# Wall thickness measurement by ultrasonography as a predictor for arteriovenous fistula function in haemodialysis patients

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
27/02/2007	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
27/02/2007	Completed	Results
Last Edited	Condition category	Individual participant data
27/02/2007	Urological and Genital Diseases	<ul><li>Record updated in last year</li></ul>

#### Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

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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

Research to investigate whether ultrasonographically and histologically measured radial artery wall thickness are comparable, and whether this wall thickness is predictive for maturation of arteriovenous fistulas in haemodialysis patients.

#### Hypothesis:

Radial artery wall thickness is predictive for arteriovenous fistula maturation and can be measured by means of ultrasonography.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Ethics approval received from the local medical ethics committee

#### Study design

Non-randomised, non-controlled, parallel group, clinical trial

#### Primary study design

Interventional

# Secondary study design

Single-centre

# Study setting(s)

Hospital

# Study type(s)

Diagnostic

#### Participant information sheet

## Health condition(s) or problem(s) studied

Chronic renal insufficiency

#### **Interventions**

Excision biopsy of a small piece of the radial artery during operation.

# Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome measure

- 1. Similarity between ulrasonographically measured and histologically measured radial artery wall thickness
- 2. Patency rates

#### Secondary outcome measures

No secondary outcome measures

#### Overall study start date

01/01/2005

#### Completion date

31/12/2005

# Eligibility

#### Key inclusion criteria

- 1. Chronic renal failure and requirement of an arteriovenous fistula
- 2. Age 18 years or older
- 3. Informed consent

#### Participant type(s)

**Patient** 

#### Age group

Adult

#### Lower age limit

18 Years

#### Sex

**Not Specified** 

## Target number of participants

30

#### Key exclusion criteria

- 1. Age less than 18 years
- 2. Placement of vascular access prosthesis
- 3. Unable to sign written informed consent

#### Date of first enrolment

01/01/2005

#### Date of final enrolment

31/12/2005

# **Locations**

#### Countries of recruitment

Netherlands

Study participating centre
University Medical Centre Utrecht (UMCU)
Utrecht
Netherlands
3508 GA

# Sponsor information

## Organisation

University Medical Centre Utrecht (UMCU) (The Netherlands)

#### Sponsor details

P.O. Box 85500 Utrecht Netherlands 3508 GA

#### Sponsor type

Hospital/treatment centre

#### Website

http://www.umcutrecht.nl/zorg/

#### **ROR**

https://ror.org/04pp8hn57

# Funder(s)

#### Funder type

Charity

#### **Funder Name**

Dutch Kidney Foundation (Nierstichting Nederland) (The Netherlands)

## Alternative Name(s)

**Dutch Kidney Foundation** 

# **Funding Body Type**

Private sector organisation

# Funding Body Subtype

Trusts, charities, foundations (both public and private)

#### Location

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