Evaluation of preoperative anxiety and fear of anesthesia using Apais score

Submission date	Recruitment status No longer recruiting	Prospectively registered		
15/02/2018		Protocol		
Registration date	Overall study status Completed	Statistical analysis plan		
19/02/2018		[X] Results		
Last Edited	Condition category	[] Individual participant data		
14/09/2018	Surgery			

Plain English summary of protocol

Background and study aims

Patients often feel anxiety prior to undergoing surgery. The reason for the anxiety can be both anesthesia used to numb the pain and the actual surgery. The preoperative anxiety that patients have can cause side effects. It is important for the reasons of anxiety to help them in future operations. The aim of this study is to investigate the preoperative patient anxiety and patients fear of anesthesia.

Who can participate?

Adults aged 18 to 95 who are scheduled to undergo elective surgery.

What does the study involve?

Participants are asked to fill out a short questionnaire at the anesthesia clinic about their feelings of anxiety prior to undergoing anesthesia. They are asked if they had received anesthesia before, if they have surgical briefing, and the cause of their anxiety.

What are the possible benefits and risks of participating?

The potential benefit is we can decrease participants anxiety with our interviews. There are no side effects of participating to our study.

Where is the study run from?
Ahi Evran University Medical Faculty (Turkey)

When is the study starting and how long is it expected to run for? February 2016 to July 2017

Who is funding the study?
Ahi Evran University Medical Faculty (Turkey)

Who is the main contact? Professor Ipek Edipoglu (Scientific) dripeks@yahoo.com

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

study no:2010-05/04

Study information

Scientific Title

Evaluation of preoperative anxiety and fear of anesthesia using the Amsterdam preoperative anxiety and information scale score: a prospective cohort stud

Study objectives

The aim of this study is to investigate how the patient's age, gender, the operation, surgical briefing, type of anesthesia recommended for the operation ahead, patient's prior anesthesia experience affect the patient's anxieties regarding anesthesia and surgery.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics Committee of Ahi Evran University on 27/04/2016, ref: 2016-05/04

Study design

Observational cross sectional cohort study

Primary study design

Observational

Secondary study design

Cross sectional study

Study setting(s)

Hospital

Study type(s)

Prevention

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Preoperative anxiety and patients' fear of anesthesia

Interventions

Interviews with the patients are performed when they came to anesthesia clinic for preoperative examination, under the supervision of a specialist anesthetist.

For the study, The Amsterdam Preoperative Anxiety and Information Scale (APAIS), which comprises 6 statements, is used. The answers are evaluated in two scales; the anxiety score and the desire for information score. Anxiety score was obtained by calculating the total scores assigned to the expressions "I am worried about the anesthesic", "The anesthesic is on my mind continually", "I am worried about the procedure", "The procedure is on my mind continually", to measure the patient's level of anxiety regarding the anesthesia and surgery. Desire for information score is obtained by calculating the total scores assigned to the expressions "I would like to know as much as possible about the anesthesic " and "I would like to know as much as possible about the procedure "to measure the patient's level of desire for information regarding the anesthesia and surgery. Higher scores indicate higher levels of anxiety and desire for information. Answers to the statements were evaluated with Likert Scale. (1-Not at all...,5-Extremely).

In addition, participants are asked whether they had received anesthesia due to any reason, if so, the type of anesthesia, and whether they received surgical briefing. Which anesthetic method recommended to the patient for the surgery they will undergo in the preoperative anesthesia clinic is recorded. Then, participants are asked about the cause of their anxiety regarding the anesthesia.

Intervention Type

Biological/Vaccine

Primary outcome measure

- 1. Demographics is measured using the patient notes at the preoperative anesthesia visit
- 2. Type of anaesthesia patient notes is measured using at the preoperative anesthesia visit
- 3. Prior experience with anaesthesia patient interviews is measured using at the preoperative anesthesia visit
- 4. Anxiety regarding anaesthesia is measured using APAIS (Amsterdam preoperative anxiety and information scale) at the preoperative anesthesia visit.

Secondary outcome measures

There are no secondary outcome measures.

Overall study start date

20/02/2016

Completion date

20/07/2017

Eligibility

Key inclusion criteria

- 1. ASA I-III
- 2. Adult patients who are scheduled to undergo elective surgery in our hospital
- 3. Adequate language skills for the interview were included in our study
- 4. 18-95 (all adults)

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Upper age limit

95 Years

Sex

Both

Target number of participants

600

Key exclusion criteria

- 1. Pediatric patients
- 2. Patients who did not give consent
- 3. Patients with mental retardation
- 4. Alzheimer's disease
- 5. Dementia or psychiatric disorders
- 6. Emergency and ASA-IV patients

Date of first enrolment

28/05/2016

Date of final enrolment

20/05/2017

Locations

Countries of recruitment

Türkiye

Study participating centre Ahi Evran University Medical Faculty

Department of Anesthesiology and Intensive Care Medicine Kirsehir Türkiye 40100

Sponsor information

Organisation

Ahi Evran University Medical Faculty

Sponsor details

Department of Anesthesiology and Intensive Care Medicine Bağbaşı Mahallesi Şht. Sahir Kurutluoğlu Cd. Merkez Kirsehir Türkiye 40100

Sponsor type

Hospital/treatment centre

ROR

https://ror.org/05rrfpt58

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Ahi Evran University Medical Faculty clinical investigations

Results and Publications

Publication and dissemination plan

Planned publication in a high-impact peer reviewed journal. Additional documents will not be available.

Intention to publish date

20/07/2018

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Dr. Ipek S. Edipoglu at dripeks@yahoo.com.

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	11/09/2018		Yes	No