

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

06 Jul 2026

### A clinical trial comparing the effects of two different brands of ibuprofen (Pedia and Ibuprofen) on ductus arteriosus closure in premature infants with patent ductus arteriosus

#### Protocol summary

##### Study aim

Comparing the effects of two different brands of ibuprofen (Pedia and Ibuprofen) on ductus arteriosus closure in premature infants with patent ductus arteriosus

##### Design

Two-arm paralleled phase 3 trial without blinding and randomization.

##### Settings and conduct

A non-blinded study will be conducted in the pediatric department of Valiasr Hospital in Birjand. One group will receive Pedia, the French brand of ibuprofen produced by Orphan pharmaceutical company, while the other group will receive the Iranian form of ibuprofen, which is a generic version produced by Caspian Tamin Pharmaceutical Company. Both groups will receive intravenous injections of the medicine. The first day's dose will be 10 mg/kg, followed by 5 mg/kg after 24 hours, and another 5 mg/kg 24 hours later. The study will involve 160 patients who will be divided into two groups, with the first 80 receiving intervention from Group 1 and the subsequent 80 receiving intervention from Group 2.

##### Participants/Inclusion and exclusion criteria

Major inclusion criteria: Preterm newborn and size of patent ductus arteriosus that responds to drug therapy. Major exclusion criteria: the presence of underlying heart diseases, major congenital anomalies, and life-threatening sepsis.

##### Intervention groups

Intervention group 1 (French Ibuprofen): Ibuprofen from the French pharmaceutical company ORPHAN, Pedia brand, will be injected intravenously at doses of 10 mg/kg on the first day, 5 mg/kg 24 hours later, and 5 mg/kg 24 hours later (the last dose). Intervention group 2 (Iranian Ibuprofen): Ibuprofen from the Iranian Caspian Tamin Pharmaceutical Company, under the Ibuprofen generic name, will be injected intravenously at doses of

10 mg/kg on the first day, 5 mg/kg 24 hours later, and 5 mg/kg 24 hours later (the last dose).

##### Main outcome variables

Ductus arteriosus closure

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20140519017756N48**

Registration date: **2023-05-15, 1402/02/25**

Registration timing: **prospective**

Last update: **2023-05-15, 1402/02/25**

Update count: **0**

##### Registration date

2023-05-15, 1402/02/25

##### Registrant information

##### Name

Mohammad Bagher Roozgar

##### Name of organization / entity

Birjand University of Medical Sciences

##### Country

Iran (Islamic Republic of)

##### Phone

+98 56 3239 5680

##### Email address

mbroozgar@bums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2023-06-05, 1402/03/15

##### Expected recruitment end date

2024-06-04, 1403/03/15

**Actual recruitment start date**

empty

**Actual recruitment end date**

empty

**Trial completion date**

empty

**Scientific title**

A clinical trial comparing the effects of two different brands of ibuprofen (Pedeia and Ibuprofen) on ductus arteriosus closure in premature infants with patent ductus arteriosus

**Public title**

Effects of two different brands of ibuprofen on ductus arteriosus closure

**Purpose**

Treatment

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

Preterm newborn A size of patent ductus arteriosus that responds to drug therapy Gestational age of 32 weeks or more Birth weight 1500 grams and less Age between 3 and 15 days

**Exclusion criteria:**

Presence of underlying heart diseases Major congenital anomalies Life-threatening sepsis Renal failure Pulmonary hemorrhage Thrombocytopenia below 60,000/mm<sup>3</sup>

**Age**

From **3 days** old to **15 days** old

**Gender**

Both

**Phase**

3

**Groups that have been masked**

*No information*

**Sample size**

Target sample size: **160**

**Randomization (investigator's opinion)**

Not randomized

**Randomization description****Blinding (investigator's opinion)**

Not blinded

**Blinding description****Placebo**

Not used

**Assignment**

Parallel

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

empty

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

Ethics Committee of Birjand University of Medical

Sciences

**Street address**

Birjand University of Medical Sciences, Ayatollah Ghaffari Street

**City**

Birjand

**Province**

South Khorasan

**Postal code**

9717853577

**Approval date**

2019-03-02, 1397/12/11

**Ethics committee reference number**

lr.bums.REC.1397.325

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

Patent ductus arteriosus

**ICD-10 code**

Q25.0

**ICD-10 code description**

Patent ductus arteriosus

**Primary outcomes****1****Description**

Ductus arteriosus closure

**Timepoint**

Prior to treatment initiation and three days post-treatment completion

**Method of measurement**

Echocardiography

**Secondary outcomes**

empty

**Intervention groups****1****Description**

Intervention group 1 (French Ibuprofen/Pedeia): Ibuprofen from the French pharmaceutical company ORPHAN, Pedeia brand, will be injected intravenously at doses of 10 mg/kg on the first day, 5 mg/kg 24 hours later, and 5 mg/kg 24 hours later (the last dose).

**Category**

Treatment - Drugs

**2****Description**

Intervention group 2 (Iranian Ibuprofen): Ibuprofen from the Iranian Caspian Tamin Pharmaceutical Company, under the Ibuprofen generic name, will be injected intravenously at doses of 10 mg/kg on the first day, 5

mg/kg 24 hours later, and 5 mg/kg 24 hours later (the last dose).

**Category**

Treatment - Drugs

**Recruitment centers**

**1**

**Recruitment center**

**Name of recruitment center**

Valiasr Hospital's Pediatrics Department

**Full name of responsible person**

Alireza Taghizadegan

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Ghaffari Ave.

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**Sponsors / Funding sources**

**1**

**Sponsor**

**Name of organization / entity**

Vice-chancellery for Research, Birjand University of Medical Sciences

**Full name of responsible person**

Dr. Mohammadreza Miri

**Street address**

Vice-chancellory for Research, Birjand University of Medical Sciences, Ayatollah Ghafari Street,

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Miri\_moh2516@yahoo.com

**Grant name**

**Grant code / Reference number**

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

Yes

**Title of funding source**

Vice-chancellery for Research, Birjand 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**

Birjand University of Medical Sciences

**Full name of responsible person**

Alireza Taghizadegan

**Position**

General Practitioner

**Latest degree**

Medical doctor

**Other areas of specialty/work**

General Practitioner

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**Person responsible for scientific inquiries**

**Contact**

**Name of organization / entity**

Birjand University of Medical Sciences

**Full name of responsible person**

Dr. Forud Salehi

**Position**

Pediatric cardiologist

**Latest degree**

Subspecialist

**Other areas of specialty/work**

Cardiology

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## Person responsible for updating data

### Contact

**Name of organization / entity**

Birjand University of Medical Sciences

**Full name of responsible person**

Mohammad Bagher Roozgar

**Position**

Translator

**Latest degree**

Master

**Other areas of specialty/work**

Others

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## Sharing plan

### Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

**Study Protocol**

Yes - There is a plan to make this available

**Statistical Analysis Plan**

Yes - There is a plan to make this available

**Informed Consent Form**

Yes - There is a plan to make this available

**Clinical Study Report**

Yes - There is a plan to make this available

**Analytic Code**

Yes - There is a plan to make this available

**Data Dictionary**

Not applicable

**Title and more details about the data/document**

Deidentified Individual Participant Data Set and other information related to research methodology

**When the data will become available and for how long**

After the paper extracted from the project is published and for 6 months ever since

**To whom data/document is available**

Researchers

**Under which criteria data/document could be used**

Research purposes

**From where data/document is obtainable**

Corresponding author

**What processes are involved for a request to access data/document**

Personal email to the corresponding author of the extracted article

**Comments**