

A double-blind randomised controlled trial of the efficacy of low level laser therapy on wound healing following nail surgery

Submission date 23/01/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 23/01/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 16/06/2014	Condition category Skin and Connective Tissue 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

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number

RDC01733

Study information

Scientific Title

Study objectives

1. To compare the efficacy of low level laser therapy with conventional wound dressing on wound healing following nail surgery
2. To show whether low level laser therapy reduces pain levels compared to conventional wound dressing
3. To evaluate the cost effectiveness of low level laser therapy in terms of drug and dressing uptake and return appointments

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Skin and connective tissue diseases

Interventions

1. Low level laser therapy
2. Conventional wound dressing

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

1. Assessments of wound size
2. Level of inflammation
3. Level of exudate

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/04/2003

Eligibility**Key inclusion criteria**

Patients referred to the chiropody clinic, who are going to have one or more toenails removed and who have given informed consent will be randomised into one of two treatment groups

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/10/2001

Date of final enrolment

01/04/2003

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Diabetic Centre

London

United Kingdom

E11 1NR

Sponsor information**Organisation**

NHS R&D Regional Programme Register - Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
NHS Executive London (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes