

Accuracy of the shoulder clinical tests, ultrasonography and magnetic resonance in the diagnosis of the supraspinatus tendon lesions in patients undergoing arthroscopy shoulder surgery

Condition category

Musculoskeletal Diseases

Date applied

02/04/2018

Date assigned

18/04/2018

Last edited

27/04/2020

Prospective/Retrospective

Prospectively registered

Overall trial status

Completed

Recruitment status

No longer recruiting

Publication status

Results

Plain English Summary

Background and study aims

The rotator cuff is the group of muscles and tendons around the shoulder joint. They keep the shoulder stable. These can be injured and tear, or get worn over time (tendinopathy).

It is important to determine the accuracy of clinical tests used to diagnosis rotator cuff injuries, to ensure complete trust in the physical exam. Currently there is an abundant use of image exams of the shoulder, such as ultrasound and MRI, to confirm the clinical diagnosis for patients with shoulder pain complaints.

This study aims to determine the right method and the accuracy of these tests to diagnose tendon problems.

Who can participate?

Adults aged over 18 years with shoulder pain following surgery

What does the study involve?

Participants complete twelve physical tests on their shoulder for examination. Within three months of this, they undergo imaging of their shoulder using ultrasound and MRI. Based on the diagnosis, participants then receive the most suitable surgical procedure or treatment on the joint, when suitable.

What are the possible benefits and risks of participating?

There are no direct benefits or risks for all participants of this study.

Where is the study run from?

1. Hospital São Paulo - Universidade Federal de São Paulo (UNIFESP) (Brazil)
2. Hospital Beneficiente Nipobrasileiro de São Paulo (Brazil)
3. Hospital e Maternidade Christóvão da Gama (Brazil)
4. Universidade de Santo Amaro (Brazil)

When is the study starting and how long is it expected to run for?

October 2016 to April 2019

Who is funding the study?

Universidade Federal de São Paulo – UNIFESP (Brazil)

Who is the main contact?

Mr Fabio Nicolao (Public)

Trial website

[\[\]](#)

Contact information

Type

Public

Primary contact

Mr Fabio Anauate Nicolao

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Contact details

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Additional identifiers

EudraCT number

ClinicalTrials.gov number

Protocol/serial number

20180402

Study information**Scientific title**

Diagnostic accuracy study of the shoulder clinical tests and ultrasonography, magnetic resonance in the supraspinatus tendon lesions in patients undergoing arthroscopy shoulder surgery

Acronym**Study hypothesis**

The empty can test and the ultrasonography may present high accuracy in the diagnosis of the supraspinatus tendon lesions.

Ethics approval

Committee on Ethics and Research of UNIFESP (Comitê de Ética e Pesquisa da UNIFESP), 27/04/2017, ref: 1662/2016

Study design

Observational prospective multicentric accuracy study

Primary study design

Observational

Secondary study design

Accuracy study

Trial setting

Hospitals

Trial type

Diagnostic

Patient information sheet

Not available in web format, please use the contact details below to request a patient information sheet: fabionicolao@terra.com.br

Condition

Rotator cuff lesions

Intervention

In this prospective accuracy study, participants undergo a number of tests, listed below:

1. Jobe's test / empty can test: test performed with the arm into medial rotation, elevation of 90°, where the patient is asked to isometrically resist a downward pressure applied by the tester. The test is positive if pain or weakness when performing the maneuver.
2. Full can test: Test performed with a neutral rotation of the shoulder (thumb pointing up), elevation of 90°, and patient is then asked to isometrically resist a downward pressure applied on the arm by the tester. The test is positive if pain or weakness when performing the maneuver.
3. Champagne toast test: Test performed with the shoulder in 30° of abduction and mild shoulder external rotation. The test is positive if pain when performing the maneuver.
4. Drop arm test: The test is applied in the plane of abduction, with the patient's arm placed passively above 90° by the tester; the support is removed, and the patient attempts to lower the arm actively. The test is positive if the patient is unable to actively lower the arm under control beyond the horizontal, and it drops to his or her side.
5. Patte's test: With the arm supported in 90° of scaption, the patient is asked to laterally rotate maximally against the tester's isometric resistance. The test is positive if pain or weakness when performing the maneuver.
6. Resisted lateral test: The patient stands, elbow at side and flexed to 90°, shoulder in neutral rotation. He or she is then asked to laterally rotate the shoulder maximally against the tester's isometric resistance, which is applied at the wrist. The test is positive if pain or weakness when performing the maneuver.
7. Hawkins' test: The upright patient's arm is passively positioned in 90° of flexion at shoulder and elbow. The tester then forcibly medially rotates the patient's shoulder. The test is positive if pain when performing the maneuver.
8. Yocum's test: The patient places the hand of the affected arm on his or her other shoulder and, keeping the point of the affected shoulder down, raises the elbow of the same limb. The test is positive if pain occurs during the maneuver.
9. Neer's sign: The tester forcibly elevates the patient's arm through scaption, preventing scapular movement by pressing down on the clavicle and acromion with the other hand. Pain constitutes a positive Neer's sign.
10. Painful arc test: The patient actively elevates, then lowers, the shoulder in the plane of the scapula to full elevation. The test is considered positive if pain during elevation, during lowering, or both.
11. Cross body adduction test: The arm is placed at 90° of flexion and then a passive horizontal adduction is performed by the examiner across the chest. The test is positive if pain occurs when performing the maneuver.
12. Speed's test: The shoulder is placed in 90° of flexion, with the elbow in extension and the supinated forearm. The patient flexes his or her shoulder against isotonic resistance applied to the patient's wrist. The test is positive if localized to the bicipital groove.

