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A randomised double-blinded cross-over trial comparing betamethasone valerate and tacrolimus ointment for the treatment of moderate to severe atopic eczema

Submission date
30/09/2004

Recruitment status
No longer recruiting

Registration date
30/09/2004

Overall study status
Completed

Last Edited
17/04/2015

Condition category
Skin and Connective Tissue Diseases

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[_] Results
[_] 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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Additional identifiers

Protocol serial number
N0263128762

Study information

Scientific Title

A randomised double-blinded cross-over trial comparing betamethasone valerate and tacrolimus ointment for the treatment of moderate to severe atopic eczema

Study objectives
Is tacrolimus more efficacious than betamethasone valerate ointment in moderate to severe atopic dermatitis?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised double-blinded cross-over trial

Primary study design
Interventional

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Skin and Connective Tissue Diseases: Atopic dermatitis

Interventions
1. Steroid ointment
2. Other ointment

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Betamethasone valerate and tacrolimus ointment

Primary outcome(s)
SCORAD score (numerical assessment score of patient symptoms, range 0-103). Ointment weight. Recording of side-effects noted.

Key secondary outcome(s))
Not provided at time of registration

Completion date
01/10/2005

Eligibility

Key inclusion criteria
100 patients from Dermatology

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
31/01/2003

Date of final enrolment
01/10/2005

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
The Middlesex Hospital
London
United Kingdom
W1N 8AA

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Government

Funder Name
University College London Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration