

Clinical Trial Protocol

Iranian Registry of Clinical Trials

03 Jul 2026

Evaluation of the effect of Atrovastatin on inflammatory markers in sulfur mustard gas induced chronic bronchitis

Protocol summary

Summary

The aim of this study is evaluation of the effect of Atrovastatin on inflammatory markers in sulfur mustard gas induced chronic bronchitis. This is a single center, double blind randomized phase III clinical trial. The main inclusion criteria are exposure to mustard gas during the Iran-Iraq war and suffering from chronic bronchitis (history of chronic cough and sputum daily for 3 months during recent 2 years). The main exclusion criteria are using statin in the last 3 months; exacerbation of symptoms in the past 4 weeks; and smoking. We will enroll 90 patients with mustard gas induced chronic bronchitis. Participants will randomly be assigned into intervention and control groups and will receive Atrovastatin (40 mg per day) and placebo (one tablet per day) respectively for 3 months. As the primary outcome of our study, the level of IL 6, TNF α , procalcitonin, and hs-CRP will be measured before and after intervention. We will compare the results between groups.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT138904144312N1**

Registration date: **2014-08-16, 1393/05/25**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2014-08-16, 1393/05/25

Registrant information

Name

Seyed Masoom Masoompour

Name of organization / entity

Shiraz University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 71 3647 4316

Email address

masoomm@sums.ac.ir

Recruitment status

Recruitment complete

Funding source

Partly by Shiraz University of Medical Sciences and partly by Project Director will be provided

Expected recruitment start date

2014-06-29, 1393/04/08

Expected recruitment end date

2015-06-29, 1394/04/08

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of the effect of Atrovastatin on inflammatory markers in sulfur mustard gas induced chronic bronchitis

Public title

Effect of Atrovastatin on lung inflammation in bronchitis

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: exposure to mustard gas during the Iran-Iraq war and have suffered from chronic bronchitis (history of chronic cough and sputum daily for 3 months during recent 2 years) Exclusion criteria: Using statin in the last 3 months of study starting time; exacerbation of symptoms in the past 4 weeks; connective tissue disease; sarcoidosis; eosinophilic granuloma; pneumoconiosis; lymphoma; carcinomatous; smokers

Age

From **45 years** old to **75 years** old

Gender

Male

Phase

3

Groups that have been masked

No information

Sample size

Target sample size: **90**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Double blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of Shiraz University of Medical Sciences

Street address

Headquarters Of Shiraz University of Medical Sciences
- Zand St - Shiraz

City

Shiraz

Postal code

Approval date

2014-06-28, 1393/04/07

Ethics committee reference number

CT-P-9378-7109

Health conditions studied

1

Description of health condition studied

Chronic bronchitis

ICD-10 code

J44

ICD-10 code description

Chronic bronchitis: obstructive

Primary outcomes

1

Description

TNF alpha level

Timepoint

before intervention - 3 month after intervention

Method of measurement

ELISA test is performed

2

Description

interleukin 6 level

Timepoint

before intervention - 3 month after intervention

Method of measurement

ELISA test is performed

3

Description

Added at 2015-08-05: Procalcitonin

Timepoint

Added at 2015-08-05: At the beginning of the intervention and 3 months after intervention

Method of measurement

Added at 2015-08-05: ELISA

Secondary outcomes

1

Description

With blood cell count

Timepoint

before intervention - 3 month after intervention

Method of measurement

Complete Blood Count

2

Description

Added at 2015-08-05: hs-CRP

Timepoint

Added at 2015-08-05: At the beginning the intervention and 3 months after intervention

Method of measurement

Added at 2015-08-05: Immunoassay

Intervention groups

1

Description

Intervention group:40 mg atorvastatin daily for 3 months

Category

Treatment - Drugs

2

Description

Control group: one placebo tablet daily for 3 months

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Mohammd Rasul Allah Clinic

Full name of responsible person

Dr Seyed Masoom Masoompour

Street address

Pasdarn Blvd, Shiraz

City

shiraz

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

shiraz University Of medical Science

Full name of responsible person

Dr Seyed Baser Hashemi

Street address

Building of Shiraz University of Medical Sciences,
Zand Ave

City

Shiraz

Grant name

Grant code / Reference number

Is the source of funding the same sponsor organization/entity?

Yes

Title of funding source

shiraz University Of medical Science

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

Shiraz University of Medical Sciences

Full name of responsible person

Dr seyed Masoom Masoompour

Position

Associate professor

Other areas of specialty/work

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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