Comparison of the Analgesic Effect of 2 Shoulder Infiltrations

NCT05408065

Status	RECRUITING
Phase	Not Applicable
Sponsor	Centre hospitalier de l'Université de Montréal (CHUM)
Enrollment	84 participants

Key Eligibility Criteria

Inclusion (6)

- A clinical examination that confirms the radiological diagnosis of moderate to severe primary shoulder osteoarthritis, stage II and above, according to the Samilson-Prieto classification.
- Patients aged between 20 and 90 years.
- A patient with bilateral shoulder osteoarthritis will choose the side of the infiltration, only one side can be chosen to participate in the study.
- The patient must have a clinical pain threshold of a minimum of 4/10 on the visual analogue scale.
- The patient must have the cognitive ability to read and fill out the questionnaires.

... and 1 more (see full listing online)

Exclusion (12)

- Presence of a transfixing rotator cuff tear assessed on MRI.
- No previous shoulder reconstruction surgery.
- Pregnant woman.
- A patient who has received a cortisone infiltration within 6 months prior to the start of the study.
- A patient who has received a platelet-rich plasma or a hyaluronic acid infiltration within 12 months prior to the start of the study.

... and 7 more (see full listing online)

Locations (1 total)

Centre Hospitalier de l'Université de Montréal (CHUM), Montreal, Quebec, Canada