

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

The Effect of 6 Weeks Resistance Training and Combined Resistance Training with Pulsed Electromagnetic Stimulation on bone biomarkers and Balance in Severe Haemophilia A with Osteoporosis

Protocol summary

Summary

This study was aimed to investigate the effects of a short-term resistance training program and pulsed electromagnetic fields (PEMFs) on bone metabolism, balance, joint function, and quality of life in severe haemophilia A with osteoporosis. Forty-three severe haemophilia A with osteoporosis were randomly assigned to resistance training (RT), combined resistance training with PEMFs (RTPEMF), PEMFs, and control groups. The RT and RTPEMF groups performed progressive resistance exercises (50-60% 1RM); in RTPEMF, the lower repetition of exercises was compensated with 30-min exposure to PEMFs. The PEMF group was exposed to 60 min of PEMFs (30 Hz and 40 Gauss). The intervention was performed 3 days weekly for 6 weeks. Bone-specific alkaline phosphatase (BALP), N-terminal telopeptide of type 1 collagen (NTX), joint function, balance, and quality of life were assessed before and after the program. After 6 weeks, joint physical conditions, balance performance and quality of life showed significant improvement in RT and RTPEMF groups. BALP increased significantly in RT and RTPEMF groups. No groups showed significant changes in NTX. The best pain relief occurred in the knee joint of the RT group. Resistance training can increase bone metabolism and improve balance performance, joint physical condition muscle strength, and quality of life in severe haemophilia A with osteoporosis.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201010174952N1**
Registration date: **2013-04-24, 1392/02/04**
Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2013-04-24, 1392/02/04

Registrant information

Name

Giti Torkaman

Name of organization / entity

Tarbiat Modares University

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Iran (Islamic Republic of)

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Recruitment status

Recruitment complete

Funding source

Tarbiat Modares university

Expected recruitment start date

2010-10-22, 1389/07/30

Expected recruitment end date

2011-11-21, 1390/08/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The Effect of 6 Weeks Resistance Training and Combined Resistance Training with Pulsed Electromagnetic Stimulation on bone biomarkers and Balance in Severe Haemophilia A with Osteoporosis

Public title

resistance training and electromagnetic fields in

osteoporosis

Purpose

Treatment

Inclusion/Exclusion criteria

Include Criteria: sever hemophilia-A; T score less than 2.5. Exclude Criteria: no history of hepatitis B and C; no HRT in the last 6 months; no acute target joint, no history of inhibitors.

Age

From **18 years** old to **35 years** old

Gender

Male

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **43**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Single blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Tarbiat modares university

Street address

ale-ahmad Ave.

City

tehran

Postal code

1411713116

Approval date

2010-10-08, 1389/07/16

Ethics committee reference number

52/99424

2

Ethics committee

Name of ethics committee

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Street address

ale-ahmad Ave.

City

tehran

Postal code

1411713116

Approval date

2010-08-20, 1389/05/29

Ethics committee reference number

52/99424

Health conditions studied

1

Description of health condition studied

hemophilia with osteoporosis

ICD-10 code

M82.8

ICD-10 code description

Osteoporosis in other diseases classified elsewhere

Primary outcomes

1

Description

NTX

Timepoint

Before and 6 weeks after intervention

Method of measurement

ELISA Kit

2

Description

CTX

Timepoint

Before and 6 weeks after intervention

Method of measurement

ELISA Kit

3

Description

Balance and Strength

Timepoint

Before and 6 weeks after intervention

Method of measurement

physical assesment and observation

4

Description

Quality of Life

Timepoint

Before and 6 weeks after intervention

Method of measurement

Questionnaire

Secondary outcomes

1

Description

joint range of motion

Timepoint

Before and 6 weeks after intervention
Method of measurement
goniameter

2

Description
joint swelling
Timepoint
Before and 6 weeks after intervention
Method of measurement
physical assesment

Intervention groups

1

Description
6-week trunk, upper and lower limb progressive resistance training; 3 days in week; 50-60% 1repeated maximum; 2 sets of 10-15 repetition
Category
Rehabilitation

2

Description
6-week trunk, upper and lower limb progressive resistance training; 3 days in week; 50-60% 1repeated maximum; 2 sets of 5-10 repetition and 30 min electromagnetic fields with frequency of 30 Hz and intensity of 40 Gauss
Category
Rehabilitation

3

Description
Control
Category
N/A

Recruitment centers

1

Recruitment center
Name of recruitment center
seydalshohada hospital
Full name of responsible person
Street address
City
Esfahan

Sponsors / Funding sources

1

Sponsor
Name of organization / entity
tarbiat modares university
Full name of responsible person

Dr Yaghoob Fathollahi
Street address
Vice-Chancellor for Reaserch Affairs,Tarbiat Modares University,Ale-Ahmad Ave.

City
Tehran

Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?

Yes
Title of funding source
tarbiat modares university
Proportion provided by this source
100

Public or private sector
empty
Domestic or foreign origin
empty
Category of foreign source of funding
empty
Country of origin
Type of organization providing the funding
empty

Person responsible for general inquiries

Contact
Name of organization / entity
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Person responsible for updating data

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City

Sharing plan

Deidentified Individual Participant Data Set (IPD)
empty
Study Protocol
empty
Statistical Analysis Plan
empty
Informed Consent Form
empty
Clinical Study Report
empty
Analytic Code
empty
Data Dictionary
empty