

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

29 May 2026

### Effect of Erythropoietin on the treatment of Anemia of prematurity

#### Protocol summary

##### Summary

This is a prospective study that determined the effect of erythropoietin in treatment of anemia of prematurity in newborns that were discharged from NICU of Emamreza hospital. We included premature infants who had Hct<30% in second week or Hct <25% in third week. Intervention group received erythropoietin 500 unites/kg twice in week along with supplementary iron. Control group received only supplementary iron for 4 weeks. Hematocrite and reticulocyte count checked at the beginning, third day after the start and one week after the end of treatment.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT138711031162N10**

Registration date: **2009-04-01, 1388/01/12**

Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

2009-04-01, 1388/01/12

##### Registrant information

###### Name

Ashraf Mohammadzadeh

###### Name of organization / entity

Neonatal Research Center of Mashhad University of  
Medical Sciences

###### Country

Iran (Islamic Republic of)

###### Phone

+98 51 1852 1121

###### Email address

mohamadzadeha@mums.ac.ir

#### Recruitment status

##### Recruitment complete

##### Funding source

Mashhad University of medical sciences

##### Expected recruitment start date

2001-05-04, 1380/02/14

##### Expected recruitment end date

2002-05-04, 1381/02/14

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Effect of Erythropoietin on the treatment of Anemia of  
prematurity

##### Public title

Effect of Erythropoietin on the treatment of Anemia of  
prematurity

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: Gestational age less than 34 weeks,  
breast milk feeding, hematocrite <30% when infants age  
was between 2 to 3 weeks or HCT<25% when infants  
age was more than 3 weeks. Exclusion criteria: Infants  
with hemolytic anemia, active bleeding or congenital  
malformation.

##### Age

No age limit

##### Gender

Both

##### Phase

2

##### Groups that have been masked

No information

##### Sample size

Target sample size: **20**

##### 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**  
Vice chancellor for research of Mashhad University of medical sciences  
**Street address**  
Ghoreshi building, Daneshgah street  
**City**  
Mashhad  
**Postal code**  
13131-91379  
**Approval date**  
2002-08-05, 1381/05/14  
**Ethics committee reference number**  
1427 ت

## Health conditions studied

### 1

**Description of health condition studied**  
Anemia of prematurity  
**ICD-10 code**  
P61.2  
**ICD-10 code description**  
Anaemia of prematurity

## Primary outcomes

### 1

**Description**  
Hematocrite and reticulocyte count  
**Timepoint**  
at the beginning, third day from the start day and one week after the end of treatment  
**Method of measurement**  
venous sampling

## Secondary outcomes

empty

## Intervention groups

### 1

**Description**  
Intervention group: erythropoietin 500 unite/kg twice in week along with supplementary iron  
**Category**  
Treatment - Drugs

### 2

**Description**  
Control group: only supplementary iron for 4 weeks  
**Category**  
Treatment - Drugs

## Recruitment centers

### 1

**Recruitment center**  
**Name of recruitment center**  
NICU of Emamreza hospital  
**Full name of responsible person**  
Dr.Ashraf mohammadzadeh  
**Street address**  
Neonatal research center- NICU- Emamreza hospital- mashhad-Iran  
**City**  
Mashhad

## Sponsors / Funding sources

### 1

**Sponsor**  
**Name of organization / entity**  
Mashhad University of medical sciences  
**Full name of responsible person**  
Dr.jalil tavakol afshari  
**Street address**  
Ghoreshi bulding-Daneshgah street  
**City**  
Mashhad  
**Grant name**  
**Grant code / Reference number**  
**Is the source of funding the same sponsor organization/entity?**  
Yes  
**Title of funding source**  
Mashhad 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**

Neonatal Research Center

**Full name of responsible person**

Dr. Ashraf mohammadzadeh

**Position**

Neonatologist, professor

**Other areas of specialty/work**

**Street address**

NRC-NICU-Emamreza hospital

**City**

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**Postal code**

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## Person responsible for scientific inquiries

### Contact

**Name of organization / entity**

Neonatal Research Center

**Full name of responsible person**

Dr. Ashraf mohammadzadeh

**Position**

Neonatologist, professor

**Other areas of specialty/work**

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## Person responsible for updating data

### Contact

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Neonatal Research Center

**Full name of responsible person**

Dr.Ashraf mohammadzadeh

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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