

# The effect of an eccentric exercise program on patients with shoulder pain and disability, which is caused by dysfunction of the rotator cuff

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<b>Registration date</b> 11/03/2014	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 11/03/2014	<b>Condition category</b> Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Shoulder pain is a very common complaint. Shoulder pain and stiffness has major effects on the use of healthcare resources and work-related costs. Exercise has been shown to help with many conditions, including tendinopathy. Tendinopathy is a term used to describe symptoms arising from a tendon (a tissue that connects muscle to bone). This study aims to find out the effect of a particular type of exercise on patients suffering from rotator cuff tendinopathy. This exercise is an eccentric exercise. Eccentric exercise is the lowering phase of an exercise, where your muscles are paying out, rather than contracting.

### Who can participate?

Adult patients suffering from tendon-related shoulder pain can participate in this study.

### What does the study involve?

Following assessment eligible patients who agree to participate will be randomly allocated to one of two groups. One of the groups will be treated using an eccentric exercise program, which will be demonstrated to them and conducted by the physiotherapist. They will also receive general advice. The other group will receive traditional physiotherapy treatment, including other forms of exercise and manual treatment techniques. All participants will be asked to complete various questionnaires at the beginning of the study, and at 6 and 12 weeks into their treatment.

### What are the possible benefits and risks of participating?

Patients will not benefit directly from taking part in this study but the information we get may provide further knowledge about managing this condition. Participation in this study should be as safe as normal physiotherapy treatment.

### Where is the study run from?

The study will be run from the Physiotherapy Department of St Vincents University Hospital, Dublin, Ireland.

When is study starting and how long is it expected to run for?  
The study is expected to run from December 2013 until July 2014.

Who is funding the study?  
Investigator initiated and funded (Ireland).

Who is the main contact?  
Mr Micheal Bailey  
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## Contact information

**Type(s)**  
Scientific

**Contact name**  
Mr Micheal Bailey

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## Additional identifiers

### Study information

**Scientific Title**  
The effect of an eccentric training program on patients clinically diagnosed with chronic unilateral rotator cuff tendinopathy

**Study objectives**  
Tendinopathy is a generic term used to describe pathology in and pain arising from a tendon; it is associated with failed healing response, in this study we will be examining tendinopathy in relation to the rotator cuff tendons of the shoulder.

The aim of this study is to evaluate the effectiveness of a structured eccentric exercise program in the management of patients suffering from unilateral shoulder pain where rotator cuff tendinopathy is seen as the primary cause when compared with regular treatment.

**Ethics approval required**  
Old ethics approval format

**Ethics approval(s)**  
Ethical Committee of St Vincent's University Hospital, 09/09/2013

## **Study design**

Single blind randomized controlled trial

## **Primary study design**

Interventional

## **Study type(s)**

Treatment

## **Health condition(s) or problem(s) studied**

Rotator Cuff Tendinopathy

## **Interventions**

Conservative management of rotator cuff tendinopathy using eccentric exercise. Patients will complete 15 repetitions in three sets of each exercise twice a day. There are three exercises to complete. Patients will be provided with written and pictorial references as to the correct technique of these exercises and guided by the physiotherapist.

The control group will be treated with traditional physiotherapy intervention, excluding the above eccentric exercise program.

The duration of each participants involvement in the study will be 12 weeks. Measures will be taken at baseline, 6 weeks, and 12 weeks.

## **Intervention Type**

Other

## **Phase**

Not Applicable

## **Primary outcome(s)**

1. Pain and Disability - SPADI
2. Health Related Quality of Life - EuroQol EQ 5D

Outcomes measured at baseline, 6 weeks and 12 weeks

## **Key secondary outcome(s)**

1. Shoulder Range of Motion - Goniometry assessed at baseline and end point
2. Patients perceived improvement - Global Rating of Change Scale assessed at 6 weeks and 12 weeks
3. Pain - VAS assessed at baseline and end point

## **Completion date**

31/07/2014

## **Eligibility**

### **Key inclusion criteria**

1. Consent to participate
2. Male and female aged from 18 65

3. Sufficient range of motion (ROM) and function of non-affected shoulder ability to perform exercises
4. Shoulder pain which distributes pain over C4/5 dermatome
5. Passive range of shoulder movement which is greater than active range of movement
6. Pain which is reduced by rest
7. Pain on resisted external rotation or pain on resisted abduction
8. Symptom duration equal to or greater than 12 weeks

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

18 years

**Sex**

All

**Key exclusion criteria**

1. Recent shoulder/upper arm surgery
2. Recent shoulder or upper arm fracture
3. Recent shoulder dislocation
4. Symptoms aggravated by C-Spine range of movement
5. Severe resting pain
6. Signs of systemic/rheumatologic cause of symptoms
7. Obvious signs of cognitive impairment
8. Inability to understand written or spoken English
9. Bilateral symptoms which will impair participation in eccentric program
10. Positive Drop Arm

**Date of first enrolment**

16/12/2013

**Date of final enrolment**

31/07/2014

**Locations****Countries of recruitment**

Ireland

**Study participating centre**

Physiotherapy Department

Dublin

Ireland

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## Sponsor information

### Organisation

University College Dublin (Ireland)

### ROR

<https://ror.org/05m7pjf47>

## Funder(s)

### Funder type

Other

### Funder Name

Investigator initiated and funded (Ireland)

## Results and Publications

### Individual participant data (IPD) sharing plan

### IPD sharing plan summary

Not provided at time of registration