Pathophysiology of Dupuytren's Contracture

Submission date 11/08/2010	Recruitment status No longer recruiting	Prospectively registeredProtocol
Registration date 11/08/2010	Overall study status Completed	 Statistical analysis plan Results
Last Edited 18/04/2017	Condition category Musculoskeletal Diseases	 Individual participant data Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s) Scientific

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers 6585

Study information

Scientific Title

Pathophysiology of Dupuytren's Contracture

Study objectives

Dupuytren's disease is a common inheritable disorder, mainly affecting the hand. The digits develop progressive flexion contractures and hand function is significantly impaired. The current mainstay of treatment is surgical excision of the affected tissues but recurrence following excision is seen in approximately 40% of patients. Replacing the palmar skin with grafts obtained from a non-palmar site on the body virtually abolishes recurrence. Based on this well-established surgical observation, we have developed a novel in vitro model that replicates these interactions between skin cells and the contractile cells responsible for Dupuytren's disease. We have also identified a molecule (tenascin-C) that may control the signalling between these cell types.

Ethics approval required

Old ethics approval format

Ethics approval(s)

MREC approved, ref: 06/Q0403/95

Study design Multicentre non-randomised interventional treatment trial

Primary study design Interventional

Secondary study design Non randomised study

Study setting(s) GP practice

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied

Topic: Generic Health Relevance and Cross Cutting Themes; Subtopic: Generic Health Relevance (all Subtopics); Disease: Surgery

Interventions Elucidate the exact role of tenascin-C in Dupuytren's disease

Intervention Type Other

Phase Not Specified

Primary outcome measure

Develop candidate therapeutic target to provide non-surgical intervention to modulate the disease

Secondary outcome measures Not provided at time of registration

Overall study start date 02/11/2006

Completion date 31/07/2011

Eligibility

Key inclusion criteria Not provided at time of registration

Participant type(s) Patient

Age group Not Specified

Sex Not Specified

Target number of participants Planned sample size: 90

Key exclusion criteria Not provided at time of registration

Date of first enrolment 02/11/2006

Date of final enrolment 31/07/2011

Locations

Countries of recruitment England

United Kingdom

Study participating centre

Imperial College London London United Kingdom W6 8LH

Sponsor information

Organisation Royal College of Surgeons of England (UK)

Sponsor details 35-43 Lincoln's Inn Fields London England United Kingdom WC2A 3PE

Sponsor type University/education

Website http://www.rcseng.ac.uk/

ROR https://ror.org/02qrg5a24

Funder(s)

Funder type Charity

Funder Name Healing Foundation

Alternative Name(s)

Funding Body Type Private sector organisation

Funding Body Subtype Other non-profit organizations

Location United Kingdom

Results and Publications

Publication and dissemination plan Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary Not provided at time of registration