# FITstrong: The feasibility of an exercise program for children who survived cancer

Submission date	Recruitment status No longer recruiting	<ul><li>Prospectively registered</li></ul>		
05/09/2007		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
05/09/2007	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
10/06/2021	Cancer			

## Plain English summary of protocol

Not provided at time of registration

# Contact information

#### Type(s)

Scientific

#### Contact name

Dr T. Takken

#### Contact details

University Medical Centre Utrecht (UMCU)/WKZ KB.02.056.0 Lundlaan 6 Utrecht Netherlands 3584EA +31 (0)30 250 4030 t.takken@umcutrecht.nl

# Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

# Study information

#### Scientific Title

The feasibility of an exercise program for children who survived cancer: the FITStrong study

#### Acronym

**FITstrong** 

#### Study objectives

Exercise training is a feasible method to improve fitness (peak oxygen uptake), muscle strength and fatigue in children who survived cancer.

Feasibility will be studied according to the number of performed training sessions and a structured interview with patients and training about their opinion on the training program.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Ethics approval received from University Medical Center Utrecht (The Netherlands) on the 2nd April 2007 (ref: 06/303).

#### Study design

A 12 week aerobic and muscle strength training program (twice a week) compared with baseline training level

#### Primary study design

Interventional

# Secondary study design

Non randomised controlled trial

#### Study setting(s)

Not specified

# Study type(s)

Quality of life

# Participant information sheet

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

Exercise in children who survived cancer

#### Interventions

Two times a week exercise training (45 minutes) starting with warm up, followed by muscle strength components and aerobic components and ended with a cool down. The participants have to perform home exercise two times per week, and at increasing muscle strength /endurance.

# Intervention Type

#### Other

#### **Phase**

**Not Specified** 

## Primary outcome measure

Feasibility of the program.

#### Timepoints:

t=0: baseline

t=1: after 12 weeks of training

t=2: 12 weeks follow-up

#### Secondary outcome measures

1. Peak oxygen uptake (VO2peak)

2. Muscle strength

#### Timepoints:

t=0: baseline

t=1: after 12 weeks of training

t=2: 12 weeks follow-up

#### Overall study start date

01/05/2007

# Completion date

01/01/2008

# **Eligibility**

# Key inclusion criteria

1. 6 - 18 years of age

2. Between 0.5 and 1.5 years since final chemotherapy

#### Participant type(s)

**Patient** 

#### Age group

Child

#### Lower age limit

6 Years

# Upper age limit

18 Years

#### Sex

Both

# Target number of participants

#### Total final enrolment

16

# Key exclusion criteria

Severe cardiomyopathy.

#### Date of first enrolment

01/05/2007

#### Date of final enrolment

01/01/2008

# Locations

#### Countries of recruitment

Netherlands

# Study participating centre

University Medical Centre Utrecht (UMCU)/WKZ

Utrecht Netherlands 3584EA

# Sponsor information

#### Organisation

University Medical Centre Utrecht (UMCU) (The Netherlands)

#### Sponsor details

Wilhelmina Children's Hospital (WKZ)
Department Pediatric Fysiotherapy
Lundlaan 6
Amsterdam
Netherlands
3584 EA
+31 (0)30 250 4030
p.j.m.helders@umcutrecht.nl

# Sponsor type

Hospital/treatment centre

#### Website

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

#### ROR

https://ror.org/04pp8hn57

# Funder(s)

# Funder type

University/education

#### **Funder Name**

Scientific College of Physiotherapy (Wetenschappelijk College Fysiotherapie [WCF]) (The Netherlands)

#### **Funder Name**

RoPaRun Foundation (Stichting RoPaRun) (The 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

# **Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		01/04/2009	10/06/2021	Yes	No