

Use of vessel sealing system (ligasure) thyroid surgery: time saving? A prospective randomised study

Submission date 11/04/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 11/04/2007	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 07/06/2012	Condition category Surgery	<input type="checkbox"/> 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Study objectives

Use of a vessel sealing system (ligasure) in thyroid surgery reduces operation time.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval received from the local medical ethics committee

Study design

Randomised, active controlled, parallel group trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Thyroidectomy

Interventions

Use of vessel sealing system (ligasure) versus conventional knotting for ligation of blood vessels during thyroid surgery.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Length of operation.

Secondary outcome measures

1. Bleeding complications
2. Nerve damage

Overall study start date

01/09/2005

Completion date

01/04/2007

Eligibility**Key inclusion criteria**

1. Indication for total hemithyroidectomy
2. Age greater than 18 years
3. Informed consent

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Not Specified

Target number of participants

20

Key exclusion criteria

1. American Society of Anaesthesiologists (ASA) IV and V patients

Date of first enrolment

01/09/2005

Date of final enrolment

01/04/2007

Locations**Countries of recruitment**

Netherlands

Study participating centre

Diakonessenhuis Utrecht

Utrecht

Netherlands

3508 TG

Sponsor information

Organisation

Diakonessenhuis Hospital Utrecht (The Netherlands)

Sponsor details

Department of Surgery

P.O. Box 80250

Utrecht

Netherlands

3508 TG

Sponsor type

Hospital/treatment centre

Website

<http://www.diakonessenhuis.nl/>

ROR

<https://ror.org/01nrpzj54>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Diakonessenhuis Utrecht (The Netherlands)

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	18/01/2012		Yes	No