Participants then undergo imaging of their shoulder using ultrasounds and magnetic resonance.

Ultrasonography: Ultrasound evaluation is performed in the coronal, axial and sagittal planes, with a 10 MHz linear transducer, characterized by the presence of tendinopathy, and partial or total rupture of the supraspinatus tendon.

The clinical tests are all done by a orthopedic specialist in shoulder and elbow surgery and the ultrasounds by a radiologist. The tests are performed individually at the hospital. Clinical tests must be performed within the 3 months before diagnostic imaging exams, either Ultrasonography or MRI. Then the participants are followed up according to their pathology to be undergone the most adequate treatment. Participants with indication of arthroscopic surgical treatment are included in this protocol.

The two different types of diagnosing rotator cuff lesions are compared. Correlation of the location and size of the supraspinatus lesion with the clinical tests measured using arthroscopy findings.

Intervention type

Mixed

Phase

Drug names

Primary outcome measure

Supraspinatus tendon tears are measured using arthroscopy findings and comparing to the magnetic resonance image and ultrasonography. Tears are classified as tendinopathy, partial tears or full thickness tears.

Secondary outcome measures

1. Level of pain is measured using the visual analogue scale after performing the empty can test (Jobe test)
2. Correlation of the location and size of the supraspinatus lesion with the clinical tests is measured using arthroscopy findings

Overall trial start date

10/10/2016

Overall trial end date

30/04/2019

Reason abandoned (if study stopped)

Eligibility

Participant inclusion criteria

1. Patients of both genders
2. Over 18 years old
2. Complaint of shoulder pain with at least 1 month of duration undergoing arthroscopy surgery

Participant type

Patient

Age group

Adult

Gender

Both

Target number of participants

100

Participant exclusion criteria

1. Cognitive alterations of the patient that prevents comprehension of the proposed treatment
2. Lack of outpatient return; image examination with more than three months of clinical evaluation of the patient
3. Traumatic or unexpected event in the affected shoulder between arthroscopy surgery, magnetic resonance image, ultrasonography and physical examination

Recruitment start date

01/05/2018

Recruitment end date

01/04/2019

Locations

Countries of recruitment

Brazil

Trial participating centre

Hospital São Paulo - Universidade Federal de São Paulo (UNIFESP)
Sao Paulo
04037020
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Trial participating centre

Hospital Beneficiente Nipobrasileiro de São Paulo
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Trial participating centre

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Sponsor information

Organisation

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Sponsor type

University/education

Website

□

GRID

[grid.412283.e \[https://www.grid.ac/institutes/grid.412283.e\]](https://www.grid.ac/institutes/grid.412283.e)

Funders

Funder type

University/education

Funder name

Universidade Federal de São Paulo - UNIFESP

Alternative name(s)

Funding Body Type

Funding Body Subtype

Location

Results and Publications

Publication and dissemination plan

Planned publication in a high-impact orthopaedic journal.

IPD sharing statement:

The datasets generated during and/or analysed during the current study will be stored in a publically accessible portal according to the last ICMJE data sharing statement.

Intention to publish date

30/05/2019

Participant level data

Stored in repository

Basic results (scientific)

See additional file ISRCTN13083925_BasicResults_27Apr2020 (added 27/04/2020)

Publication list

Publication citations

Additional files

- [ISRCTN13083925_BasicResults_27Apr2020.pdf](#) [/editorial/retrieveFile/a705f94d-e8a6-48ae-940f-3004a7fddoc3/35069] Uploaded 27/04/2020

Editorial Notes

27/04/2020: The basic results of this trial have been uploaded as an additional file.