

Pilot study on the effects of oral immunonutrition on the quality of life in patients receiving palliative treatment for oesophageal and proximal gastric cancer.

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 05/03/2020	Condition category Cancer	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
N0055143763

Study information

Scientific Title

Pilot study on the effects of oral immuno-nutrition on the quality of life in patients receiving palliative treatment for oesophageal and proximal gastric cancer.

Study objectives

Does EPA supplementation affect patients' quality of life and survival with known esophageal and proximal gastric cancer who are receiving non-curative treatments?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Cancer: Gastrointestinal

Interventions

Patients, after obtaining informed consent, are randomly assigned one of the nutritional supplements, both of similar calorific value but one contains EPA.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Measurements of QoL, weight etc. are made at monthly interval, follow-up until death.

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/06/2004

Eligibility

Key inclusion criteria

Patients with histologically diagnosed oesophageal or proximal gastric cancer that are not receiving curative treatment.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/06/2002

Date of final enrolment

30/06/2004

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

North Cumbria Acute Hospitals NHS Trust

Carlisle

United Kingdom

CA2 7HY

Sponsor information**Organisation**

Department of Health

Funder(s)**Funder type**

Government

Funder Name

North Cumbria Acute Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration