# The Groningen Novice Running 2 (GRONORUN 2) Project

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To determine the effect of a preconditioning (PRECON) program on the development of running related injuries in novice runners compared to a traditional training program (CON) without a preconditioning program.

Ethical review Approved WMO

**Status** Pending

Health condition type Other condition
Study type Interventional

## **Summary**

## ID

NL-OMON31410

#### Source

**ToetsingOnline** 

**Brief title**GRONORUN 2

#### **Condition**

Other condition

## **Synonym**

overuse injuries, running injuries

## **Health condition**

blessures van het skeletspierstelsel van de onderste extremiteit

## Research involving

Human

## **Sponsors and support**

**Primary sponsor:** Universitair Medisch Centrum Groningen **Source(s) of monetary or material Support:** ZONMW

## Intervention

**Keyword:** Prevention, Randomized controlled trial, Risk factors, Running injuries

#### **Outcome measures**

### **Primary outcome**

The primary outcome of the GRONORUN 2 trial is the number of running related injuries (RRI's) in both groups. Definition of a RRI in this trial is; running related musculoskeletal ailment of the lower extremity or back, causing a restriction of running for at least one week, i.e. three consecutive training sessions.

## **Secondary outcome**

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# **Study description**

## **Background summary**

Running is a popular form of recreational exercise. Although running has positive effects on health and fitness, the risk of a running related injury (RRI) has to be considered. The incidence of RRI\*s is high and varies from 30-79%. In novice runners an RRI can negatively affect future physical activity, so the prevention of running injuries especially in novice runners is important.

## Study objective

To determine the effect of a preconditioning (PRECON) program on the development of running related injuries in novice runners compared to a traditional training program (CON) without a preconditioning program.

## Study design

A two arm randomized controlled trial

#### Intervention

A 4 week preconditioning program, with walking and hopping exercise, prior to a 10 week training program in a group of novice runners compared to a 10 week training program without a preconditioning program.

## Study burden and risks

There is no additional risk for participants

Time to fill in the database and/or questionnaires will be 180 minutes in 9 months

## **Contacts**

#### **Public**

Universitair Medisch Centrum Groningen

Hanzeplein 1 9700 RB Groningen NI

## **Scientific**

Universitair Medisch Centrum Groningen

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# **Trial sites**

## **Listed location countries**

**Netherlands** 

# **Eligibility criteria**

## Age

Adults (18-64 years) Elderly (65 years and older)

## Inclusion criteria

Age 18 - 65 years
In last year no running experience
No known cardiovascular disease
No pain from the lower back of lower extremity in the three months prior to inclusion

## **Exclusion criteria**

Pain from the lower back of lower extremity in the three months prior to inclusion

# Study design

## **Design**

Study type: Interventional

Intervention model: Parallel

Allocation: Randomized controlled trial

Masking: Open (masking not used)

**Primary purpose:** Prevention

## Recruitment

NL

Recruitment status: Pending

Start date (anticipated): 01-12-2007

Enrollment: 540

Type: Anticipated

# **Ethics review**

Approved WMO

Application type: First submission

# **Study registrations**

## Followed up by the following (possibly more current) registration

No registrations found.

# Other (possibly less up-to-date) registrations in this register

No registrations found.

## In other registers

Register ID

CCMO NL19668.042.07