

Managing Nerve Injury Risk During Cryoablation Procedures

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Introduction

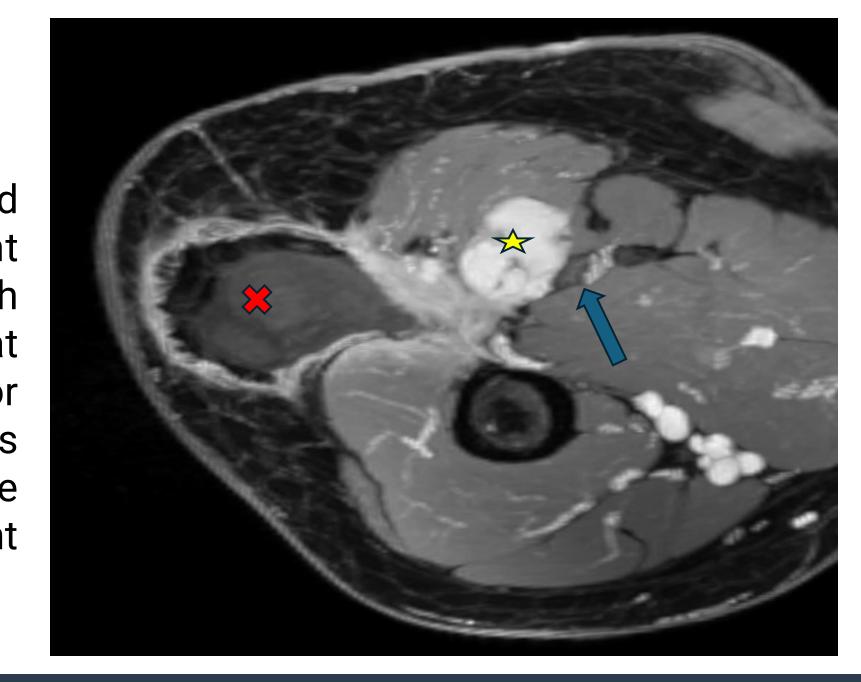
- Cryoablation is a minimally invasive technique that uses compressed argon gas to freeze tumors to temperatures below -20 °C, causing tumor necrosis.
- It is an established treatment for numerous pathologies including renal, pulmonary, and musculoskeletal tumors.
- A key challenge arises when lesions are adjacent to neurovascular structures, where freezing can result in neuropathy and long-term functional deficits (1).
- This learning exhibit aims to educate interventional radiologists on practices for preventing nerve injury during cryoablation procedures.

Key Anatomical Considerations

- Proximity to major nerve pathways
- Sciatic nerve, brachial plexus, femoral nerve, spinal nerve roots, and peripheral branches.
- A margin of >1 cm is recommended between the cryoablation ice ball and critical nerve structures (2).
- Vascular structures can act as **heat sinks**, increasing risk of injury during freezing (3).
- Anatomic variants can increase complexity and should be assessed on pre-procedural imaging.

Symptomatic Desmoid Tumor

Figure 1. T1-weighted contrast-enhanced axial MRI of a desmoid tumor. The patient previously underwent ablation, with nonenhancing tissue and surrounding fat necrosis (red x). Recurrent/residual tumor is visible along the medial edge as enhancing tissue (yellow star). The blue arrow highlights the adjacent neurovascular bundle.



Techniques and Clinical Pearls

- 1. Pre-procedural Imaging & Cryoprobe Trajectory
- a. Use CT and/or MRI to map the lesion and identify nearby nerves and vessels.
- b. Plan a trajectory that maximizes tumor coverage and minimizes proximity to neurovascular structures.
- 2. Hydrodissection and CO₂ Pneumodissection
 - a. Hydrodissection: Inject **sterile saline or D5W** to displace nerves away from the ablation zone, aiming for a ≥1 cm margin.
 - b. CO₂ pneumodissection: Is less conductive and used for regions where **fluid can alter anatomy**.
- 3. Intraoperative Neuromonitoring (EMG, SSEP, MEP)
 - a. Real time **detection of nerve signal changes** allows the interventional radiologist to adjust probe position or stop treatment to prevent permanent damage.
- b. Electromyography (EMG): Detects spontaneous **muscle activity** when the ice ball approaches a motor nerve
- c. Somatosensory Evoked Potentials (SSEP): Monitors **sensory pathway** conduction during freezing, identifying changes in peripheral sensory nerves.
- d. Motor Evoked Potentials (MEP): Loss or reduction of amplitude may indicate **motor deficit** due to cryoablation.
- 4. Freeze-Thaw Cycle Modifications
- a. Use **shorter freeze times** or single freeze cycles when nerves are within high-risk proximity.
- b. Continuously monitor ice ball expansion with real-time imaging.

