

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

Comparison of the efficacy of intra-muscular Botulinum toxin type A with oral Tizanidine in the treatment of upper limb spasticity and functional improvement due to cerebral stroke.

Protocol summary

Summary

Our aim in this study is comparison of the Botulinum toxin type A with oral Tizanidine in the treatment of spasticity due to stroke. The key inclusion criteria was to have a history of cerebral stroke for at least 3 months ago and the key exclusion criteria was to have a systemic disorder. In a randomized unicentral single blinded study 68 patients from both genders with upper limb spasticity referring to the neurology clinic of Emam Reza hospital, Tabriz, Iran, were assigned in two equally-sized groups to receive oral Tizanidine or intra-muscular Botulinum toxin. For the first group, Tizanidine will be started with 2 mg/day and added 2 mg/week until the dose of 16 mg/day in week 16. Patients will be assessed in week 16. In Botulinum group, patients will be injected two doses of Botulinum into the upper limb muscles with a 3-4 month time gap. Limb spasticity in Ashworth scale and Limb function improvement in ARAT scale in the start of the study for all the patients, week 16 (for Tizanidine) and 3 months after the last injection (for Botulinum) will be measured and the results will be compared within and between the groups.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2012112411563N1**

Registration date: **2012-12-06, 1391/09/16**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2012-12-06, 1391/09/16

Registrant information

Name

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Name of organization / entity

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Recruitment status

Recruitment complete

Funding source

Tabriz University of Medical Sciences

Expected recruitment start date

2012-04-04, 1391/01/16

Expected recruitment end date

2012-12-06, 1391/09/16

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the efficacy of intra-muscular Botulinum toxin type A with oral Tizanidine in the treatment of upper limb spasticity and functional improvement due to cerebral stroke.

Public title

Comparison of the Botulinum toxin type A with oral Tizanidine in the treatment of spasticity due to stroke

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: (to have a history of cerebral stroke for

at least 3 months ago; severity of spasticity more than 2 in Ashworth scale; to be older than 35 years old)
Exclusion criteria: (patients with impairment in level or content of consciousness; to have systemic disorders such as Diabetes, hyperthyroidism, electrolyte disorders, renal failure; to be older than 70 years old)

Age

From **35 years** old to **70 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **68**

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**

Ethics Committee of Tabriz University of Medical Sciences

Street address

Vice chancellor for research, Tabriz University of Medical Sciences, Daneshgah Square

City

Tabriz

Postal code**Approval date**

2012-04-03, 1391/01/15

Ethics committee reference number

9122

Health conditions studied**1****Description of health condition studied**

Spasticity due to cerebral stroke

ICD-10 code

I69.3

ICD-10 code description

Sequelae of cerebral infarction

Primary outcomes**1****Description**

Limbs Spasticity

Timepoint

in the start of the study for all the patients, week 16 (for Tizanidine) and 3 months after the last injection (for Botulinum)

Method of measurement

Based on Ashworth scale

2**Description**

Limb functional improvement

Timepoint

in the start of the study for all the patients, week 16 (for Tizanidine) and 3 months after the last injection (for Botulinum)

Method of measurement

Based on ARAT scale

Secondary outcomes

empty

Intervention groups**1****Description**

For Tizanidine group: Tizanidine will be started with 2 mg/day orally and added 2 mg/week until the dose of 16 mg/day in week 16.

Category

Treatment - Drugs

2**Description**

In Botulinum group, patients will be injected two doses of Botulinum (not more than 1000 units/dose) into the upper limb muscles with a 3-4 month time gap.

Category

Treatment - Drugs

Recruitment centers**1****Recruitment center****Name of recruitment center**

Neurology clinic of Emam Reza hospital

Full name of responsible person

Mohammad Yazdchi

Street address**City**

Tabriz

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Tabriz University of Medical Sciences

Full name of responsible person

Alireza Ostad Rahimi

Street address

Vice chancellor for research, Tabriz University of Medical Sciences, Daneshgah square

City

Tabriz

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

Yes

Title of funding source

Tabriz University of Medical Sciences

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**

Tabriz University of Medical Sciences

Full name of responsible person

Zahra Ghasemi

Position

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Person responsible for scientific

inquiries

Contact**Name of organization / entity**

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Full name of responsible person

Mohammad Yazdchi

Position

Associate professor/Neurologist

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Web page address

Person responsible for updating data

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Full name of responsible person

Mehdi Hemmati

Position

General physician

Other areas of specialty/work**Street address****City****Postal code****Phone****Fax****Email**

mehdi.hemmati@rocketmail.com

Web page address

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