Preoperative risk estimation for onco-geriatric patients.

No registrations found.

Ethical review	Not applicable
Status	Suspended
Health condition type	-
Study type	Observational non invasive

Summary

ID

NL-OMON29384

Source
NTR

Brief title
PREOP

Health condition

Surgical oncology
Elderly

Sponsors and support

Primary sponsor: No sponsors
Source(s) of monetary or material Support: No funding source

Intervention

Outcome measures

Primary outcome

- Primary endpoint will be 30 day morbidity and mortality.

Secondary outcome

- Secondary endpoints will be length of hospital stay and the number of additional specialists involved in patient care in the 30 days after surgery.

Study description

Background summary

-

Study objective

This study will be performed to test the predictive value of the GFI, the VES-13 and the timed “up and go” test compared to components of PACE in elderly patients of the age of 70 years and above undergoing surgery for a solid tumour.

Study design

Data will be collected preoperatively and up to 30 days postoperatively

Intervention

The combination of components of PACE, the timed “up and go” test, the VES-13 and the GFI will be administered to every patient within two weeks prior to the surgical procedure. This can be done by a nurse in the preoperative clinic and will take approximately 30 minutes.

Patient data will be recorded during the length of the hospital stay, the patient file will be used to retrieve these data.

Preoperative data:

- medication used

- living situation

- comorbidity

- nutritional status.

Perioperative data:

- type of surgery

- duration of anaesthesia

- blood loss

- length of hospital stay

- postoperative complications

- consultation by other specialist.

Follow up: at the postoperative visit to the outpatient clinic additional data on morbidity occurring during the first 30 days postoperatively can be recorded as well as any additional specialists that were involved in patient care in the first 30 postoperative days.

Contacts

Public
University Medical Center Groningen (UMCG), Department of Surgery,
P.O. Box 30.001
B.L. Leeuwen, van
Groningen 9700 RB
The Netherlands
+31 (0)50 3612301
Scientific
University Medical Center Groningen (UMCG), Department of Surgery,
P.O. Box 30.001
B.L. Leeuwen, van
Groningen 9700 RB
The Netherlands
+31 (0)50 3612301

Eligibility criteria

Inclusion criteria

- Patients aged >70 years undergoing elective surgery for a solid tumour under general anaesthesia will be included for this study.

- Patients will be stratified according to tumour localisation: intra cavity (e.g. colorectal, gynaecological) vs superficial (e.g. breast, head & neck, melanoma).

Exclusion criteria

- Patients not able to give informed consent

Study design

Design

Study type:	Observational non invasive
Intervention model:	Parallel
Allocation:	Non-randomized controlled trial
Masking:	Open (masking not used)
Control:	N/A , unknown

Recruitment

NL

Recruitment status:	Suspended
Start date (anticipated):	01-09-2008
Enrollment:	362
Type:	Anticipated

Ethics review

Not applicable

Application type:	Not applicable

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
NTR-new	NL1497
NTR-old	NTR1567
Other	:
ISRCTN	ISRCTN wordt niet meer aangevraagd

Study results

Summary results
N/A