

Minimally invasive oesophageal adenocarcinoma sentinel node biopsy 4.0

Submission date 05/04/2012	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 24/07/2012	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 06/07/2018	Condition category Cancer	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

<http://cancerhelp.cancerresearchuk.org/trials/a-study-looking-finding-first-lymph-nodes-cancer-may-spread-people-having-surgery-cancer-food-pipe>

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Protocol serial number

11479

Study information

Scientific Title

A prospective, observational study evaluating a minimally invasive technique of identifying the sentinel lymph nodes in patients undergoing surgery for oesophageal adenocarcinoma

Study objectives

Oesophagectomy for cancer carries some of the highest morbidity and mortality risks of any elective surgical procedure. The incidence of oesophageal adenocarcinoma in the Western world is increasing dramatically and shows no signs of slowing down. Endoscopic surveillance of Barrett's oesophagus is also anticipated to increase numbers of patients with high-grade dysplasia of the oesophagus who may require resectional surgery. Studies indicate that the routinely performed extended lymphadenectomy contributes significantly to the risks of surgery. However, determining which patients do not require a radical lymphadenectomy is limited by the relatively low sensitivity and specificity of pre-operative staging investigations. Minimally invasive techniques such as laparoscopic gastric mobilisation and thoracoscopic oesophageal mobilisation have been developed to reduce surgical trauma. These techniques need to be equally as radical in terms of the lymphadenectomy although this can be difficult in some cases. The sentinel lymph node (SLN) concept is that if the first draining lymph node in proximity to a cancer is clear of cancer cells then no other nodes should be involved.

A minimally invasive resection technique coupled to a sensitive minimally invasive SLN assessment could potentially reduce surgical morbidity by identifying patients who could have less extensive surgery without compromising oncological clearance. The SLN status might be the determining factor in deciding whether a patient is indeed suitable for minimally invasive surgery. Furthermore, the technique could also be applied to tailor surgery for those patients having open resections and determine the lymph node status for patients having endoscopic mucosal and submucosal resections for very superficial adenocarcinoma.

Ethics approval required

Old ethics approval format

Ethics approval(s)

NRES Committee North East County Durham & Tees Valley, First MREC 04/10/2011, ref: 11/NE/0262

Study design

Observational trial

Primary study design

Observational

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Upper Gastro-Intestinal Cancer

Interventions

Laparoscopic gamma probe identification of abdominal sentinel lymph nodes following endoscopic peritumoural technetium injection.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

1. Sentinel lymph node identification
2. Numbers of patients in whom the abdominal sentinel lymph node is identified after study completion

Key secondary outcome(s)

No secondary outcome measures

Completion date

01/10/2013

Eligibility

Key inclusion criteria

1. All patients with potentially curable lower third oesophageal adenocarcinoma or oesophagogastric junction adenocarcinoma planned to undergo curative twostage oesophagectomy
2. Male and female participants

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Incurable disease
2. Not suitable for laparoscopy

Date of first enrolment

20/02/2012

Date of final enrolment

01/10/2013

Locations

Countries of recruitment

United Kingdom

England

Study participating centre
Royal Victoria Infirmary
Newcastle upon Tyne
United Kingdom
NE1 4LP

Sponsor information

Organisation
Newcastle upon Tyne Hospitals NHS Foundation Trust (UK)

ROR
<https://ror.org/05p40t847>

Funder(s)

Funder type
University/education

Funder Name
Bupa Foundation

Alternative Name(s)

Funding Body Type
Private sector organisation

Funding Body Subtype
Trusts, charities, foundations (both public and private)

Location
United Kingdom

Funder Name
Royal College of Surgeons of England

Alternative Name(s)

RCS England, RCS ENG, The Royal College of Surgeons of England, RCS

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Abstract results	results	16/09/2016		No	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes