

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jun 2026

### Evaluation of Blood Inflammatory Markers and Their Comparison with Exercise Capacity Indices in Patients with Chronic Obstructive Pulmonary Disease Following Short-Term Pulmonary Rehabilitation

#### Protocol summary

##### Study aim

Evaluation of Blood Inflammatory Markers and Their Comparison with Exercise Capacity Indices in Patients with Chronic Obstructive Pulmonary Disease Following Short-Term Pulmonary Rehabilitation

##### Design

In this study, which will be conducted in phase III of a clinical trial, a total of 206 patients will be enrolled according to the inclusion and exclusion criteria. After obtaining written informed consent, they will be randomly assigned (using permuted block randomization) in a single-blind manner into two groups: intervention and control.

##### Settings and conduct

This study is a phase III clinical trial conducted in a randomized (using permuted block randomization) and single-blind manner on patients with chronic obstructive pulmonary disease (COPD) who are referred to Masih Daneshvari Hospital.

##### Participants/Inclusion and exclusion criteria

The inclusion criteria for the study include a definite diagnosis of chronic obstructive pulmonary disease based on the Global Initiative for Chronic Obstructive Lung Disease (GOLD) criteria, age above 18 years, stability of the patient's clinical condition at the time of entry into the study, and the ability to perform functional tests. Patients who develop acute respiratory symptoms or infection during the rehabilitation period, patients with a history of exacerbation, or those who are unwilling to cooperate will be excluded from the study.

##### Intervention groups

In the control group, patients receive respiratory physiotherapy and bedside exercises. In the intervention group, in addition to active care, training in caregiving techniques, and bedside physiotherapy, patients receive the full rehabilitation program and participate in a pulmonary rehabilitation program.

##### Main outcome variables

Platelet-to-lymphocyte ratio, erythrocyte sedimentation rate, neutrophil-to-lymphocyte ratio

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20210813052172N4**

Registration date: **2026-05-05, 1405/02/15**

Registration timing: **prospective**

Last update: **2026-05-05, 1405/02/15**

Update count: **0**

##### Registration date

2026-05-05, 1405/02/15

##### Registrant information

##### Name

Masoumeh ZoghAli

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 21 3662 4225

##### Email address

masoumezoghali@sbmu.ac.ir

##### Recruitment status

**Not yet recruiting**

##### Funding source

##### Expected recruitment start date

2026-06-22, 1405/04/01

##### Expected recruitment end date

2027-03-21, 1406/01/01

##### Actual recruitment start date

empty

**Actual recruitment end date**

empty

**Trial completion date**

empty

**Scientific title**

Evaluation of Blood Inflammatory Markers and Their Comparison with Exercise Capacity Indices in Patients with Chronic Obstructive Pulmonary Disease Following Short-Term Pulmonary Rehabilitation

**Public title**

Assessment of the Effect of Pulmonary Rehabilitation in Patients with Chronic Obstructive Pulmonary Disease

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Definitive diagnosis of COPD based on GOLD criteria  
Age over 18 years  
Stability of the patient's clinical condition at the time of study entry  
Ability to perform functional tests

**Exclusion criteria:**

Exacerbation of the disease within the past two weeks  
Acute respiratory symptoms or infection during pulmonary rehabilitation period  
Incomplete training sessions for any reason  
Lack of willingness to cooperate in performing functional tests or completing questionnaires

**Age**

From **18 years** old

**Gender**

Both

**Phase**

3

**Groups that have been masked**

- Outcome assessor
- Data analyser

**Sample size**

Target sample size: **206**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

The method used is permuted block randomization. For this purpose, two treatment groups, A and B, are defined in blocks as AB and BA. Then, numbers from a random table in the range 0 to 9 are considered. Numbers 0 to 4 are assigned to block AB, and numbers 5 to 9 are assigned to block BA. Random numbers are then selected from the table. If the number 0 appears, it corresponds to block AB, and therefore two individuals enter this block, such that the first individual receives treatment A and the second individual receives treatment B. In the same way, the treatment groups for the remaining participants are determined. Although in this method the number of observations in both groups will be equal, because of the small block sizes, there is a high probability that the person conducting the study may predict the treatment group assignment. To solve this problem, the randomization list is prepared before the start of the study by a blinded individual who is not

part of the treatment team. Additionally, the block sizes will be increased.

**Blinding (investigator's opinion)**

Single blinded

**Blinding description**

Given the nature of the study, which involves pulmonary rehabilitation, blinding of participants and therapists is not feasible, as patients will be aware of the type of treatment they receive. To minimize potential bias, the outcome assessor and the statistical data analyst will be blinded to group allocation. For this purpose, patient information will be provided to the outcome assessor and the statistical analyst in a coded format, and they will not have access to information regarding the type of intervention. Data analysis will be conducted solely based on the assigned codes

**Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

**Ethics committees****1****Ethics committee****Name of ethics committee**

Ethics Committee of School of Medicine, Shahid Beheshti University of Medical Sciences

