

Pharmacokinetic study of adjuvant capecitabine after resection of pancreatic adenocarcinoma

Submission date 15/06/2009	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 28/08/2009	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 20/03/2019	Condition category Cancer	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol

<http://www.cancerhelp.org.uk/trials/a-study-looking-capecitabine-after-surgery-for-cancer-pancreas-cap-001>

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

NCT00854477

Secondary identifying numbers

CAP001

Study information

Scientific Title

A pharmacokinetic study of adjuvant capecitabine in patients who have undergone proximal pancreatico-duodenectomy for resection of pancreatic adenocarcinoma

Acronym

CAP001

Study objectives

1. To establish the pharmacokinetics (PK) of capecitabine in patients who have undergone proximal pancreatico-duodenectomy, i.e. the action of drug capecitabine in the body over a period of time, including the processes of absorption, distribution, localisation in tissues, biotransformation and excretion
2. To ensure equivalent capecitabine exposure when compared to previous studies using patients who have not undergone such surgery

Ethics approval required

Old ethics approval format

Ethics approval(s)

London, Northwick Park Hospital REC Committee, 07/08/2009, REC ref: 09/H0718/27

Study design

Multicentre non-randomised single-arm open-label study

Primary study design

Interventional

Secondary study design

Non randomised study

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied

Pancreatic adenocarcinoma

Interventions

This is a clinical trial to evaluate the PK of adjuvant capecitabine in patients who have undergone proximal pancreatico-duodenectomy. The study also aims to establish the toxicity profile of capecitabine in these patients, to identify any dose limiting toxicities (DLT), and to ensure equivalent capecitabine exposure when compared to previous studies using patients who have

not undergone such surgery. Screening tests will consist of demographic details, complete medical history, physical exam, vital signs, tumour serum markers, haematology and biochemistry tests. There will also be an electrocardiogram (ECG), faecal elastase measurement and a serum or urine pregnancy test (for women of childbearing potential). Haematology and biochemistry (including CA19.9) will be repeated prior to each study drug administration.

All patients will receive 8 cycles of oral capecitabine chemotherapy at a dose of 1250 mg/m², administered twice daily at 12-hourly intervals for 14 consecutive days out of a 21-day cycle. Total proposed duration of therapy is 24 weeks, assuming patients commence all cycles without delay. Capecitabine and its metabolites (DFCR, DFUR and 5-FU) plasma levels will be measured during cycles 1 and 3 in all patients.

An optional pharmacogenetic blood sample will be collected prior to the start of chemotherapy treatment. Treatment should continue for 8 cycles unless there is evidence of disease progression, or unacceptable toxicity.

Intervention Type

Drug

Phase

Phase I

Drug/device/biological/vaccine name(s)

Capecitabine

Primary outcome measure

To establish the PK of capecitabine in patients who have undergone proximal pancreaticoduodenectomy, measured at week 7.

Secondary outcome measures

1. To establish the toxicity profile of capecitabine in these patients and to identify any dose limiting toxicities (DLT), measured at week 24
2. To ensure equivalent capecitabine exposure when compared to previous studies using patients who have not undergone such surgery, measured at PK analysis

Overall study start date

01/08/2009

Completion date

31/08/2012

Eligibility

Key inclusion criteria

1. Complete macroscopic resection for pathologically proven ductal adenocarcinoma (or poorly differentiated/undifferentiated carcinoma) of the pancreas (R0 or R1 resection)
2. Surgery must have included a proximal pancreatico-duodenectomy
3. Histological confirmation of the primary diagnosis and examination of all resection margins
4. At least 4 weeks since surgery, fully recovered from the operation and all surgical wounds fully healed

5. Aged greater than or equal to 18 years, either sex
6. World Health Organisation (WHO) performance status of less than or equal to 2
7. Haematological and biochemical indices (these measurements must be performed within one week prior to the patient being registered on the study):
 - 7.1. Haemoglobin (Hb) greater than or equal to 9.0 g/dl (patients may be transfused to this level, however, Hb must be above 9.0 g/dl before registration)
 - 7.2. Neutrophils greater than or equal to $1.5 \times 10^9/l$
 - 7.3. Platelets (Plts) greater than or equal to $100 \times 10^9/l$
 - 7.4. Serum bilirubin less than or equal to 1.5 x upper normal limit (ULN)
 - 7.5. Alanine amino-transferase (ALT) and/or aspartate amino-transferase (AST) less than or equal to 2.0 x ULN (if both are measured, both must be less than or equal to 2.0 x ULN)
 - 7.6. Calculated creatinine clearance greater than or equal to 50 ml/min (uncorrected value) or isotope clearance measurement greater than or equal to 50 ml/min
8. Female patients of child-bearing potential must have a negative serum or urine pregnancy test within two weeks prior to enrolment and agree to use appropriate medically approved contraception for four weeks prior to entering the trial, during the trial, and for six months afterwards
9. Male patients must agree to use appropriate medically approved contraception during the trial and for six months afterwards
10. Written, informed consent provided
11. Ability of the patient to co-operate with treatment and follow up must be ensured
12. Patients receiving oral anti-coagulation prior to entry into the study must be converted to low molecular weight heparin in light of the interaction between capecitabine and warfarin

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

12

Key exclusion criteria

1. Patients with pancreatic lymphoma or other histological diagnosis
2. Macroscopically remaining tumour (R2 resection)
3. Evidence of malignant ascites, peritoneal or liver metastasis, or spread to other distant abdominal or extra-abdominal organs
4. History of confirmed ischaemic heart disease, concurrent congestive heart failure or prior history of class III/IV cardiac disease
5. Concurrent mechanical or malabsorptive disorders precluding affective oral administration of the drug (excluding malabsorption related directly to proximal pancreatic-duodenectomy)
6. Pregnancy or lactation
7. Patients known to be serologically positive for hepatitis B, hepatitis C or human immunodeficiency virus (HIV)

8. Patients who are high medical risks because of non-malignant systemic disease including active uncontrolled infection
9. Any other serious medical or psychological condition precluding adjuvant treatment
10. Patients with any other condition which in the Investigator's opinion would not make the patient a good candidate for the clinical trial

Date of first enrolment

01/11/2009

Date of final enrolment

06/12/2011

Locations

Countries of recruitment

England

United Kingdom

Study participating centre**Oncology Centre**

Cambridge

United Kingdom

CB2 0QQ

Sponsor information

Organisation

Cambridge University Hospitals NHS Foundation Trust (UK)

Sponsor details

R&D Department, Box 146

Addenbrooke's Hospital

Hills Road

Cambridge

England

United Kingdom

CB2 0QQ

Sponsor type

Hospital/treatment centre

Website

http://www.cuh.org.uk/addenbrookes/addenbrookes_index.html

ROR

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

Funder(s)

Funder type

Industry

Funder Name

Roche (UK)

Alternative Name(s)

F. Hoffmann-La Roche Ltd, F. Hoffmann-La Roche & Co, F. Hoffmann-La Roche AG, Roche Holding AG, Roche Holding Ltd, Roche Holding, Roche Holding A.G., Roche Holding, Limited, F. Hoffmann-La Roche & Co.

Funding Body Type

Government organisation

Funding Body Subtype

For-profit companies (industry)

Location

Switzerland

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?
HRA research summary			28/06/2023	No	No