

Effects of Rosiglitazone on fluid retention and insulin resistance in response to Head-out immersion in individuals with type 2 diabetes and impaired glucose tolerance.

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 15/05/2012	Condition category Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

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

N0295122840

Study information

Scientific Title

Study objectives

Type 2 diabetes is linked to molecular mechanisms of insulin resistance. Rosiglitazone restores impaired sodium excretion in type 2 diabetes by causing increased insulin sensitivity in the kidneys and within the peripheral vasculature.

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

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

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

Nutritional, Metabolic, Endocrine: Diabetes

Interventions

Prospective Randomised Controlled Trial.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Insulin sensitivity before and after treatment and sodium excretion.

Secondary outcome measures

Not provided at time of registration

Overall study start date

17/07/2002

Completion date

10/12/2004

Eligibility

Key inclusion criteria

Patients with Type 2 Diabetes and impaired glucose tolerance (48).

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

17/07/2002

Date of final enrolment

10/12/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Diabetes Centre
Rugby
United Kingdom
CV22 5PX

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
University Hospitals Coventry and Warwickshire NHS Trust (UK)

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	01/04/2008		Yes	No