ISRCTN60370678 https://doi.org/10.1186/ISRCTN60370678

Comparison of the effectiveness of routinely used treatments for external genital warts

Submission date	Recruitment status
30/09/2004	No longer recruiting
Registration date	**Overall study status**
30/09/2004	Completed
Last Edited	**Condition category**
08/01/2009	Skin and Connective Tissue Diseases

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Jackie S Sherrard

Contact details
GUM
The Harrison Department
Radcliffe Infirmary
Woodstock Road
Oxford
United Kingdom
OX2 6HE
+44 (0)1865 224343

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0176052132

Study information

Scientific Title

Study objectives
This study aims to compare the routinely available and used treatments in a methodological way, assessing time to clearance of macroscopic warts.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Prospective randomised controlled study

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Skin and Connective Tissue Diseases: Genital warts

Interventions
Interventions will be five different treatments:
1. Podophyllin 25% in tincture of benzoin
2. Trichloacetic acid
3. Cryotherapy
4. Trichloacetic acid and Podophyllin 25%
5. Podophyllin 25% and Cryotherapy

Intervention Type
Other

Phase
Not Specified

Primary outcome measure

1. Number of treatments required to achieve clearance of external genital warts
2. If no clearance, percentage reduction in surface area of warts after treatment eight

Secondary outcome measures
Not provided at time of registration

Overall study start date
02/11/1999

Completion date
01/03/2004

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
2270 patients

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
02/11/1999

Date of final enrolment
01/03/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
GUM
Oxford

United Kingdom
OX2 6HE

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
Oxford Radcliffe Hospitals 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2007		Yes	No