

Cryoneurolysis of the Suprascapular Nerve for Perioperative Pain Control After Receiving a Reverse Total Shoulder Arthroplasty (RTSA)

NCT07125833

Status	RECRUITING
Phase	Not Applicable
Sponsor	North Texas Medical Research Institute, PLLC
Enrollment	20 participants

Key Eligibility Criteria

Inclusion (7)

- Male patients or female who are not pregnant and do not plan on future pregnancy during trial participation
 - between 30-85 years of age
 - BMI less than or equal to 45
 - Currently setup for an elective primary reverse total shoulder arthroplasty due to primary osteoarthritis
 - Ability to provide informed consent to participate in the clinical trial
- ... and 2 more (see full listing online)

Exclusion (9)

- poorly controlled comorbidities that would not allow surgical intervention such as poorly controlled diabetes (HbA1C \gt 8.0) renal insufficiency (eGFR \lt 60) poorly controlled CV disease such as CHF that is NYHA class 3 and 4
 - inability to receive the intervention including contraindications:
 - Cryoglobulinemia, paroxysmal cold hemoglobinuria, cold urticaria, Raynaud's disease, and open and/or infected wounds at or near the treatment site
 - patients with history of total joint infection ever or any infection in the last 6 months
 - ASA score \gt 3 and Outpatient Arthroplasty Risk Assessment (OARA; medical risk stratification scoring system to help determine day surgery vs inpatient)1 score \gt 80.
- ... and 4 more (see full listing online)

Locations (1 total)

Orthopaedic Specialists of Dallas, Rockwall, Texas, United States

<https://clinicaltrials.gov/study/NCT07125833>

DISCLAIMER: This document is for informational purposes only and does not constitute medical advice. Always consult your healthcare provider before enrolling in any clinical trial. Information may not be up to date — verify details at ClinicalTrials.gov. Generated by ClinicalTrialsFinder.org.