

Clinical Trial Protocol

Iranian Registry of Clinical Trials

01 Jun 2026

Evaluation of short term protective effect of aggressive hydration therapy with lactated ringer's solution on pancreatitis after endoscopic retrograde cholangiopancreatography

Protocol summary

Summary

150 patient candidate for endoscopic retrograde cholangiopancreatography in 5 months period with inclusion criteria(Choledochobilliary Stone; Dilatation and Obstruction of Choledochobilliary) and without exclusion criteria(Acute Cholangitis; Pregnancy; End Organ Failure; Age above 70 years) divided to two groups, case and control randomizely. That's obvious for researcher exclusively. For case group aggressive hydration and for control group routine hydration therapy will be done. Both groups discontinue intravenous fluids when they are able to eat a normal diet. The aim of this study is evaluation of aggressive hydration therapy in prevention of post endoscopic retrograde cholangiopancreatography pancreatitis as primary end point.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201410161213N3**

Registration date: **2014-10-19, 1393/07/27**

Registration timing: **prospective**

Last update:

Update count: **0**

Registration date

2014-10-19, 1393/07/27

Registrant information

Name

Manouchehr Khoshbaten

Name of organization / entity

Tabriz University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 41 1334 7554

Email address

khoshbatenm@tbzmed.ac.ir

Recruitment status

Recruitment complete

Funding source

Vice-chancellor for research -Tabriz University of Medical Science

Expected recruitment start date

2014-11-06, 1393/08/15

Expected recruitment end date

2015-04-20, 1394/01/31

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Evaluation of short term protective effect of aggressive hydration therapy with lactated ringer's solution on pancreatitis after endoscopic retrograde cholangiopancreatography

Public title

Effect of aggressive hydration therapy in prevention of complication of endoscopic therapeutic intervention on biliary duct.

Purpose

Prevention

Inclusion/Exclusion criteria

(Inclusion criteria: Indication of endoscopic retrograde cholangiopancreaticography; consent of enrollment in study and Exclusion criteria: Acute cholangitis; sepsis; pregnancy; age above 70 years; chronic pancreatitis; acute pancreatitis due to biliary stone; risk of fluid

overload; peripheral edema; pulmonary edema; electrolyte disturbance for example Na over 150 or below 130; history of endoscopic retrograde cholangiopancreatography performance with sphinctrotomy.)

Age

From **18 years** old to **70 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **150**

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

Tabriz University of Medical Sciences , affiliated to the Ministry of Health, Treatment and Medical

Street address

Central bulding of medical science university, St. Goltasht, St. Azadi ,Tabriz

City

Tabriz

Postal code

Approval date

2014-10-12, 1393/07/20

Ethics committee reference number

93104

Health conditions studied

1

Description of health condition studied

Pancreatitis

ICD-10 code

K85

ICD-10 code description

Acute pancreatitis

Primary outcomes

1

Description

Post endoscopicretrograde cholangiopancreatography pancreatitis

Timepoint

2,8 and 24 hours after intervation

Method of measurement

Questionnaire

Secondary outcomes

1

Description

Hyper amylasemia

Timepoint

2, 8 and 24 after intervation

Method of measurement

U/L

2

Description

Pain

Timepoint

2, 8 and 24 after intervation

Method of measurement

Visual analouge pain scale

Intervention groups

1

Description

In case group lactated ringer serum administer with dose of 20 cc per kilogram of body mass and continue with 3 cc per kilogram of body mass until 8 hours.

Category

Treatment - Drugs

2

Description

In control group lactated ringer serum administer with dose of 1.5 cc per kilogram of body mass until 8 hours.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Tabriz University of Medical Sciences

Full name of responsible person

Ali reza shayegan nezhad

Street address

Endoscopy ward, first floor, Imam Reza Hospital, St.

Golgasht, St.Azadi

City

Tabriz

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Vice-chancellor for research -Tabriz University of Medical Science

Full name of responsible person

Dr.Mohammad Reza Rashidi

Street address

Third floor, Central building No.2, Tabriz University of Medical Sciences ,St.Golgasht

City

Tabriz

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Vice-chancellor for research -Tabriz University of 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

Tabriz University of Medical Sciences

Full name of responsible person

Ali Reza Shaygan nezhad

Position

Gastroentrology and hepatology assistant

Other areas of specialty/work

Street address

Endoscopy ward, first floor, Imam Reza Hospital, St. Golgasht, St.Azadi, Tabriz

City

Endoscopy ward, first floor, Imam Reza Hospital, St. Golgasht, St.Azadi, Tabriz

Postal code

Phone

+98 41 3334 7554

Fax

+98 41 3334 7554

Email

ali.shayegan@yahoo.com

Web page address

Person responsible for scientific inquiries

Contact

Name of organization / entity

Tabriz University of Medical Sciences

Full name of responsible person

Ali Reza Shaygan Nezhad

Position

Gastroentrology and hepatology assistant

Other areas of specialty/work

Street address

Endoscopy ward, first floor, Imam Reza Hospital, St. Golgasht, St.Azadi, Tabriz

City

Tabriz

Postal code

Phone

+98 41 3334 7554

Fax

+98 41 3334 7554

Email

Ali.shayegan@yahoo.com

Web page address

Person responsible for updating data

Contact

Name of organization / entity

Tabriz University of Medical Sciences

Full name of responsible person

Ali Reza Shaygan Nezhad

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Gastroentrology and hepatology assistant

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Street address

Endoscopy ward, first floor, Imam Reza Hospital, St. Golgasht, St.Azadi, Tabriz

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ali.shayegan@yahoo.com

Web page 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