

Skeletal Muscle Metabolism during Rest, Exercise and Recovery in Patients with Cystic Fibrosis and Healthy Controls

Published: 13-05-2011

Last updated: 27-04-2024

To compare the oxygenation and oxidative metabolism in skeletal muscles of patients with CF with healthy controls to provide insight in the metabolism in patients with CF.

Ethical review	Approved WMO
Status	Recruitment stopped
Health condition type	Muscle disorders
Study type	Observational non invasive

Summary

ID

NL-OMON36204

Source

ToetsingOnline

Brief title

Muscle Metabolism in CF

Condition

- Muscle disorders
- Congenital respiratory tract disorders

Synonym

mucoviscidosis

Research involving

Human

Sponsors and support

Primary sponsor: Universitair Medisch Centrum Utrecht

Source(s) of monetary or material Support: verzoek tot co-financiering is ingediend (d.d. 12-04-2011)

Intervention

Keyword: Cystic Fibrosis, muscle metabolism, oxygen delivery, oxygen utilization

Outcome measures

Primary outcome

- * time of PCr recovery (31P MRS)
- Resting, exercise and recovery phoscreatine (PCr)(31P MRS)
- Resting, exercise and recovery Inorganic phosphorus (Pi)(31P MRS)
- Resting, exercise and recovery Pi:PCr ratio(31P MRS)
- Resting, exercise and recovery (Adenosine triphosphate) ATP(31P MRS)
- Resting, exercise and recovery ATP:PCr ratio(31P MRS)Change in concentration deoxygenated hemoglobin. (NIRS)
- Change in concentration oxygenated hemoglobin. (NIRS)
- Change in concentration total hemoglobin. (NIRS)
- Change in concentration deoxygenated hemoglobin minus change in concentration oxygenated hemoglobin. (NIRS)
- The T1/2 (s) change in oxygenated hemoglobin in the recovery phase. (NIRS)

Secondary outcome

- Peak workload (Wpeak)
- Resting and peak oxygen uptake (VO₂peak/kg (ml.min⁻¹.kg⁻¹)
- Resting and peak minute ventilation (VEpeak) (l.min⁻¹)
- Resting and peak respiratory exchange ratio (RER = VCO₂/VO₂)
- Resting and peak heart rate (HR; bpm)
- Resting and peak SpO₂

Study description

Background summary

Patients with CF have a reduced exercise tolerance. Studies show an interrelationship between lung function, muscle mass, energy expenditure, (respiratory) muscle function and exercise capacity in patients with CF. Up to date, there is no scientific evidence or consensus whether the muscle dysfunction is caused by an intrinsic abnormality in muscle energy metabolism and/or is hampered due to impaired oxygen delivery to these muscles, which might be mediated by an increased blood flow demand of the respiratory muscles. Therefore, it remains unclear whether the impaired muscle function is limited by oxygen delivery or oxygen utilization. We hypothesise that patients with CF have impaired oxidative metabolism and/or reduced/slowed oxygen delivery to the skeletal muscles compared to healthy controls.

Study objective

To compare the oxygenation and oxidative metabolism in skeletal muscles of patients with CF with healthy controls to provide insight in the metabolism in patients with CF.

Study design

This is an case-control study. Ten patients with CF and ten healthy controls will be recruited. In two separate test-sessions, thigh muscles of a population of patients with CF and healthy age-matched controls will be measured during rest, incremental cycling exercise, and recovery with NIRS (first test) and, in another test session on a separate day, with ³¹P MRS (second test). The ergometer is made of non-ferrous components, based on the prototype developed by Jeneson et al. for in-magnet bicycling exercise testing (Jeneson et al, 2009).

Before the first (NIRS) test, subjects perform a lungfunction test and adipose tissue thickness is measured. Subjects are asked to fill in a daily activities questionnaire and a MRI and maximal exercise test (only in healthy controls) questionnaire two weeks before the first (NIRS) test day.

Study burden and risks

No burden and risks are reported concerning exercise in a ³¹P-MRS scanner or with NIRS. Participation in this study will take 25 minutes to fill in the questionnaires and will cost 2,5 hours in total on the two separate test sessions. Insight in the muscle metabolism from patients with CF will help to aim and institute interventions as exercise training.

Contacts

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adolescents (12-15 years)

Adolescents (16-17 years)

Inclusion criteria

Patients with CF:

- Ambulant patients with CF
- free from acute pulmonary or gastro-intestinal exacerbation
- from 12 to 18 years

Healthy controls:

- from 12 to 18 years
- free from constraints in performing a maximal exercise test

Exclusion criteria

Patients with CF:

- Resting oxygen saturation (SpO₂) < 94%
- FEV₁ predicted < 80%
- Ineligible to perform an exercise test
- Not familiar with the Dutch language
- Presence of contra-indications for ³¹P MRS measurements (assessed by a standardised questionnaire as previously used in METC 08-267/K)
- Participation in DO-IT or having participated less than six months ago in DO-IT (METC 09-114/K); Healthy controls:
- Ineligible to perform a maximal exercise test (assessed by a standardised questionnaire as previously used in METC 10-468)
- Not familiar with the Dutch language
- Presence of contra-indications for ³¹P MRS measurements (assessed by a standardised questionnaire as previously used in METC 08-267/K)

Study design

Design

Study type:	Observational non invasive
Intervention model:	Other
Allocation:	Non-randomized controlled trial
Masking:	Open (masking not used)
Control:	Active
Primary purpose:	Diagnostic

Recruitment

NL	
Recruitment status:	Recruitment stopped
Start date (anticipated):	31-08-2011
Enrollment:	20
Type:	Actual

Ethics review

Approved WMO	
Date:	13-05-2011
Application type:	First submission

Review commission:	METC Universitair Medisch Centrum Utrecht (Utrecht)
Approved WMO	
Date:	27-06-2011
Application type:	Amendment
Review commission:	METC Universitair Medisch Centrum Utrecht (Utrecht)

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	NL35697.041.11