**Street address**

Shahid Abbas Arabi St., Yemen St., Shahid Chamran Highway

**City**

Tehran

**Province**

Tehran

**Postal code**

1956944413

**Approval date**

2026-02-17, 1404/11/28

**Ethics committee reference number**

IR.SBMU.MSP.REC.1404.763

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

Chronic obstructive pulmonary disease

**ICD-10 code**

J44.9

**ICD-10 code description**

Chronic obstructive pulmonary disease, unspecified

**Primary outcomes**

## 1

### **Description**

Neutrophil-to-lymphocyte ratio

### **Timepoint**

Before starting pulmonary rehabilitation and after completing 10 days of pulmonary rehabilitation

### **Method of measurement**

Complete Blood Count

## 2

### **Description**

Platelet-to-lymphocyte ratio

### **Timepoint**

Before starting pulmonary rehabilitation and after completing 10 days of pulmonary rehabilitation

### **Method of measurement**

Complete Blood Count

## 3

### **Description**

Erythrocyte sedimentation

### **Timepoint**

Before starting pulmonary rehabilitation and after completing 10 days of pulmonary rehabilitation

### **Method of measurement**

Complete Blood Count

## **Secondary outcomes**

## 1

### **Description**

Exercise capacity

### **Timepoint**

Before initiation pulmonary rehabilitation and after 10 days pulmonary rehabilitation

### **Method of measurement**

Six minute walk test

## 2

### **Description**

Strength and endurance of the lower limb muscles

### **Timepoint**

Before starting pulmonary rehabilitation and after completing 10 days of pulmonary rehabilitation

### **Method of measurement**

Sit to stand test

## 3

### **Description**

Dyspnea

### **Timepoint**

Before starting pulmonary rehabilitation and after completing 10 days of pulmonary rehabilitation

### **Method of measurement**

Borg scale questionnaire

## 4

### **Description**

Fatigue

### **Timepoint**

Before starting pulmonary rehabilitation and after completing 10 days of pulmonary rehabilitation

### **Method of measurement**

Borg scale questionnaire

## 5

### **Description**

Quality of life

### **Timepoint**

Before starting pulmonary rehabilitation and after completing 10 days of pulmonary rehabilitation

### **Method of measurement**

St. George's Questionnaires and Chronic Obstructive Pulmonary Disease Assessment Test (CAT)

## **Intervention groups**

## 1

### **Description**

Intervention group: In addition to active care, patients receive education on care techniques and bedside physiotherapy, undergo a full rehabilitation program, and participate in a structured 10-session pulmonary rehabilitation program.

### **Category**

Rehabilitation

## 2

### **Description**

Control group: Patients receive active care and only basic respiratory education and care techniques (teaching sitting posture, simple diaphragmatic breathing, pursed-lip breathing), education and counseling (medication management, breathlessness management, teaching breathlessness control), and respiratory physiotherapy and bedside exercises (range-of-motion exercises and strengthening of respiratory muscles in a seated position in bed, isometric exercises, and prescribed exercises to maintain daily activities).

### **Category**

Rehabilitation

## **Recruitment centers**

## 1

### **Recruitment center**

#### **Name of recruitment center**

Masih Daneshvari Hospital

#### **Full name of responsible person**

Masoumeh Zoghali

#### **Street address**

Masih Daneshvari Hospital, Daarabad

#### **City**

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masoumezoghali@gmail.com

## Sponsors / Funding sources

### 1

#### Sponsor

**Name of organization / entity**  
Shahid Beheshti University of Medical Sciences  
**Full name of responsible person**  
Afshin Zarghi  
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Shahid Abbas Arabi St., Yemen St., Shahid Chamran  
Highway  
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zarghi@sbmu.ac.ir

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

Shahid Beheshti University of Medical Sciences

#### Proportion provided by this source

100

#### Public or private sector

Public

#### Domestic or foreign origin

Domestic

#### Category of foreign source of funding

empty

#### Country of origin

#### Type of organization providing the funding

Academic

## Person responsible for general inquiries

#### Contact

**Name of organization / entity**  
Shahid Beheshti University of Medical Sciences  
**Full name of responsible person**  
Maryam Sadat Mirenayat  
**Position**  
Associate professor  
**Latest degree**  
Subspecialist  
**Other areas of specialty/work**

Internal Medicine

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## Person responsible for scientific inquiries

#### Contact

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## Person responsible for updating data

#### Contact

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#### Position

Researcher

#### Latest degree

Master

#### Other areas of specialty/work

Medical Biotechnology

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zahirireyhane@gmail.com

**Sharing plan****Deidentified Individual Participant Data Set (IPD)**

Undecided - It is not yet known if there will be a plan to make this available

**Study Protocol**

Undecided - It is not yet known if there will be a plan to make this available

**Statistical Analysis Plan**

Undecided - It is not yet known if there will be a plan to make this available

**Informed Consent Form**

Undecided - It is not yet known if there will be a plan to make this available

**Clinical Study Report**

Undecided - It is not yet known if there will be a plan to make this available

**Analytic Code**

Undecided - It is not yet known if there will be a plan to make this available

**Data Dictionary**

Undecided - It is not yet known if there will be a plan to make this available