

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

Comparison of the view of the larynx, the time spent for intubation and the success rate of tracheal intubation with a gladoscope and Sanyar video laryngoscope in adults patients with difficult airway during induction of anesthesia

Protocol summary

Study aim

Determining the time required for tracheal intubation and with Sanyar video laryngoscope success rate of intubation among patients with difficult airway

Design

A single-blind randomized clinical trial with control group, and parallel groups, on 70 patients. Computer-generated random table method was used for randomization

Settings and conduct

A single-blind clinical trial study is performed on patients with difficult airway who candidate for surgery , and need laryngoscopy and tracheal intubation at Sina Hospita Blinding is the person who collects the data and the analyst.

Participants/Inclusion and exclusion criteria

Inclusion criteria: All patients who are candidates for surgery; Ages 18-70 who need tracheal intubation and have difficult airway criteria. Non-inclusion criteria: Patients with a history of an easy documented tracheal intubation (first attempt success); history of unsuccessful intubation and failed ventilation mask; known unstable cervical spine injury; age less than 18 years; or patients who have an emergency surgery

Intervention groups

Intervention group: Patients who are candidates for surgery and need tracheal intubation in the operating room, after monitoring and injecting anesthetic drugs, laryngoscopy is performed with a Sanyar video laryngoscope. Control group: Laryngoscopy is performed with GlideScope laryngoscopy.

Main outcome variables

View of the larynx, duration of intubation, success rate of intubation in the first attempt, during laryngoscopy

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20130304012695N15**

Registration date: **2023-01-25, 1401/11/05**

Registration timing: **prospective**

Last update: **2023-01-25, 1401/11/05**

Update count: **0**

Registration date

2023-01-25, 1401/11/05

Registrant information

Name

mohammadreza khajavi

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 21 6312 1220

Email address

khajavim@tums.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2023-04-04, 1402/01/15

Expected recruitment end date

2024-06-19, 1403/03/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison of the view of the larynx, the time spent for intubation and the success rate of tracheal intubation with a gladoscope and Sanyar video laryngoscope in adults patients with difficult airway during induction of anesthesia

Public title

Evaluation of the speed and accuracy of laryngoscopy with the Sanyar video laryngoscope in patients with airway problems

Purpose

Other

Inclusion/Exclusion criteria

Inclusion criteria:

Patients aged 18-70 years old Candidates for elective surgeries need tracheal intubation Having difficult airway criteria

Exclusion criteria:

Patients with a history of easy documented tracheal intubation History of failed intubation and failed ventilation mask Known unstable cervical spine injury Patients having an emergency surgery.

Age

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

Gender

Both

Phase

N/A

Groups that have been masked

- Participant
- Investigator
- Data analyster

Sample size

Target sample size: **70**

Randomization (investigator's opinion)

Randomized

Randomization description

For the randomization of patients who meet the inclusion criteria, the method of four blocks including intervention and control groups will be used. The preparation of randomization sequences will be done using the Random Generator software and the created sequences will be given to a trained staff member of the operating room who is not a member of the research group. The researchers of this study will not be aware of the existing sequences and arrangement of the blocks. After the patient enters the operating room, the trained person removes the first sequence from the special box of this study and according to the predetermined protocol, if it is S, it will be transferred to the intervention group, and if it is G, it will be transferred to the control group.

Blinding (investigator's opinion)

Single blinded

Blinding description

Patients participating in the study are not aware of the type of laryngoscope for intubation. The name of the person collecting the research information is not on the list of researchers. The analyzer is not involved in the

research process.

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of the School of Medicine, Tehran University of Medical Sciences

Street address

Sina hospital, Hasan Abad Sq., Emam Khomeni st.

City

Tehran

Province

Tehran

Postal code

1136746911

Approval date

2023-01-24, 1401/11/04

Ethics committee reference number

IR.TUMS.SINAHOSPITAL.REC.1401.099

Health conditions studied

1

Description of health condition studied

intra tracheal intubation

ICD-10 code

T88.4XXA

ICD-10 code description

Failed or difficult intubation, initial encounter

Primary outcomes

1

Description

Glottic view

Timepoint

At the time of laryngoscopy

Method of measurement

According to Cormac criteria

2

Description

Duration of intubation

Timepoint

From the moment the laryngoscope enters the mouth to observe the expiratory wave of the capnograph

Method of measurement

With stopwatch and in seconds

3

Description

Successes rate in first try

Timepoint

During laryngoscopy and intubation

Method of measurement

Divide the number of successful intubation attempts by the total number of attempts

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: Intubation is done with a Sanyar video laryngoscope

Category

Treatment - Devices

2

Description

Control group: Tracheal Intubation is performed by direct laryngoscopy

Category

Treatment - Devices

Recruitment centers

1

Recruitment center

Name of recruitment center

Sina Hospital

Full name of responsible person

Mohammad Reza khajavi

Street address

Sina Hospital Hassan Abad sq, Emnam Khomini st.

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Tehran University of Medical Sciences

Full name of responsible person

Akbar Fotouhi

Street address

Central building of Tehran University of Medical sciences, Ghods st., Keshavarz blv.

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Grant name

Grant code / Reference number

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

Yes

Title of funding source

Tehran 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

Tehran University of Medical Sciences

Full name of responsible person

Mohammad Reza Khajavi

Position

Anesthesiologist

Latest degree

Specialist

Other areas of specialty/work

Anesthesiology

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Person responsible for scientific inquiries

Contact

Name of organization / entity

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Web page address

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

Not applicable

Data Dictionary

Not applicable

Title and more details about the data/document

Main study outcome data

When the data will become available and for how long

Six months after the end of the study

To whom data/document is available

University and industry researchers

Under which criteria data/document could be used

Share experiences to increase the use of video laryngoscope

From where data/document is obtainable

khajavim@tums.ac.ir Dr.khajavi

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

Through Dr. Khajavi e-mail address: khajavim@tums.ac.ir

Comments