

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

Comparison the effect of applying “expiratory rib cage compression” and “manual percussion” before suction on respiratory parameters in patients under mechanical ventilation

Protocol summary

Study aim

Comparison the effect of applying “expiratory rib cage compression” and “manual percussion” before suction on respiratory parameters in patients under mechanical ventilation

Design

Randomized cross-over Clinical Trials, with A and B intervention groups will be performed on 46 patients. The letter A is written on 23 cards and the letter B on the other 23 cards. One of these two cards is randomly selected for each patient. Patients with card A enter Group A and patients with card B enter Group B.

Settings and conduct

The intensive care units of Kowsar Hospital will be used as a research environment. Both interventions will be performed by the researcher on the same day with an interval of 3 hours (to neutralize the previous intervention) and changes in study variables will be evaluated and recorded in the 4 interval before, 1,5, 25 minutes after endotracheal suctioning.

Participants/Inclusion and exclusion criteria

Inclusion criteria: intubated patients under mechanical ventilation for at least 48 h, not needing more than 2 times suction per shift, patients with stable hemodynamic parameters, reversible gastric secretion less than 100 ml before Intervention, aged 18 to 65 years Exclusion criteria: Respiratory distress, pneumothorax and Fractures of ribs, sternum and chest, chest burns and surgery, dysrhythmia and heart disease, Abscess and lung cyst.

Intervention groups

Intervention Group A: First, the Manual percussion and then the expiratory rib cage compression will be performed. Intervention Group B: First, the expiratory rib cage compression and then Manual percussion will be performed. Both interventions will be performed by the researcher in one day, observing a time interval of 3

hours (to neutralize the previous intervention).

Main outcome variables

End tidal CO₂ (ET-CO₂), arterial oxygen saturation, respiratory rate, pulmonary Compliance and airway resistance

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20191030045281N1**

Registration date: **2020-06-25, 1399/04/05**

Registration timing: **registered_while_recruiting**

Last update: **2020-06-25, 1399/04/05**

Update count: **0**

Registration date

2020-06-25, 1399/04/05

Registrant information

Name

Maryam Rasuli

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 87 3424 0418

Email address

maryam.rasuli@muk.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2020-05-21, 1399/03/01

Expected recruitment end date

2020-09-22, 1399/07/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison the effect of applying “expiratory rib cage compression” and “manual percussion” before suction on respiratory parameters in patients under mechanical ventilation

Public title

Comparison the effect of applying “expiratory rib cage compression” and “manual percussion” before suction on respiratory parameters

Purpose

Prevention

Inclusion/Exclusion criteria**Inclusion criteria:**

intubated patients Patients under mechanical ventilation with volumetric mode For at least 48 hours under mechanical ventilation not needing more than 2 times suction per shift patients with stable hemodynamic parameters reversible gastric secretion less than 100 ml before Intervention Age between 18 and 65 years

Exclusion criteria:

Respiratory distress pneumothorax and Fractures of the ribs, sternum and chest chest burns and surgery dysrhythmia and heart disease Abscess and lung cyst

Age

From **18 years** old to **65 years** old

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **46**

Randomization (investigator's opinion)

Randomized

Randomization description

Group A is a group that will be carried out first by manual percussion and then expiratory rib cage compression. Group B group that will be performed first by expiratory rib cage compression and then manual percussion . The letter A is written on 23 cards and the letter B on the other 23 cards. One of these two cards is randomly selected for each patient. Patients with card A enter Group A and patients with card B enter Group B.

Blinding (investigator's opinion)

Not blinded

Blinding description**Placebo**

Not used

Assignment

Crossover

Other design features**Secondary Ids**

empty

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

Ethics committee of Kurdistan University of Medical Sciences

Street address

Kurdistan University of Medical Sciences Campus, Pasdaran Ave, Sanandaj

City

Sanandaj

Province

Kurdistan

Postal code

13446-66177

Approval date

2020-03-14, 1398/12/24

Ethics committee reference number

IR.MUK.REC.1398.293

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

Patients under mechanical ventilation

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

end Tidal CO2

Timepoint

Exhaled CO2 is measured 4 times: before and minutes 1, 5 and 25 minutes after endotracheal suctioning.

