

A study of lapatinib in combination with oxaliplatin and capecitabine in oesophageal and gastric cancers

Submission date	Recruitment status	<input checked="" type="checkbox"/> Prospectively registered
27/05/2011	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
27/05/2011	Completed	<input checked="" type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
25/10/2022	Cancer	

Plain English summary of protocol

<https://www.cancerresearchuk.org/about-cancer/find-a-clinical-trial/a-trial-looking-lapatinib-with-chemotherapy-for-cancer-of-the-food-pipe-and-stomach-leo>

Contact information

Type(s)

Scientific

Contact name

Dr Paula Kareclas

Contact details

Addenbrookes Hospital
Oncology Clinical Trials S4, Box 279
Hills Road
Cambridge
United Kingdom
CB2 0QQ

Additional identifiers

Clinical Trials Information System (CTIS)

2010-019602-16

Protocol serial number

9848

Study information

Scientific Title

A study of Lapatinib in combination with oxaliplatin and capecitabine in Early HER-2 overexpressing Oesophageal and gastric cancers

Acronym

LEO

Study objectives

This study will test whether the ex vivo molecular response on a pre-treatment biopsy can predict the molecular response on a biopsy taken after 10 days of treatment with lapatinib. The study will also report observations of patterns of radiological, functional imaging and pathological response associated with molecular response

Ethics approval required

Old ethics approval format

Ethics approval(s)

South West 3 REC, 26/10/2010, 10/H0106/73

Study design

Non-randomised, interventional, observational qualitative trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Upper Gastro-Intestinal Cancer; Oesophagus, Stomach

Interventions

1. All 25 patients will be treated with lapatinib, there is no control arm
2. Fresh biopsies will be taken prior to lapatinib treatment start and 10 days post treatment start
3. All patients will have blood tests and renal function assessment prior to study entry and then after 31, 52 and 72 days of treatment
4. Treatment with lapatinib, oxaliplatin and capecitabine
5. All patients have a CT scan of thorax and abdomen prior to therapy and then another one after 3 cycles of chemotherapy for tumour assessment
6. All patient undergo a baseline whole body FDG-PET/CT examination at study entry and then 10 days after start of lapatinib therapy
7. Follow up length: 24 month(s)
8. Study entry : registration only

Intervention Type

Other

Phase

Phase IV

Primary outcome(s)

1. Molecular response
2. Timepoint(s): 10 days post treatment with lapatinib

Key secondary outcome(s)

1. FDG PET response timepoint(s): 10 days post treatment with lapatinib
2. Objective radiological response timepoint(s): 72 days post treatment
3. Pathological complete response timepoint(s): 102 days post treatment

Completion date

31/07/2015

Eligibility

Key inclusion criteria

1. Histologically confirmed gastric or oesophageal adenocarcinoma
2. Human Epidermal growth factor Receptor 2 (HER-2) 3+ on IHC OR HER-2 2+ on IHC but shown to have HER-2 amplification by (Fluorescent In-Situ Hybridization) FISH
3. Decision to treat with curative intent
4. Deemed to require chemotherapy prior to surgery using standard management algorithms
5. Ability to swallow oral medication
6. Baseline 18FDG PET/CT scan showing no evidence of distant metastases
7. Adequate haematological parameters: ANC = $1.0 \times 10^9/L$; WBC = $3.0 \times 10^9/L$; Plt = $100 \times 10^9/L$; haemoglobin (Hb) = 9g/dL (can be post-transfusion)
8. Adequate renal function (Measured or calculated creatinine clearance = 60 ml/min- if calculated the Cockcroft-Gault equation (Appendix A) to be used
9. Adequate liver function: serum Bilirubin = $1.5 \times ULN$; ALT/AST = $1.5 \times ULN$; ALP = $2.5 \times ULN$
10. Age >18 years
11. Women of child bearing potential using medically approved contraception (Postmenopausal women must have been amenorrheic for at least 12 months to be considered of non-childbearing potential)
12. Male patients using barrier contraceptives during the trial and for 6 months after the completion of the trial
13. Target gender: male & female
14. Lower age limit 18 years

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Total final enrolment

10

Key exclusion criteria

1. Advanced disease not amenable to surgery
2. Previous diagnosis of malignancy (excluding adequately treated Cervical carcinoma in situ or Basal cell carcinoma of the skin)
3. Abnormal Cardiac function (LVEF below normal as measured by echocardiogram or MUGA scan)
4. History of clinically significant cardiac disease e.g. symptomatic coronary artery disease, uncontrolled cardiac dysrhythmia or myocardial infarction within the last 12 months)
5. History of interstitial lung disease (e.g., pneumonitis or pulmonary fibrosis) or evidence of interstitial lung disease on baseline chest CT scan
6. Known peripheral neuropathy >Grade 1 (absence of deep tendon reflexes as the sole neurological abnormality does not render the patient ineligible)
7. Inability to give informed consent
8. Hypersensitivity to lapatinib or oxaliplatin or capecitabine
9. Prior treatment with chemotherapy or lapatinib or other specific anticancer therapy
10. Squamous cell carcinomas, unclear differentiation type, sarcomas, carcinoid or GIST
11. Known positive tests for human immunodeficiency virus (HIV) infection, hepatitis C virus, acute or chronic active hepatitis B infection
12. Pregnancy/breastfeeding
13. Subjects who have current active hepatic or biliary disease (with exception of patients with Gilbert's syndrome, asymptomatic gallstones, liver metastases or stable chronic liver disease per investigator assessment)
14. Treatment with another investigational agent within 30 days of commencing study treatment
15. Known or suspected dihydropyrimidine dehydrogenase deficiency (DPD)
16. Galactose intolerance, Lapp lactase deficiency or glucose-galactose malabsorption

Date of first enrolment

17/06/2011

Date of final enrolment

16/10/2013

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Addenbrooke's Hospital

Cambridge

United Kingdom

CB2 0QQ

Study participating centre
University College London Hospital
235 Euston Road
London
United Kingdom
NW1 2BU

Sponsor information

Organisation
Cambridge University Hospitals NHS Foundation Trust (UK)

ROR
<https://ror.org/04v54gj93>

Funder(s)

Funder type
Industry

Funder Name
GlaxoSmithKline

Alternative Name(s)
GlaxoSmithKline plc., GSK plc., GlaxoSmithKline plc, GSK

Funding Body Type
Government organisation

Funding Body Subtype
For-profit companies (industry)

Location
United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

The protocol has not specified these data sharing requirements, and patients have not consented to data sharing (recruited from 2011–2013) therefore this is not applicable. The trial is currently in the process of being archived.

IPD sharing plan summary

Not expected to be made available

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	03/11/2015		Yes	No
HRA research summary			28/06/2023	No	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
Plain English results			25/10/2022	No	Yes