

Can breathing pure oxygen at high pressure improve exercise-induced muscle injury recovery in baseball players?

Submission date
14/03/2019

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
22/03/2019

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
11/02/2020

Condition category
Musculoskeletal Diseases

☐ Individual participant data

Plain English summary of protocol

Background and study aims

Hyperbaric oxygen therapy (HBOT) involves a person breathing pure oxygen at higher pressure than normal air pressure. Air normally contains about 21% oxygen and 78% nitrogen, so HBOT provides more oxygen to the body than is normally breathed in. It is thought that HBOT might results in muscle injuries healing more quickly, which would be an advantage to those playing sport competitively or professionally. This study aims to investigate whether HBOT improved recovery of muscle soreness and strains caused by exercise in baseball players.

Who can participate?

Professional or amateur male baseball players in training or the baseball season who have a muscle injury caused by exercise.

What does the study involve?

The participants will be randomly allocated to one of two groups. Both groups will spend 10 sessions of 100 minutes over 5 weeks in a pressurised chamber, where the pressure will be raised to above normal. One group will breathe pure oxygen while in the chamber and the other will breathe normal air. Oxygen and nitrogen are colourless gases with no smell so the participants will not be able to tell which gas they are breathing. Before the first session, after the fifth session, after the last session and 2 weeks after the last session, the participants will be asked give a blood sample and to rate their muscle pain and how much it affects their daily activities.

What are the possible benefits and risks of participating?

Those who breathe pure oxygen might experience earlier recovery of the muscle injury. There is also a small risk of side effects caused by absorbing too much oxygen.

Where is the study run from?

Kaohsiung Chang Gung Memorial Hospital, Taiwan, Republic of China

When is the study starting and how long is it expected to run for?
August 2013 to August 2016

Who is funding the study?
Chang Gung Research Fund (Taiwan)

Who is the main contact?
Miss Chen-Yu Chen

Contact information

Type(s)
Public

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Additional identifiers

Clinical Trials Information System (CTIS)
Nil known

ClinicalTrials.gov (NCT)
Nil known

Protocol serial number
Chang Gung Research Fund grant CMRPG8D0411

Study information

Scientific Title
Hyperbaric oxygen therapy influence of high-intensity athletes in vivo metabolic indicators

Study objectives
HBOT could facilitate the early recovery of exercise-related muscular injury and could therefore be beneficial for elite athletes.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 21/08/2013, Institutional Review Board (IRB) of Chang Gung Medical Foundation (123 Dinghu Rd, Guishan Township, Taoyuan County, Taiwan (R.O.C.); +886 3 3196200 ext 3707/3703; merlinchi@cgmh.org.tw), ref: 102-2994B

Study design

Prospective randomized double-blind controlled study

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Exercise-related muscular injury

Interventions

41 participants were recruited between June 2014 and December 2015 and were divided into study group and control group with 20 and 21 participants. All the participants received either HBOT or placebo sessions twice in a week. The study group and the control group stayed in a hyperbaric chamber pressurized to 2.5 and 1.3 atm, and breathed pure oxygen and general air, respectively. The duration of each session was 100 mins, and 10 sessions were completed in 5 weeks for each participant.

Intervention Type

Supplement

Primary outcome(s)

1. Serum creatine phosphokinase (CPK)
2. Serum glutamic-oxaloacetic transaminase (GOT)
3. Serum myoglobin
4. Blood urine nitrogen (BUN)
5. Serum lactate

Data were collected before the treatment (T1), end of 5th HBOT (T2), end of 10th HBOT (T3), and 2 weeks after the 10th HBOT (T4).

Key secondary outcome(s)

1. Pain intensity assessed using the Brief Pain Inventory before the treatment (T1), end of 5th HBOT (T2), end of 10th HBOT (T3), and 2 weeks after the 10th HBOT (T4)
2. Pain interference assessed using the Brief Pain Inventory before the treatment (T1), end of 5th HBOT (T2), end of 10th HBOT (T3), and 2 weeks after the 10th HBOT (T4)

Completion date

19/08/2016

Eligibility

Key inclusion criteria

1. Aged 20 years or older
2. Diagnosed with prolonged (more than 2 weeks) exercise-induced muscular soreness or pain with grade I muscle strain of the extremities
3. Currently under intensive and regular baseball training or regular baseball season

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Total final enrolment

41

Key exclusion criteria

1. Pneumothorax
2. Upper respiratory tract infection
3. Recently received chest or ear surgery
4. Claustrophobic

Date of first enrolment

07/07/2014

Date of final enrolment

11/09/2015

Locations**Countries of recruitment**

Taiwan

Study participating centre

Kaohsiung Chang Gung Memorial Hospital

123, Ta Pei Road, Niao Sung District

Kaohsiung

Taiwan

83301

Sponsor information

Organisation

Chang Gung Memorial Hospital

ROR

<https://ror.org/02verss31>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Chang Gung Medical Foundation

Alternative Name(s)**Funding Body Type**

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Taiwan

Results and Publications

Individual participant data (IPD) sharing plan

The datasets generated and/or analyzed during the current study during this study will be included in the subsequent results publication.

IPD sharing plan summary

Other

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	29/05/2019	11/02/2020	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes