

Clinical Trial Protocol

Iranian Registry of Clinical Trials

04 Jun 2026

The Effect of Oral Zinc on Reduction of Mortality in Newborns with Neonatal Sepsis

Protocol summary

Study aim

The Effect of Oral Zinc on Morbidity and Mortality in Neonatal Sepsis

Design

. Two arm parallel group randomised double blinded trial

Settings and conduct

The aim of this study was to evaluate the effect of oral zinc sulfate on mortality and morbidity in neonatal sepsis admitted to NICU of Shahrekord Hajar Hospital during 2016-2017. Parents of neonates and data analyzer and committee on safety and supervision of data were kept unaware of the groups their neonate was allocate to.

Participants/Inclusion and exclusion criteria

neonates with sepsis are enrolled and neonates with congenital anomaly ,GA<32 weeks, NEC excluded from trial

Intervention groups

The first group received the oral zinc sulfate for 10 days, together with the antibiotic therapy, and the second group (the control group) received the classic treatment of sepsis.

Main outcome variables

mortality morbidity Hospitalization period

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20180915041040N2**

Registration date: **2019-08-16, 1398/05/25**

Registration timing: **retrospective**

Last update: **2019-08-16, 1398/05/25**

Update count: **0**

Registration date

2019-08-16, 1398/05/25

Registrant information

Name

Roya Choopani

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 38 3222 0016

Email address

choopani.r@skums.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2016-06-10, 1395/03/21

Expected recruitment end date

2017-10-02, 1396/07/10

Actual recruitment start date

2016-06-12, 1395/03/23

Actual recruitment end date

2017-08-23, 1396/06/01

Trial completion date

2017-11-12, 1396/08/21

Scientific title

The Effect of Oral Zinc on Reduction of Mortality in Newborns with Neonatal Sepsis

Public title

The Effect of Oral Zinc on Neonatal Sepsis

Purpose

Supportive

Inclusion/Exclusion criteria

Inclusion criteria:

neonate less than 28 day old neonate with sepsis

Exclusion criteria:

parental dissatisfaction threatening congenital anomalies asphyxia(resuscitation at birth more than 15 minutes or 10-minute Apgar score less than 5) Gestational age less than 32 weeks neonatal necrotizing enterocolitis

Age

From **1 day** old to **28 days** old

Gender

Both

Phase

3

Groups that have been masked

- Participant
- Care provider
- Outcome assessor
- Data analyser
- Data and Safety Monitoring Board

Sample size

Target sample size: **60**

Actual sample size reached: **60**

Randomization (investigator's opinion)

Not randomized

Randomization description

Blinding (investigator's opinion)

Double blinded

Blinding description

Parents of neonates and data analyzer and committee on safety and supervision of data were kept unaware of the groups their neonate was allocate to.

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Shahrekord University of Medical Sciences

Street address

Hajar hospital, parastar street

City

Shahrekord

Province

Chahar-Mahal-va-Bakhtiari

Postal code

8815874119

Approval date

2017-05-29, 1396/03/08

Ethics committee reference number

IR.SKUMS.REC.1396.49

Health conditions studied

1

Description of health condition studied

neonatal sepsis

ICD-10 code

ICD-10 code description

2

Description of health condition studied

neonatal sepsis

ICD-10 code

ICD-10 code description

Primary outcomes

1

Description

Mortality

Timepoint

Neonatal period

Method of measurement

Number

Secondary outcomes

1

Description

Hospitalization period

Timepoint

Neonatal period

Method of measurement

Day

Intervention groups

1

Description

Intervention group:

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Hajar Hospital

Full name of responsible person

Roya Choupani

Street address

Hajar Hospital ,Parastar Street

City

Shahrekord

Province

Chahar-Mahal-va-Bakhtiari

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity
Shahre-kord University of Medical Sciences
Full name of responsible person
Dr.Kamal Solati
Street address
Kashani Street,
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vcrt@skums.ac.ir
Web page address
<https://skums.ac.ir>

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Shahre-kord 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
Shahre-kord University of Medical Sciences
Full name of responsible person
Roya.Choopani
Position
Associate Professor
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Subspecialist
Other areas of specialty/work
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Person responsible for updating data

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

No - There is not a plan to make this available

Justification/reason for indecision/not sharing IPD

There are no information

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

No - There is not 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