

Gene-lifestyle interactions and complex traits Involved in elevated disease risk

Submission date 05/10/2017	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 17/11/2017	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 17/11/2017	Condition category Circulatory System	☐ Individual participant data ☐ Record updated in last year

Plain English summary of protocol

Background and study aims

Since 1986, all residents of the northern Swedish county of Västerbotten who turned 40, 50 and /or 60 years are invited to visit the primary care centre. The aim of this study is to use the extensive data collected as part of this ongoing health survey in order to address questions related to lifestyle and genetic risk factors in cardiometabolic disease (e.g., heart disease and type 2 diabetes).

Who can participate?

Adults aged 40, 50, and 60 from Västerbotten

What does the study involve?

Patients complete a detailed lifestyle questionnaire and provide blood samples. The blood sample is used to measure blood glucose (sugar), blood lipids (fat, cholesterol), and genetic information. The data is used to find out whether lifestyle/environmental factors affect the link between genetics and human character (e.g. body mass index, blood glucose).

What are the possible benefits and risks of participating?

Participants receive some health-related advice as part of the background study, which may be perceived as a benefit. Risks include those related to the handling of sensitive personal data. There are no significant treatment-related risks.

Where is the study run from?

Primary Care Center in Västerbotten (Sweden)

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

January 1986 to December 2025

Who is funding the study?

1. Swedish Research Council (Sweden)
2. Novo Nordisk (Denmark)
3. Swedish Heart-Lung Foundation (Sweden)
4. Swedish Diabetes Association (Sweden)

Who is the main contact?
Prof. Paul Franks

Contact information

Type(s)
Scientific

Contact name
Prof Paul Franks

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Additional identifiers

Protocol serial number
Dnr 2015-197-32M

Study information

Scientific Title
Gene-lifestyle interactions and complex traits involved in elevated disease risk: a prospective population-based cohort study

Acronym
GLACIER

Study objectives
The GLACIER study was designed to estimate the separate and combined effects of genetic and lifestyle factors in cardiometabolic disease.

Ethics approval required
Old ethics approval format

Ethics approval(s)
The Regional Ethics Review Board in Umeå, 21/10/2010, ref: Dnr 2015-197-32M

Primary study design
Observational

Study design
Prospective population-based cohort study

Study type(s)

Other

Health condition(s) or problem(s) studied

Cardiovascular disease and type 2 diabetes

Interventions

All participants are asked to complete a questionnaire concerning various lifestyle factors including diet and to donate a blood sample for for later research purposes. The blood samples were then measured for blood glucose, blood lipid profiles (e.g. triglyceride, HDL-cholesterol) and for genetic information. Body weight and height were measured. The study is focused on whether lifestyle/environmental factors influence the association between genetics and human character (e.g. body mass index, blood glucose).

Intervention Type

Mixed

Primary outcome(s)

Many hundreds of exposure and outcome variables including type 2 diabetes, myocardial infarction and stroke, derived from health survey data every 10 years

Key secondary outcome(s)

Secondary outcomes include quantitative cardiometabolic traits (e.g., blood pressure, glucose, and lipids, and anthropometry) usually assessed in the year of each participant's 40, 50, 60th or 70th birthdays. Examinations occurred between 1986 and 2007

Completion date

31/12/2025

Eligibility**Key inclusion criteria**

All individuals 40, 50, and 60 years of age in the county's population are invited

Participant type(s)

Mixed

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

People who are not able to consent or not living in the county

Date of first enrolment

01/01/1986

Date of final enrolment

01/01/2020

Locations

Countries of recruitment

Sweden

Study participating centre

Primary Care Center in Västerbotten

SE-901 87

Sponsor information

Organisation

Umeå University Department of Biobank Research

ROR

<https://ror.org/05kb8h459>

Funder(s)

Funder type

Government

Funder Name

Vetenskapsrådet

Alternative Name(s)

Swedish Research Council, VR

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

Sweden

Funder Name

Novo Nordisk

Alternative Name(s)

Novo Nordisk Global

Funding Body Type

Private sector organisation

Funding Body Subtype

For-profit companies (industry)

Location

Denmark

Funder Name

Hjärt-Lungfonden

Alternative Name(s)

Swedish Heart-Lung Foundation, Hjärt Lungfonden

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Sweden

Funder Name

Diabetesförbundet

Alternative Name(s)

Swedish Diabetes Foundation, Svenska Diabetesförbundet

Funding Body Type

Private sector organisation

Funding Body Subtype

Associations and societies (private and public)

Location

Sweden

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study is not expected to be made available due to consenting constraints.

IPD sharing plan summary

Not expected to be made available