

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

The comparative study of the effectiveness of teaching methods of workshop and multimedia on knowledge, attitude and performance of nurses about blood

Protocol summary

Summary

The purpose of this study is to compare the effect of two methods, multimedia and workshops, on knowledge, attitude and performance of nurses in the case of blood transfusion which is a randomized clinical trial with a control group. The members of this research are 111 people which are based on the determined sample size of nurses working in Rasoul Akram, Firoozgar and Firoozabadi teaching Hospitals. In case of lack of cooperation, the participant will be excluded from the study. To participate to the study 37 people will be selected from each hospital which two hospitals would be put in the intervention group and one hospital in the control group. At first the questionnaire of demographic, characteristics, knowledge, attitudes and performance will be completed as a self-report (pretest) then, it would be presented in the two hospitals which type of training is determined between the two types of training (workshops and multimedia). Two weeks after the intervention, in order to assess the primary outcome, self-reported questionnaires will be complete by nurses, as the previous, in the condition of transfusion, then will collect and evaluate. Also, the flash containing educational content and two educational posters would be offer to the control group as to comply with ethical issues then questionnaires would be analyzed by descriptive and interventional statistical tests.

General information

Acronym

Not

IRCT registration information

IRCT registration number: **IRCT2016081625333N1**

Registration date: **2016-12-10, 1395/09/20**

Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2016-12-10, 1395/09/20

Registrant information

Name

Maryam Janamiri

Name of organization / entity

Iran School of Nursing and Midwifery, Iran University of Medical Science

Country

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+98 21 8820 1880

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Recruitment status

Recruitment complete

Funding source

Research Assistance of Iran University of Medical Sciences-The International Pardis

Expected recruitment start date

2015-11-18, 1394/08/27

Expected recruitment end date

2016-08-16, 1395/05/26

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The comparative study of the effectiveness of teaching methods of workshop and multimedia on knowledge, attitude and performance of nurses about blood

Public title

Effectiveness of teaching methods of workshop and multimedia on knowledge, attitude and performance of nurses about blood transfusion

Purpose

Other

Inclusion/Exclusion criteria

Inclusion criteria: The nurse would be in direct contact with patients; also at least would have 6 months of experience in the emergency, surgery or oncology parts; additionally, employees should be excepted in Official, Contractual or project personnel. Exclusion criteria: During time of the study, In the case of lack of cooperation or nurse's displacement of the emergency, surgery or oncology parts nurses will be excluded from the study.

Age

No age limit

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **111**

Randomization (investigator's opinion)

N/A

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

Not blinded

Blinding description**Placebo**

Not used

Assignment

Parallel

Other design features

Not

Secondary Ids

empty

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

Ethics Committee of Iran University of Medical Sciences

Street address

Iran University of Medical Sciences, Between The Highway Martyr Chamran and Martyr Sheikh Fazlullah Nuri, Martyr Hemmat Highway ,Tehran, Iran.

City

Tehran

Postal code

1997613883

Approval date

2015-11-18, 1394/08/27

Ethics committee reference number

IR.IUMS.REC.1394 .9213655201

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

Nurse education

ICD-10 code

N/A

ICD-10 code description

N/A

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

knowledge, attitude and performance of nurses

Timepoint

Pre test and Two Week's after intervention

Method of measurement

questionnaire

Secondary outcomes

empty

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

In the training workshops, including a four-hour session, three groups separately will be trained by method of lecture, group discussion, questions and answers, using the slide and two educational posters about educational purposes, the necessity of education, history of blood transfusions, procedures and actions required before blood transfusion which includes (verify the patient's identity, provide the conditions before taking delivery of products from blood bank, check apparent characteristics of submitted products, the process of blood transition) and the Instruction of dealing with the most common Acute complications related to blood transfusion which includes (fever, hives, dyspnea, hypotension). During the meeting, researcher will encourage nurses to have group discussion, questions and answers related to ambiguities. At end of the session, researcher and participants will summarize the subject. The nurses also will be asked to study the two posters during these two weeks. Also The Telephone number of research will be given to caregivers for solving the problems and answer to questions. In the training session, the researcher after introduction of her will divide the participants into groups of eight to ten people and then will explain the training program, the aim of the intervention and the importance of their cooperation. In addition to that the researcher will give some explanation about the process of blood transfusion and the important role of nurses in this process.

Category

Other

2

Description

In the method of teaching with multimedia, educational content which is saved on the flash will be offer to the participants.

Category

Other

3

Description

For the control group, training will not be given by researcher. In future they will receive routine training which includes saved educational content on flash and the two educational posters.

Category

Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Rasoole Akram Hospital

Full name of responsible person

Forough Rafii

Street address

Iran School of Nursing and Midwifery, Yasami St., Vali Asr Avenue.

City

Tehran

2

Recruitment center

Name of recruitment center

Firoozgar Hospital

Full name of responsible person

Forough Rafii

Street address

Iran School of Nursing and Midwifery, Yasami St., Vali Asr Avenue.

City

Tehran

3

Recruitment center

Name of recruitment center

Firoozabadi Hospital

Full name of responsible person

Forough Rafii

Street address

Iran School of Nursing and Midwifery, Yasami St., Vali Asr Avenue.

City

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Research Assistance of Iran University of Medical Sciences

Full name of responsible person

Syed Ali Javad Mousavi

Street address

Iran University of Medical Sciences, Between The Highway Martyr Chamran and Martyr Sheikh Fazlullah Nuri, Martyr Hemmat Highway ,Tehran, Iran.

City

Tehran

Grant name

Grant code / Reference number

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

Yes

Title of funding source

Research Assistance of Iran University 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

Iran School of Nursing and Midwifery, Iran University of Medical Science

Full name of responsible person

Forough Rafii

Position

Professor

Other areas of specialty/work

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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