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Home visits by a nurse practitioner of patients who have undergone an operation for esophageal carcinoma: a randomized study

Submission date	Recruitment status	[_] Prospectively registered
28/04/2006	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
28/04/2006	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
11/01/2021	Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

NL585, NTR641

Study information

Scientific Title
Home visits by a nurse practitioner of patients who have undergone an operation for esophageal carcinoma: a randomized study

Acronym
VETO study

Study objectives
Have home visits performed by a nurse practitioner, compared with standard medical follow-up, a positive impact on the quality of life of patients who have undergone an operation for esophageal carcinoma?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Ethics approval received from the local medical ethics committee

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Esophageal cancer

Interventions
Nurse-led follow-up by home visits versus standard medical follow-up (out-patient clinic).

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Quality of Life.

Secondary outcome measures
1. Patient satisfaction
2. Costs

Overall study start date
15/01/2004

Completion date
01/05/2007

Eligibility

Key inclusion criteria
1. Patients have undergone esophageal cancer surgery
2. Informed consent

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
100

Total final enrolment
109

Key exclusion criteria
Patients are unable to understand the Dutch language.

Date of first enrolment
15/01/2004

Date of final enrolment
01/05/2007

Locations

Countries of recruitment
Netherlands

Study participating centre
Erasmus Medical Center
Rotterdam
Netherlands
3000 CA

Sponsor information

Organisation
Erasmus Medical Center, Department of Gastroenterology and Hepatology (The Netherlands)

Sponsor details
P.O. Box 2040
Rotterdam
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Sponsor type
University/education

ROR
https://ror.org/018906e22

Funder(s)

Funder type
Charity

Funder Name
Dutch Digestive Diseases Foundation

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	13/01/2009	11/01/2021	Yes	No