

# Influence of RYANodyne receptor 1 mutations On Pulmonary arterial Pressure And ventilation During isocapnic hypoxia.

No registrations found.

<b>Ethical review</b>	Positive opinion
<b>Status</b>	Recruiting
<b>Health condition type</b>	-
<b>Study type</b>	Interventional

## Summary

### ID

NL-OMON25834

### Source

Nationaal Trial Register

### Brief title

Ryan op pad

### Health condition

Malignant hyperthermia  
hypoxia  
hypoxic ventilatory response  
hypoxic pulmonary vasoconstriction

## Sponsors and support

**Primary sponsor:** LUMC

**Source(s) of monetary or material Support:** LUMC

## Intervention

## Outcome measures

### Primary outcome

1. Systolic PAP;
2. HVR.

### Secondary outcome

1. Systolic and diastolic cardiac functions;
2. Vital and ventilatory parameters.

## Study description

### Background summary

The ryanodine receptor plays a crucial role in the development of the so called hypoxic pulmonary vasoconstriction. Malignant hyperthermia patients have a mutated ryanodine receptor. In this study we want to investigate the influence of the ryanodine receptor on the human response to hypoxia.

### Study objective

1. What are the normoxic ventilation parameters and systolic PAP in MH patients?
2. What is the influence of hypoxia on the systolic PAP on MH patients?

### Study design

1 hour.

### Intervention

One hypoxic period of 1 hour.

## Contacts

### Public

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## Eligibility criteria

### Inclusion criteria

Healthy Malignant Hyperthermia patients with a phenotypic high susceptibility and a proven causative mutation.

The major inclusion criteria are that the subject has echo evidence of tricuspid regurgitation during systole, which is not clinically relevant but in fact can be demonstrated in most normal individuals.

### Exclusion criteria

1. Obesity (BMI > 30);
2. Presence of medical disease: heart-, lung-, liver-, kidney- and lung disease; diabetes;
3. Presence of psychiatric disease:
  - A. History of chronic alcohol or drug use;
  - B. Possibility of pregnancy;
  - C. Lactation.

## Study design

## Design

Study type:	Interventional
Intervention model:	Parallel
Allocation:	Non-randomized controlled trial
Masking:	Single blinded (masking used)
Control:	Placebo

## Recruitment

NL	
Recruitment status:	Recruiting
Start date (anticipated):	01-05-2011
Enrollment:	24
Type:	Anticipated

## Ethics review

Positive opinion	
Date:	17-02-2011
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
NTR-new	NL2656

**Register**

NTR-old

Other

ISRCTN

**ID**

NTR2784

METC LUMC / CCMO : P11.013 / NL35083.058.11;

ISRCTN wordt niet meer aangevraagd.

## Study results

**Summary results**

N/A