizure, and / or a coma
- No definitive treatment is recommended yet but early intervention, benzodiazepines, close clinical monitoring, and in refractory cases “CSF exchange-washout” is recommended

References

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