Pre-Ablation MEPs

MEPs During 1st Ablation

Record 1

Figure 2. Intraoperative neurophysiological monitoring (IONM) traces for motor evoked potentials (MEPs) and somatosensory evoked potentials (SSEPs) across multiple muscle and nerve distributions. Pre-ablation baseline recordings (left) are compared to signals during the first ablation (right), allowing for real-time monitoring.

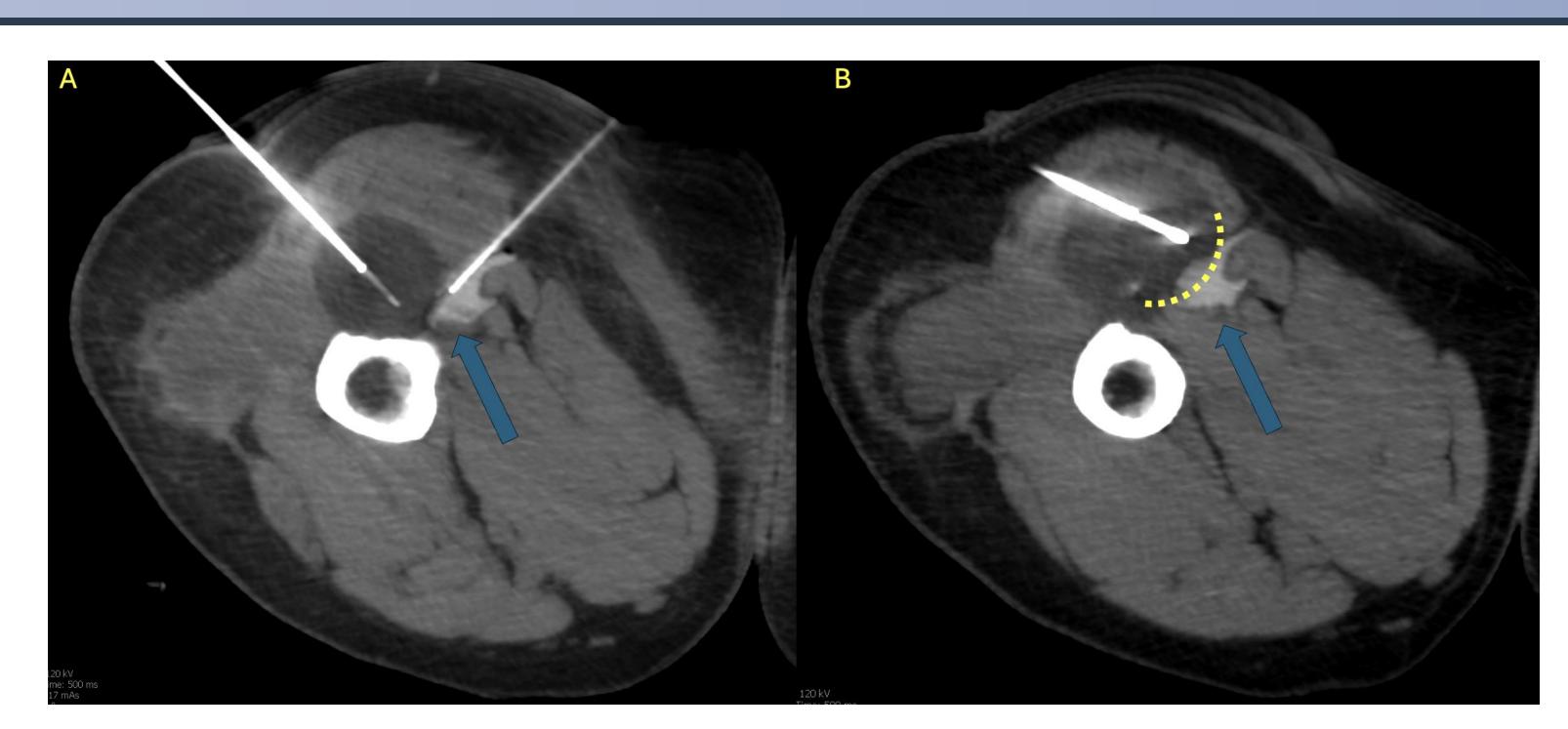


Figure 3. Cryoablation with Varian Ellipsis spherical cryoprobes. (A) Posterolateral probe targeting the superior and deep aspect of the mass. Hydrodissection with a Chiba needle allowed for displacement of the sciatic nerve and profunda femoris branches. (B) Second probe positioned at the inferior/superficial margin. A portion of the hypodense ablation zone overlaps the hyperdense injected contrast (curved dotted yellow line), displacing the critical neurovascular bundle inferomedially (blue arrow).

Conclusion

- The **risk of nerve injury** during cryoablation near neurovascular structures **can be mitigated** through careful planning and protective techniques
- Techniques: **preprocedural** cryoprobe planning, **hydrodissection**, **intraprocedural neuromonitoring**, and freeze-thaw adjustment.
- Applying these evidence-based techniques can significantly reduce nerve injury risk and ultimately improving patient outcomes in interventional oncology.

References

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