

A randomised controlled trial of larval therapy versus standard care in the management of necrotic wounds with and without methicillin-resistant *Staphylococcus aureus* (MRSA)

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 22/09/2014	Condition category Injury, Occupational Diseases, Poisoning	<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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0034136469

Study information

Scientific Title

Study objectives

To compare the clinical and cost effectiveness of larval therapy versus standard care.

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

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Necrotic wounds with or without methicillin-resistant *Staphylococcus aureus*

Interventions

Randomised controlled trial of larval therapy versus standard care

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Efficacy of treatment

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/10/2002

Completion date

30/09/2005

Eligibility

Key inclusion criteria

All patients over 18 years with necrotic wounds, in hospital or the community

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/10/2002

Date of final enrolment

30/09/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Barnsley District General Hospital NHS Trust

Barnsley

United Kingdom

S75 2EP

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Government

Funder Name

Barnsley District General Hospital NHS Trust (UK)

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