

Resistance training and leucine supplementation : effects on performance and body composition of untrained subjects.

Submission date
30/09/2005

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
30/09/2005

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
23/08/2012

Condition category
Nutritional, Metabolic, Endocrine

☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0436146689

Study information

Scientific Title

Study objectives

To investigate whether long-term leucine supplementation in combination with resistance training can induce higher increases in strength and muscle mass of untrained subjects compared to a placebo group undertaking resistance training only.

To investigate potential decreases in muscle mass after four weeks following the end of the supplementation period

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised placebo controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Nutritional, Metabolic, Endocrine: Leucine supplementation

Interventions

Randomised controlled trial

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Bone density

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/12/2003

Completion date

01/09/2004

Eligibility

Key inclusion criteria

Healthy male subjects

Participant type(s)

Patient

Age group

Adult

Sex

Male

Target number of participants

30 male volunteers

Key exclusion criteria

1. Athletes or trained subjects
2. Having diabetes
3. Heart disease
4. High blood pressure or any other chronic or serious illness
5. Having taken anabolic steroids or any other sports supplements at least 2 months before the beginning of the study

Date of first enrolment

01/12/2003

Date of final enrolment

01/09/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

School of leisure and sports studies

Leeds

United Kingdom

LS1 3HE

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

+44 (0)20 7307 2622

dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Government

Funder Name

Leeds Teaching Hospitals NHS Trust (UK)

Funder Name

NHS R&D Support Funding

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	results	01/03/2011		Yes	No