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

| Submission date              | Recruitment status  No longer recruiting | <ul><li>Prospectively registered</li></ul> |  |  |
|------------------------------|--|--|--|--|
| 30/09/2005                   |  | ☐ Protocol                                 |  |  |
| Registration date 30/09/2005 | Overall study status Completed           | Statistical analysis plan                  |  |  |
|                              |  | [X] Results                                |  |  |
| Last Edited                  | Condition category                       | Individual participant data                |  |  |
| 23/08/2012                   | Nutritional. Metabolic. Endocrine        |  |  |  |

### Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Mr T Ispoglou

### 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

### Kev 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              |