

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

28 May 2026

### The effect of personalized parenteral nutrition on Intensive care unit (ICU) patients

#### Protocol summary

##### Study aim

Determination of the effect of personalized parenteral nutrition on ICU patients

##### Design

Randomized clinical trial, 2-blind trial, 60 ICU patients candidate for parenteral nutrition, Intervention and control groups through randomization, 30 patients as intervention group & 30 patients as control group.

##### Settings and conduct

We attend at ICU of Razi hospital in Rasht and complete consent form, information will be required using questionnaires: Medical history, Anthropometric measurement, Dietary intake, Biochemical indices (at baseline and after 14 days). This is a Double-blind trial study, which intervention and control groups are blinded to intervention assignment and nurses who assess and collect data on outcome, are blinded to groups' assignment.

##### Participants/Inclusion and exclusion criteria

Study will be done in the Rasht Razi hospital. 60 ICU patients who are candidate for parenteral nutrition, will be selected and enter to the study in the intervention and control groups through simple randomized selection. Inclusion criteria: Informed consent, need to parenteral nutrition at least 24-h of hospitalization and exclusion criteria: no tendency, no parenteral nutrition indication for more than 1 week, death, not having severe or middle malnutrition.

##### Intervention groups

We avoid feeding the lipid to intervention group in the first of hospitalization. Required protein will be calculated upon patient's status and determined by required percent of amino acids & Dextrose. We use daily relevant vials for providing vitamins & minerals. Control group daily receive low calorie of intra venous diet (800-1000 calories) upon hospital's routine. We usually administer Dextrose every day, providing 70% and administer Aminoven or Intralipid every other day, providing 30% of patient's calorie. All of measurements repeated after one

week.

##### Main outcome variables

Urea, Creatinine, CRP, ICU hospitalization, Survival rate

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20151226025699N4**

Registration date: **2020-06-03, 1399/03/14**

Registration timing: **prospective**

Last update: **2020-06-03, 1399/03/14**

Update count: **0**

##### Registration date

2020-06-03, 1399/03/14

##### Registrant information

##### Name

Saeid Doaei

##### Name of organization / entity

National Nutrition and Food Technology Research Institute

##### Country

Iran (Islamic Republic of)

##### Phone

+98 21 6643 6744

##### Email address

sdoaei@sbmu.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2020-06-04, 1399/03/15

##### Expected recruitment end date

2020-06-18, 1399/03/29

##### Actual recruitment start date

empty  
**Actual recruitment end date**  
empty  
**Trial completion date**  
empty

**Scientific title**  
The effect of personalized parenteral nutrition on Intensive care unit (ICU) patients

**Public title**  
The effect of parenteral nutrition on ICU patients

**Purpose**  
Supportive

**Inclusion/Exclusion criteria**  
**Inclusion criteria:**  
Consent to participation Need to parenteral nutrition At least 24-h of hospitalization

**Exclusion criteria:**  
Having no indication for parenteral nutrition for more than 1 week Unstable hemodynamic conditions

**Age**  
From **18 years** old

**Gender**  
Both

**Phase**  
N/A

**Groups that have been masked**

- Participant
- Data analyser

**Sample size**  
Target sample size: **60**

**Randomization (investigator's opinion)**  
Randomized

**Randomization description**  
Patients of ICU are assigned to the intervention and control groups through simple individualized randomization by online software (randomizer.org).

**Blinding (investigator's opinion)**  
Double blinded

**Blinding description**  
The subjects (both the case and control groups) are blinded to intervention assignment. Also, nurses who assess and collect data on outcome, are blinded to groups' assignment.

**Placebo**  
Not used

**Assignment**  
Parallel

**Other design features**

**Secondary Ids**  
empty

**Ethics committees**

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

Cancer research center of Shahid beheshti University of Medical Sciences

**Street address**  
Tehran, Tajrish square, Tajrish martyrs hospital,Cancer research center

**City**  
Tajrish

**Province**  
Tehran

**Postal code**  
1985717413

**Approval date**  
2019-11-12, 1398/08/21

**Ethics committee reference number**  
IR.SBMU.CRC.REC.1398.015

**Health conditions studied**

1  
**Description of health condition studied**  
Gastrointestinal hemorrhage

**ICD-10 code**  
K92.2

**ICD-10 code description**  
Gastrointestinal hemorrhage, unspecified

**Primary outcomes**

1  
**Description**  
Urea  
**Timepoint**  
Baseline, 14-day after intervention  
**Method of measurement**  
using standard kits and the researchers gather these information from the lab test section of the ICU sheets.

2  
**Description**  
Creatinine  
**Timepoint**  
Baseline, 14-day after intervention  
**Method of measurement**  
using standard kits and the researchers gather these information from the lab test section of the ICU sheets.

3  
**Description**  
C reactive protein (CRP)  
**Timepoint**  
Baseline, 14-day after intervention  
**Method of measurement**  
using standard kits and the researchers gather these information from the lab test section of the ICU sheets.