Method of measurement

Using Capnography Probe

2**Description**

Oxygen saturation (SPO2)

Timepoint

Oxygen saturation (SPO2) is measured 4 times: before and minutes 1, 5 and 25 minutes after endotracheal suction

Method of measurement

Using a pulse oximeter device (via monitor view)

3**Description**

Pulmonary compliance

Timepoint

Pulmonary compliance is measured 4 times: before and minutes 1, 5 and 25 minutes after endotracheal suction

Method of measurement

Through viewing the ventilator screen

4

Description

Airway resistance

Timepoint

Airway resistance measured 4 times: before and minutes 1, 5 and 25 minutes after endotracheal suction

Method of measurement

Through viewing the ventilator screen

5

Description

Respiratory rate

Timepoint

Respiratory rate is measured 4 times: before and minutes 1, 5 and 25 minutes after endotracheal suction

Method of measurement

Total spontaneous breathing and ventilator-provided breathing per minute by viewing the ventilator screen

Secondary outcomes

empty

Intervention groups

1

Description

Group A is the group that will do the manual percussion first and then expiratory rib cage compression. In the manual percussion procedure, the hands are cupped and all fingers are folded and the wrist joint is opened and performed at speeds of 100 to 120 times per minute. In the expiratory rib cage compression, gradually compressed the end and lateral parts of patients' rib cage during expiration by his hands. This technique was bilaterally performed, and in the end of expiration, the compression was released from the patients' rib cage to let them have a free inspiration. Both interventions will be performed by the researcher in one day with a 3-hour interval (to neutralize the previous intervention), respectively. Respiratory parameters: increased end-expiratory CO₂, pulmonary compliance and hemoglobin saturation, and respiratory rate and airway resistance in patients undergoing mechanical ventilation in the ICU will be assessed before and 1, 5 and 25 minutes after endotracheal suctioning.

Category

Prevention

2

Description

Group B is the group that first will do the expiratory rib cage compression and then manual percussion before

the suction. In the manual percussion procedure, the hands are cupped and all fingers are folded and the wrist joint is opened and performed at speeds of 100 to 120 times per minute. In the expiratory rib cage compression, gradually compressed the end and lateral parts of patients' rib cage during expiration by his hands. This technique was bilaterally performed, and in the end of expiration, the compression was released from the patients' rib cage to let them have a free inspiration. Both interventions will be performed by the researcher in one day with a 3-hour interval (to neutralize the previous intervention), respectively. Respiratory parameters: increased end-expiratory CO₂, pulmonary compliance and hemoglobin saturation, and respiratory rate and airway resistance in patients undergoing mechanical ventilation in the ICU will be assessed before and 1, 5 and 25 minutes after endotracheal suctioning.

Category

Prevention

Recruitment centers

1

Recruitment center

Name of recruitment center

Kosar Medical Center of Sanandaj

Full name of responsible person

Maryam Rasuli

Street address

Kowsar hospital, Pasdaran Ave, Azadi Square

City

Sanandaj

Province

Kurdistan

Postal code

6617983476

Phone

+98 87 3361 1234

Email

kowsar@muk.ac.ir

Web page address

Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Sanandaj University of Medical Sciences

Full name of responsible person

Khaled Rahmani

Street address

Vice Chancellor for Research and Technology,
Kurdistan University of Medical Sciences, Pasdaran
Blvd., Sanandaj, Kurdistan

City

Sanandaj

Province

Kurdistan

Postal code

13446-66177

Phone

+98 87 3366 4653

Email

khaledrahmani111@gmail.com

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

Yes

Title of funding source

Sanandaj 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**

Sanandaj University of Medical Sciences

Full name of responsible person

Maryam Rasuli

Position

MSc Student of Critical Care Nursing

Latest degree

Bachelor

Other areas of specialty/work

Nursery

Street address

Narges 5 Suleiman Bag Street 37

City

Baneh

Province

Kurdistan

Postal code

66918-64496

Phone

+98 87 3424 0418

Email

maryam.rasuli@muk.ac.ir

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

Sanandaj University of Medical Sciences

Full name of responsible person

Dr. Mohammad Fathi

Position

Assistant Professor

Latest degree

Ph.D.

Other areas of specialty/work

Nursery

Street address

Kurdistan University of Medical Sciences Campus,
Pasdaran Ave, Sanandaj

City

Sanandaj

Province

Kurdistan

Postal code

66177-13446

Phone

+98 87 3366 4653

Fax

+98 87 3366 4654

Email

fathi_sanana@yahoo.com

Person responsible for updating data**Contact****Name of organization / entity**

Sanandaj University of Medical Sciences

Full name of responsible person

Maryam Rasuli

Position

Nursing Expert- MSc Student of Critical Care Nursing

Latest degree

Bachelor

Other areas of specialty/work

Nursery

Street address

No. 37, Narges Alley 5, Suleiman Bag Ave,

City

Baneh

Province

Kurdistan

Postal code

66918-64496

Phone

+98 87 3424 0418

Email

m.rbaneh61@gmail.com

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