

The effectiveness of peri-operative use of human surgical sealant (Quixil) in reducing the post-operative seroma formation in breast cancer patients

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
30/09/2004	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
30/09/2004	Completed	<input type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
05/12/2014	Cancer	<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Mr T Vijayganesh

Contact details

Good Hope Hospital
Sutton Coldfield
Birmingham
United Kingdom
B75 7RR

-
abc@email.com

Additional identifiers

Protocol serial number

N0523136364

Study information

Scientific Title

The effectiveness of peri-operative use of human surgical sealant (Quixil) in reducing the post-operative seroma formation in breast cancer patients

Study objectives

The trial proposes to establish the usefulness of 'Quixil' in reducing the incidence of post-operative seroma in breast cancer operations

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Cancer: Breast

Interventions

1. Quixil
2. No Quixil

Intervention Type

Procedure/Surgery

Primary outcome(s)

Reduction in seroma formation of more than 30%

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/12/2004

Eligibility

Key inclusion criteria

100 patients undergoing mastectomy or wide local excision with axillary clearance for biopsy proven breast cancer. 50 to each arm.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Female

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/08/2003

Date of final enrolment

01/12/2004

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Good Hope Hospital

Birmingham

United Kingdom

B75 7RR

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Good Hope Hospital NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration