

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

19 Jun 2026

### Assessing the effect of probiotics on intestinal feeding tolerance in preterm infants hospitalized in 17 shahrivar hospital of Rasht

#### Protocol summary

##### Summary

Objectives :To assess the effect of probiotics on intestinal feeding tolerance in preterm infants hospitalized in 17 Shahrivar hospital of Rasht, Design: A double blind controlled clinical trial, Participants including major eligibility criteria: 68 preterm infants with gestational age less than 36 weeks and 1000 - 2500 gram birth weight. Neonates with congenital malformations will be excluded from the study, Intervention: Eligible neonates will be assessed in two groups of treatment (oral probiotics) and control (normal saline), main outcome measures (variables): Mean duration of reaching full enteral feeding will be compared in two groups.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT201706129075N2**  
Registration date: **2017-09-11, 1396/06/20**  
Registration timing: **registered\_while\_recruiting**

Last update:

Update count: **0**

##### Registration date

2017-09-11, 1396/06/20

##### Registrant information

##### Name

Seyyedeh Zohreh Jalali

##### Name of organization / entity

Guilan University of Medical Science

##### Country

Iran (Islamic Republic of)

##### Phone

+98 13 1324 5126

##### Email address

z\_jalali@gums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

The Vice Chancellor of Research of Guilan University of Medical Sciences

##### Expected recruitment start date

2017-08-01, 1396/05/10

##### Expected recruitment end date

2018-04-01, 1397/01/12

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Assessing the effect of probiotics on intestinal feeding tolerance in preterm infants hospitalized in 17 shahrivar hospital of Rasht

##### Public title

Assessing the effect of probiotics on intestinal feeding tolerance in preterm infants hospitalized in 17 shahrivar hospital of Rasht

##### Purpose

Supportive

##### Inclusion/Exclusion criteria

Main inclusion criteria: Preterm infants with gestational age less than 36 weeks; Birth weight between 1000 and 2500 gram; Age less than or equal to two weeks with enteral feeding. Main exclusion criteria: Neonates with congenital malformations (congenital heart diseases; gastrointestinal obstruction; omphalocele; gastroschisis); congenital metabolism errors; Grade two and three of asphyxia; newborns of addicted mothers; age more than two weeks.

##### Age

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

##### Gender

Both

## Phase

2-3

## Groups that have been masked

No information

## Sample size

Target sample size: 68

## 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 Guilan University of Medical Sciences

##### Street address

Gaz Square

##### City

Rasht

##### Postal code

##### Approval date

2017-04-29, 1396/02/09

##### Ethics committee reference number

IR.GUMS.REC.1396.40

## Health conditions studied

### 1

#### Description of health condition studied

preterm birth

#### ICD-10 code

P07.3

#### ICD-10 code description

Other preterm infants

## Primary outcomes

### 1

#### Description

Mean duration of reaching full enteral feeding

#### Timepoint

at the discharge

#### Method of measurement

day

## Secondary outcomes

empty

## Intervention groups

### 1

#### Description

In the intervention group, administering oral probiotics with a dose of 1 drop/ kg diluted with up to 0.5 cc saline solution

#### Category

Rehabilitation

### 2

#### Description

In the control group, 0.5 cc of the normal saline solution will be administered every 12 hours and will be continued to reach 120 cc / kg

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

17 Shahrivar Hospital

##### Full name of responsible person

Dr Seyyedeh Zohre Jalali

##### Street address

Siadati Street

##### City

Rasht

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

The Vice Chancellor of Resrach of Guilan niversity of Medical Sciences

##### Full name of responsible person

Dr Shadman Nemati

##### Street address

Gaz Square

##### City

Rasht

#### Grant name

#### Grant code / Reference number

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

Yes

#### Title of funding source

The Vice Chancellor of Resrach of Guilan niversity 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**

Guilan University of Medical Sciences

**Full name of responsible person**

Dr Seyyedeh Zohre Jalali

**Position**

Assistant Professor

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Assistant Professor

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Guilan University of Medical Sciences

**Full name of responsible person**

Dr Seyyedeh Zohreh Jalali

**Position**

Assistant Professor

**Other areas of specialty/work****Street address**

Shahid Siadati Street

**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*