

Dextrose Hydrodissection for Post-Breast Cancer Chest Wall and Axillary Tightness.

NCT07378319

Status	RECRUITING
Phase	Not Applicable
Sponsor	Taichung Veterans General Hospital
Enrollment	15 participants

Plain Language Summary

This study tests dextrose prolotherapy — injections of concentrated sugar water into connective tissue — as a treatment for persistent chest wall and underarm tightness or pain that continues more than 3 months after breast cancer surgery and has not improved with standard physical therapy.

****You may be eligible if...****

- You have had breast cancer surgery (mastectomy or lumpectomy), with or without lymph node removal
- You still have significant chest or underarm pain, tightness, or movement restriction at least 3 months after surgery
- Your symptoms haven't significantly improved after 12 weeks of standard physical therapy
- You are 18 or older

****You may NOT be eligible if...****

- Your breast cancer has come back or spread
- You currently have active lymphedema requiring treatment
- You have a bleeding disorder or are on blood thinners that would make injections unsafe
- You have an active infection or unhealed wound at the injection site
- You are allergic to dextrose
- You are pregnant

Talk to your doctor to see if this trial is right for you.

Key Eligibility Criteria

Inclusion (6)

- History of breast cancer surgery (mastectomy or breast-conserving surgery), with or without axillary lymph node dissection
 - Persistent ipsilateral chest wall and/or axillary pain, tightness, or movement restriction lasting e 3 months after surgery;
 - Clinically significant baseline symptom severity, defined as Quick Disabilities of the Arm, Shoulder and Hand (QuickDASH) questionnaire e 25 and/or pain intensity e 5/10 on the Numeric Rating Scale (NRS);
 - Completion of 12 weeks of standard physical therapy with no significant improvement, defined as a change in QuickDASH score from baseline to completion of therapy below the minimal clinically important difference (MCID; <15 points) and/or pain improvement < 2 points on the NRS .
 - Age e 18 years;
- ... and 1 more (see full listing online)

Exclusion (7)

- Evidence of local cancer recurrence or metastatic disease;
- Active lymphedema requiring ongoing decongestive therapy at the time of enrollment;
- Shoulder pathology, including intra-articular or peri-articular conditions (e.g., adhesive capsulitis with marked capsular restriction, or acute rotator cuff tear,) judged by clinical assessment to be the dominant cause of symptoms.

<https://clinicaltrials.gov/study/NCT07378319>
• Active infection, skin lesion, or unhealed surgical wound at the intended injection site;

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.

- Known bleeding disorders or anticoagulant use contraindicating injection;
- ... and 2 more (see full listing online)

Locations (1 total)

Taichung Veterans General Hospital, Taichung, Taiwan, Taiwan