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A prospective within-patient randomised controlled trial of the treatment of vitiligo with the 308-nm excimer laser: a pilot study

Submission date
12/09/2003

Recruitment status
No longer recruiting

Registration date
12/09/2003

Overall study status
Completed

Last Edited
14/05/2018

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
N0064122126

Study information

Scientific Title

A prospective within-patient randomised controlled trial of the treatment of vitiligo with the 308-nm excimer laser: a pilot study

Study objectives
The main purpose of this study is to determine the efficacy of the 308-nm excimer laser in the treatment of facial vitiligo which is a very cosmetically disabling disorder. This will be a prospective within patient randomised controlled trial and will investigate and quantify the repigmentation achieved with this treatment. We shall also be monitoring the side effects of the treatment if any. The other aspect of the study is to determine if the repigmentation achieved is sustained over a duration of time.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Single centre, patient volunteers, prospective, controlled, therapeutic, observational, single-blind, devices, longitudinal, invasive procedures, randomised study.

Primary study design
Interventional

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Skin and Connective Tissue Diseases: Facial vitiligo

Interventions
Not provided at time of registration

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
Not provided at time of registration

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

Completion date
01/06/2005

Eligibility

Key inclusion criteria
17 to 48 patient volunteers will be recruited.

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Child

Sex
Not Specified

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
22/04/2003

Date of final enrolment
01/06/2005

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
City Hospital
Birmingham
United Kingdom
B18 7QH

Sponsor information

Organisation
Department of Health (UK)

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Sandwell and West Birmingham Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration