

REPORT FOR RESEARCH OR TEACHING INVOLVING HUMANS

Human Research Ethics Committee

HUMAN ETHICS APPROVAL NUMBER

H6129

Please submit in hard copy a signed original to: Research Office, Room 128, Faculty Science & Engineering Building (DB17), James Cook University, Townsville, Qld, 4811. Please also email a pdf of this report to ethics@jcu.edu.au.

1	TITLE OF PROJECT	Specific phys	siothe	erapy ma	nageme	nt for su	ubacromial imp	pingement
2	CATEGORY	4				4		
3	PERIOD DURING WH	ICH ACTIVITIES	SREG	UIRING	ETHICS	APPRO	VAL OCCURR	ED
	COMMENCEMENT DATE	26.08.2015			FINISH D	ATE	9.12.2016	
4	STATUS OF PROJECT (Please tick)	Completed	√	In Pro	gress	Ab	andoned	Not Commenced

5	PRINCIPAL INVESTIGATOR'S DETAILS							
	Last Name, First name and Title	ESN ¹	Orgu	Discipline/Sch	ool or Institution (Country)			
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5a	SUPERVISOR DETAILS 1 (if applicable)						
	Last Name, First name and Title	ESN ¹	Orgu	Discipline/Sch	ool or Institution (Country)		
	Gordon, Susan Professor			School of He University	alth Sciences, Flinders		
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5b	SUPERVISOR DETAILS 2 (if applicable)	ERVISOR DETAILS 2 (if applicable)			
	Last Name, First name and Title	ESN ¹	Orgu	Discipline/Sch	ool or Institution (Country)
	Watt, Kerrianne Associate Professor	E	2110	College of Pu Sciences	blic Health, Medical & Vet
	Email	Phone			Fax
- b	kerrianne.watt@jcu.edu.au	4781 60	97		

¹ Indicate if the Researcher is currently an Employee or a **S**tudent of JCU, or a researcher who is **N**ot affiliated with JCU. If the project involves international cooperation, please specify the country.

PLEASE A	NSWER ALL	QUESTIONS	S:						
6 COMPLIA	ANCE								
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If NO, please	provide deta	ils below:				Yes		No	
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NUMBER C	F INDIGENC	US PARTIC	IPANTS INVO	DLVED IN T	HE STUDY	·		-	
Male		Female		Children		То	tal		
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7 UNEXPE	CTED ADVE	RSE EFFEC	TS or EVENT	'S	\\				
unexpected	events that of	ccurred of wh	suffer any un sich the Huma s? Could thes	n Research E	Ethics Com	mittee s	should I	be awar	
If YES, pleas	se provide det	ails below:				Yes		No	V
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8 COMPLA	AINTS								
Were there a with the stu		s from partic	ipants, or any	other organi	sation/com	munity	group	etc invo	lved
If YES, pleas	se provide det	ails below:				Yes		No	
					10.100.27		-		

11 **CERTIFICATION PRINCIPAL INVESTIGATOR**

I declare that the statements made in this report ar	re correct	
Adelen Land	HELEN LAND	17,01,2017
Signature (Principal Investigator)	Name	Date
Please note: if the principal investigator is an Honours or H	igher Degree Student, the supervisor must als	o sian this report

AUTHORISATION by SUPERVISOR(S) (Supervisor(s) must sign this declaration) 12

I/We certify that the statements made in this report are correct.. Kundff Kerrianne Watt 17/1/17 Signature (Supervisor) Name Date

13 AUTHORISATION by DEAN OF COLLEGE/DELEGATE:

certify that the statements made in this report are o	correct	
Digitally signed by Lee Stewart DN: cn=Lee Stewart, c=JCU, ou=CHS, email=leestewart@cueduau, c=AU Date: 2017 012 01 114349 + 10'00'	Prof Lee Stewart, Dean CHS	20/01
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9 STATEMENT OF REPORT

Please provide a brief statement of the outcomes and conduct of project.

Recruitment commenced on the 18th August 2015. Trial registration accepted with the Australian New Zealand Clinical Trials Registry (ANZCTR): 12615001303538 on the 30th November 2015. The final participant recruited on the 7th September 2016 with the twelve week trial successfully completed on the 30th November 2016. One hundred and fifty two volunteers were assessed by the primary investigator for eligibility, with 79 failing to meet the eligibility criteria (outside age limits, cervical and/or thoracic signs present, frozen shoulder, constant pain limiting active glenohumeral range above 90 degrees) and four electing to not participate in the trial (excessive distance needing to travel for treatments or inability to attend all required treatment sessions during the allocated period) Sixty nine volunteers consented to participate in the trial and were randomly allocated to a treatment B or treatment C on attendance at their first scheduled treatment session. Twenty three participants were allocated to treatment group A, 22 to treatment group B and 24 to treatment group C. Recruitment continued until 20 participants had completed the 12 week trial period in each of the three groups. Three participants randomised to treatment group A, two randomised to treatment group B and two randomised to treatment group C did not complete the 12 week trial period ceasing to attend scheduled treatment sessions. Two participants randomised to treatment group C were withdrawn from the trial with one identified as early stage adhesive capsulitis and the other with tendon calcification Data input into SPSS completed by 10th January 2017. Currently reviewing statistical analysis of results with Dr Watt. Preparing draft RCT paper following the CONSORT statement guidelines.

10 PUBLICATIONS

Please provide a reference list of all publications generated from the project. (Please indicate status of the publications, i.e. submitted, accepted etc)

"Specific physiotherapy management for subacromial impingement" to submit for publication with Musculoskeletal Science and Practice in May 2017.