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

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Additional identifiers

Protocol serial number
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

Study type(s)
Not Specified

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(s)
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

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

Completion date
01/03/2004

Eligibility

Key inclusion criteria
Not provided at time of registration

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
02/11/1999

Date of final enrolment
01/03/2004

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
GUM
Oxford
United Kingdom
OX2 6HE

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Government

Funder Name
Oxford Radcliffe Hospitals NHS Trust (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?
Results article	results	01/06/2007		Yes	No