

DRN 725: Chronic Disease Resource Centre Diabetes Division

Submission date 19/11/2012	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 27/03/2013	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 27/03/2013	Condition category Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

The study aims to recruit individuals with diabetes or a pre-diabetic condition. We will collect information and blood samples from patients once a year at a time convenient to the individual and the research team (normally at the same time as their annual diabetes check). The information collected from this study will be used in research into how diabetes affects individuals in the setting of it being a chronic disease. The aim of this study is to establish a database of information of adult participants diagnosed with impaired glucose tolerance. This database will be used to explore the psycho-social aspects of diabetes in the context of it being a chronic disease. Also, the database will be used to develop partnerships with other researchers to advance diabetes research for the benefit of individuals with diabetes.

Who can participate?

Individuals who are 18 years or older with confirmed diabetes or abnormal glucose tolerance. Individuals must be willing and able to attend annual appointments at clinical study recruitment site centre.

What does the study involve?

The study involves an annual study visit where non-invasive clinical measurements will be taken, along with a questionnaire, blood and urine samples.

What are the possible benefits and risks of participating?

As this study does not involve any intervention then the risks to participants are minimal.

Where is the study run from?

The study is currently run from the Queen Elizabeth Hospital Birmingham, UK. There is a network of other centres throughout the West Midlands who are also participating in this study.

When is the study starting and how long is it expected to run for?

The study started in September 2012 and is expected to run until April 2022.

Who is funding the study?

Novo Nordisk A/S (USA)

Who is the main contact?
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Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
12984

Study information

Scientific Title
Chronic Disease Resource Centre: What is the natural history of diabetes and its complications?

Study objectives
The aim of this study is to establish a prospective, observational multi-ethnic cohort of adult participants diagnosed with impaired glucose tolerance. The cohort will encourage academic collaboration for patient benefit and will be characterised by anthropometry, questionnaires, biological samples and co-morbidities forming a resource for epidemiological, clinical and health service research.

The study will aim to recruit all eligible participants from across the West Midlands Birmingham area. A 10 year study will be undertaken to recruit and follow-up 1000 participants, concentrating initially on University Hospitals Birmingham NHS Foundation Trust (UHBFT), before but could be opening to other NHS clinical centres trusts interested in collaborating.

In addition to providing information about the natural history and progression of diabetes, development of complications and co-morbidities it will provide unique information for use in the detection of biomarkers to help predict the progression of complications and to monitor the effectiveness of interventions.

Also, the study will try to accurately categorise patients with potentially different types of diabetes in order to determine whether any factors significantly affect their long-term morbidity or mortality.

In summary the overarching aims and objectives of this study is:

1. To understand the natural history of diabetes in our local population. To determine the incidence of other chronic medical conditions in patients with diabetes.
2. Provide a repository of information for future analysis to help determine potential biomarkers of chronic diseases related to diabetes.

More details can be found at: <http://public.ukcrn.org.uk/Search/StudyDetail.aspx?StudyID=12984>

Ethics approval required

Old ethics approval format

Ethics approval(s)

NRES Committee West Midlands - Staffordshire, 8th May 2012, ref: 12/WM/0089

Study design

Non-randomised observational cohort study

Primary study design

Observational

Secondary study design

Cohort study

Study setting(s)

Hospital

Study type(s)

Quality of life

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

Diabetes Research Network

Interventions

The study is observing changes in participants with diabetes in several domains: depression, quality of life, eating behaviour, risk of sleep apnoea, activities of daily living, employment and productivity, as well as developing a repository of biological samples. The tests will include routine clinical assessment, bone, liver, renal, thyroid blood tests, and questionnaires. Patients will also be invited to undertake more significant testing including analysis of saliva samples, AGE reader, pancreatic stimulation tests.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Micro and macro vascular complications measured at 10 years

Secondary outcome measures

Qualitative and questionnaire based assessment of diabetes related endpoints measured at 10 years

Overall study start date

06/09/2012

Completion date

01/04/2022

Eligibility

Key inclusion criteria

1. Male or female, age ≥ 18 years
2. Confirmed diagnosis of diabetes or prediabetes
3. Individuals freely able to attend appointments at the Chronic Disease Research Centre
4. Individuals who are able to undergo the appropriate investigations

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

UK Sample Size: 1000

Key exclusion criteria

1. Individuals unable to attend for research visits
2. Individuals who in the clinicians opinion is not appropriate for recruitment into this study

Date of first enrolment

06/09/2012

Date of final enrolment

01/04/2022

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

University of Birmingham

Birmingham

United Kingdom

B15 2TT

Sponsor information

Organisation

University of Birmingham (UK)

Sponsor details

Department of Primary Care & General Practice

Primary Care Clinical Sciences Building

Edgbaston

Birmingham

England

United Kingdom

B15 2TT

Sponsor type

University/education

Website

<http://www.birmingham.ac.uk/>

ROR

<https://ror.org/03angcq70>

Funder(s)

Funder type

Industry

Funder Name

Novo Nordisk A/S (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