

Clinical Trial Protocol

Iranian Registry of Clinical Trials

13 Jun 2026

Determination of mgso4 infusion effect for treatment of patient with acute pulmonary edema arrived rasol and sina hospital from khordad 1391 to khordad 1392

Protocol summary

Summary

Loop diuretics and thiazides are mainstay of cardiac acute pulmonary edema treatment. These drugs lead to excretion of Mg in urine, that decreasing Mg in intra and extracellular fluids. In past trial reported depletion of Mg in intra and extracellular fluid by one month treatment of lasix or hydrochlorotiazid. and so Mg depletion leads to dysrhythmia, neuromuscular dysfunction, hypokalemia and hypocalcemia that resistance to treatment in CHF patient under treatment by diuresis. also MgSo4 infusion (by depletion both catecholamine release and peripheral vascular resistance) leads to both repletion and improving noncardiac pulmonary edema (in pheochromocytoma). It causes to improvement of respiratory failure and decreased intubation in severe asthma attack. In this trial, goal is whether that MgSo4 infusion in decreased hospitalization duration, decreased intubation, decreased dysrhythmia incidence that should be treat in ED, improvement both SaO2 and dyspnea score (verbal quantitative scale) in 6 hours after treatment initiation? this trial is doing randomized single blind clinical trial. Patients with acute cardiac pulmonary edema divided in 2 groups randomizely after that explained about trial and side effect of Mgso4 infusion. one group name is intervention that giving to patient 1 gr Mgso4 in 100 cc normal saline for 20 min, another group name is control that 100 cc normal saline infused for 20 min as placebo. both groups patients received Loop diuretics or thiazides in last month ago. if intervention patients show symptoms of Mgso4 toxicity (eg: patella areflexia or pulmonary depression) or with renal failure, exit this group. Primary outcome measure are MgSo4 infusion in decreased hospitalization duration, decreased intubation, decreased dysrhythmia incidence that should be treat in ED, improvement both SaO2 and dyspnea score (verbal quantitative scale) in 6 hours after treatment initiation

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201203089235N1**
Registration date: **2012-10-16, 1391/07/25**
Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2012-10-16, 1391/07/25

Registrant information

Name

Seyyed Mahdi Zia Ziabari

Name of organization / entity

Tehran University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 13 1551 7531

Email address

sm-ziabari@razi.tums.ac.ir

Recruitment status

Recruitment complete

Funding source

publicdomain

Expected recruitment start date

2012-06-05, 1391/03/16

Expected recruitment end date

2013-06-06, 1392/03/16

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Determination of mgso4 infusion effect for treatment of patient with acute pulmonary edema arrived rasol and sina hospital from khordad 1391 to khordad 1392

Public title

Determination of mgso4 infusion effect for treatment of patient with acute pulmonary edema arrived rasol and sina hospital from khordad 1391 to khordad 1392

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria:patient with acute pulmonary edema

Exclusion criteria:presence of mgso4 side effects

Age

No age limit

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: 40

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Single blinded

Blinding description

Placebo

Used

Assignment

Single

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Tehran university medical science

Street address

Tehran porsina street north aspect of tehran university

City

Tehran

Postal code

1417613151

Approval date

2012-06-05, 1391/03/16

Ethics committee reference number

130/272/91/5

Health conditions studied

1

Description of health condition studied

Acute pulmonary edema

ICD-10 code

150.1

ICD-10 code description

with mention of heart disease NOS or heart failure

Primary outcomes

1

Description

hospitalization duration

Timepoint

for 6 hours after trial initiation

Method of measurement

Compare of hospitalization duration in two groups

2

Description

dysrhythmia incidence that should be treat in ED

Timepoint

for 6 hours after trial initiation

Method of measurement

Compare of demand to intubation in two groups

3

Description

necessary for intubation

Timepoint

for 6 hours after trial initiation

Method of measurement

Compare of incidence of dysrhythmia that demand treatment in two groups

4

Description

improvement both SaO2 and dyspnea score(verbal quantitative scale) in 6 hours after treatment initiation

Timepoint

for 6 hours after trial initiation

Method of measurement

Compare of Sao2 improvement more than 94% and dyspnea score (by verbal quantitative scale pathway) for 6 hours after treatment initiation in two groups

Secondary outcomes

1

Description

Decrease respiratory rate under 8/min

Timepoint

for 6 hours from tretment initiation

Method of measurement

Account respiratory rate in 1 minute

2

Description

Diminish of deep tendon reflex

Timepoint

for 6 hours after treatment initiation

Method of measurement

Monitoring of deep tendon reflex

Intervention groups

1

Description

In intervention group start treatment by o2 until o2Sat of patient >94% (minimum by 4 lit/min o2), plus Amp lasix 1 mgr/kg and Amp nitroglycerin 5 mic gr/min started and titrated 5 mic gr/min if patients blood pressure permitted, and Amp morphine sulfate 0.05 mgr/kg for attention of patient blood pressure (preferly systolic blood pressure > 90 mmhg). these treatment are basic treatment of acute pulmonary edema. in addition to in intervention group gives 1 gr Mgso4 in 100 cc normal saline for 20 min.

Category

Treatment - Drugs

2

Description

In control group start treatment by o2 until o2Sat of patient >94% (minimum by 4 lit/min o2), Amp lasix 1 mgr/kg and Amp nitroglycerin by 5 mi cgr/min started and titrated 5 mi cgr/min if patients blood pressure permitted, and Amp morphine sulfate 0.05 mgr/kg for attention of patient blood pressure (preferly systolic blood pressure > 90 mm hg). these treatment are basic treatment of acute pulmonary edema. but in control group we administrate basic treatment of acute pulmonary edema and not infused Mgso4. as we administrate 100 cc normal saline in 20 min for placebo instead Mgso4

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Rasoul and Sina hospital

Full name of responsible person

Seyyed Mahdi Zia Ziabari M.D

Street address

City

Rssht

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Tehran university medical science

Full name of responsible person

Dr. Akbar Fotohi

Street address

Teran Rasol hospital

City

Tehran

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Tehran university 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

Tehran university medical science

Full name of responsible person

Dr.Seyyed Mahdi Zia Ziabari M.D

Position

Emergency medicine assistant

Other areas of specialty/work

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Tehran university medical science

Full name of responsible person

Dr.Peyman Hafezi Moghadam

Position

Professor assistant of emergency medicine

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Tehran university medical science

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Position

Emergency medicine assistant

Other areas of specialty/work**Street 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