4  
**Description**

Serum albumin

### **Timepoint**

Baseline, 14-day after intervention

### **Method of measurement**

using standard kits and the researchers gather these information from the lab test section of the ICU sheets.

## **Secondary outcomes**

### 1

#### **Description**

ICU hospitalization

#### **Timepoint**

Baseline, 14-day after intervention

#### **Method of measurement**

Patients documents

### 2

#### **Description**

Survival rate

#### **Timepoint**

During 14-day intervention

#### **Method of measurement**

Patients documents

### 3

#### **Description**

Weight

#### **Timepoint**

During 14-day intervention

#### **Method of measurement**

This anthropometric indicator will be gathered from the patients records.

### 4

#### **Description**

Body mass index (BMI)

#### **Timepoint**

During 14-day intervention

#### **Method of measurement**

This anthropometric indicator will be gathered from the patients records.

### 5

#### **Description**

Malnutrition status

#### **Timepoint**

During 14-day intervention

#### **Method of measurement**

It will be evaluated by weight & BMI criteria and by the level of serum albumin. We also use the Nutritional Risk Screening (NRS 2002) criterion for more evaluation.

## **Intervention groups**

### 1

#### **Description**

After the gain of real weight and evaluation of disease status and chemistry results, the amount of required calorie will be estimated according to the ASPEN guidelines and then, the appropriate diet will be designed. We avoid to feeding the lipid in the first of hospitalization. The required protein will be calculated according to the patient's status and determined by the required percent of amino acids and Dextrose. We also will use the daily relevant vials for providing the vitamins and required minerals. Finally, we will insert the diet with relevant advises in to the conference form and deliver to the relevant specialist doctor for final acceptance.

#### **Category**

Treatment - Drugs

### 2

#### **Description**

Control group will daily receive low calorie of parenteral diet, including 800 to 1000 calories according to the hospital's routine. We usually administer Dextrose every day, which provide 70% of patient's calorie and administer Aminoven or Intralipid every other day, which provide 30% of patient's calorie. All of measurements repeated after one week.

#### **Category**

Placebo

## **Recruitment centers**

### 1

#### **Recruitment center**

##### **Name of recruitment center**

Razi hospital

##### **Full name of responsible person**

Siamak Rimaz

##### **Street address**

RaziSardare Jangal Boulevard

##### **City**

Tehran

##### **Province**

Guilan

##### **Postal code**

41448

##### **Phone**

+98 13 3355 0028

##### **Email**

sdoaee@yahoo.com

## **Sponsors / Funding sources**

### 1

#### **Sponsor**

##### **Name of organization / entity**

Shahid Beheshti University of Medical Sciences

##### **Full name of responsible person**

Pr. Mohammad Esmaeil Akbari

**Street address**

Chamran Highway Yeman street-Arabi street

**City**

Tehran

**Province**

Tehran

**Postal code**

1985717413

**Phone**

+98 21 23871

**Email**

info@sbmu.ac.ir

**Grant name****Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

**Title of funding source**

Shahid Beheshti University of Medical Sciences

**Proportion provided by this source**

50

**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

**2****Sponsor****Name of organization / entity**

Rasht University of Medical Sciences

**Full name of responsible person**

Dr. Arsalan salari

**Street address**

Parastar Street

**City**

Rasht

**Province**

Guilan

**Postal code**

41887-94755

**Phone**

+98 13 3334 6489

**Email**

salari@gums.ac.ir

**Grant name****Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

No

**Title of funding source**

Rasht university of medical sciences

**Proportion provided by this source**

50

**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**

Shahid Beheshti University of Medical Sciences

**Full name of responsible person**

Pr. Mohammad Esmaeil Akbari

**Position**

Super-specialist of Endocrine and Cancer surgery

**Latest degree**

Subspecialist

**Other areas of specialty/work**

Others

**Street address**

Tehran, Tajrish square, Tajrish martyrs hospital, Cancer research center

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**Postal code**

1985717413

**Phone**

021 22748001-2

**Email**

me-akbari@sbmu.ac.ir

**Person responsible for scientific inquiries****Contact****Name of organization / entity**

Shahid Beheshti University of Medical Sciences

**Full name of responsible person**

Dr. Saeid Doaei

**Position**

Tehran shahid beheshti university of Medical Sciences

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Nutrition

**Street address**

Sardare Jangal Boulevard

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**Province**

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sdoaei@sbmu.ac.ir

## Person responsible for updating data

sdoaei@sbmu.ac.ir

### Contact

**Name of organization / entity**

Shahid Beheshti University of Medical Sciences

**Full name of responsible person**

Dr. Saeid Doaei

**Position**

Ph.D

**Latest degree**

Ph.D.

**Other areas of specialty/work**

Nutrition

**Street address**

Shahid Beheshti University of Medical Sciences,  
Tehran, IRAN.

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009821

**Phone**

+98 21 6643 6744

**Email**

## Sharing plan

**Deidentified Individual Participant Data Set (IPD)**

Undecided - It is not yet known if there will be a plan to make this available

**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**

Undecided - It is not yet known if there will be 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