A Pilot study to evaluate the Clinical Application of Ultrasound Elastography

Submission date	Recruitment status	Prospectively registered
18/06/2010	No longer recruiting	<pre>Protocol</pre>
Registration date	Overall study status	Statistical analysis plan
18/06/2010	Completed	Results
Last Edited	Condition category	Individual participant data
29/08/2013	Surgery	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

5260

Study information

Scientific Title

Acronym

Ultrasound Elastography

Study objectives

Ultrasound elastography is a non-invasive ultrasound method which evaluates the deformability of a structure. The principle of elastography is the utilisation of tissue compression producing varying levels of strain within different tissue types. 'Strain images' obtained can then be analysed to assess the mechanical properties of tissues and infer their size and nature.

Ethics approval required

Old ethics approval format

Ethics approval(s)

MREC approved (ref: 07/H0306/90)

Study design

Single centre non-randomised interventional diagnosis, prevention and screening trial

Primary study design

Interventional

Secondary study design

Non randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Prevention

Participant information sheet

Health condition(s) or problem(s) studied

Topic: Generic Health Relevance and Cross Cutting Themes; Subtopic: Generic Health Relevance (all Subtopics); Disease: Other

Interventions

Ultrasound elastography

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

Assess the ability to obtain strain images from different tissues of the body, e.g. abdomen, pelvis, head and neck, breast, testes and soft tissues

Secondary outcome measures

Evaluate accuracy of ultrasound elastography compared with standard ultrasound

Overall study start date

01/08/2007

Completion date

01/12/2010

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

Planned sample size: 600

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/08/2007

Date of final enrolment

01/12/2010

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Addenbrooke's Hospital

Cambridge United Kingdom CB2 0QQ

Sponsor information

Organisation

Cambridge University Hospitals NHS Foundation Trust (UK)

Sponsor details

Addenbrookes Hospital Hills Road Cambridge England United Kingdom CB2 0QQ

Sponsor type

Hospital/treatment centre

Website

http://www.cuh.org.uk/addenbrookes/addenbrookes index.html

ROR

https://ror.org/04v54gj93

Funder(s)

Funder type

Research council

Funder Name

Engineering and Physical Sciences Research Council (EPSRC) (UK)

Alternative Name(s)

UKRI Engineering and Physical Sciences Research Council, Engineering and Physical Sciences Research Council - UKRI, Engineering & Physical Sciences Research Council, EPSRC

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Funder Name

The Wellcome Trust (UK)

Results and Publications

Publication and dissemination planNot 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