A prospective, phase III, controlled, multicentre, randomised clinical trial comparing combination gemcitabine and capecitabine therapy with concurent and sequential chemoimmunotherapy using a telomerase vaccine in locally advanced and metastatic pancreatic cancer

Submission date	Recruitment status No longer recruiting	[X] Prospectively registered		
28/11/2005		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
14/12/2005	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
27/07/2022	Cancer			

Plain English summary of protocol

https://www.cancerresearchuk.org/about-cancer/find-a-clinical-trial/a-trial-looking-at-a-vaccine-called-gv1001-for-pancreatic-cancer-that-has-spread

Study website

http://www.cancer.gov/clinicaltrials/CRUK-TELOVAC-V4

Contact information

Type(s)

Scientific

Contact name

Dr Gary Middleton

Contact details

Royal Surrey County Hospital Egerton Road Guildford United Kingdom GU2 7XX

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

NCT00425360

Secondary identifying numbers

N/A

Study information

Scientific Title

A prospective, phase III, controlled, multicentre, randomised clinical trial comparing combination gemcitabine and capecitabine therapy with concurent and sequential chemoimmunotherapy using a telomerase vaccine in locally advanced and metastatic pancreatic cancer

Acronym

TeloVac

Study objectives

In patients with locally advanced or metastatic pancreatic adenocarcinoma, does the addition of telomerase vaccine GV1001, when given concurrently or sequentially, to combination gemcitabine and capecitabine, improve survival over treatment with combination gemcitabine and capecitabine alone.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethical review pending

Study design

Phase III, prospective, open-label, controlled, multicentre, randomised clinical trial comparing combination gemcitabine and capcitabine with GV1001 in pancreatic cancer with follow-up.

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Locally advanced and metastatic pancreatic cancer

Interventions

Arm 1 - Gemcitabine and capecitabine therapy: Gemcitabine will be administered on day one, eight and 15 followed by seven days rest. Per oral capecitabine will be administered morning and evening for 21 days followed by seven days rest. Gemcitabine and capecitabine therapy cycles will be repeated every four weeks until withdrawal from trial treatment.

Arm 2 - Gemcitabine and capecitabine then sequential GV1001 therapy: Patients will receive two cycles of combination gemcitabine and capecitabine before commencing GV1001 alone. Each of the two cycles of combination gemcitabine and capecitabine consists of:

Gemcitabine will be administered on day one, eight and 15 followed by seven days rest. Per oral Capecitabine will be administered morning and evening for 21 days followed by seven days rest. Following completion of gemcitabine and capecitabine therapy, GV1001 will be administered intradermally on Monday, Wednesday and Friday during week eight and once weekly during weeks nine, ten, 11, 13 and 17. Thereafter, vaccinations will follow a once monthly schedule until withdrawal from trial treatment.

Arm 3 - Concurrent Gemcitabine, Capecitabine and GV1001 therapy: Gemcitabine will be administered on day one, eight and 15 followed by seven days rest. Per oral Capecitabine will be administered morning and evening for 21 days followed by seven days rest. Gemcitabine and Capecitabine therapy cycles will be repeated every four weeks until withdrawal from trial treatment. GV1001 will be administered intradermally on Monday, Wednesday and Friday during week one followed by a once weekly schedule for weeks two, three, four, six and ten. Thereafter, GV1001 will be administered once monthly until withdrawal from trial treatment.

Intervention Type

Drug

Phase

Phase III

Drug/device/biological/vaccine name(s)

Gemcitabine, capecitabine and telomerase vaccine

Primary outcome measure

Length of survival

Secondary outcome measures

- 1. Time to Progression
- 2. Quality of life
- 3. Clinical Benefit Response
- 4. Objective response rates according to RECIST criteria
- 5. Toxicity
- 6. Survival and response according to Delayed Type Hypersensitivity

Overall study start date

01/04/2006

Completion date

15/03/2013

Eligibility

Key inclusion criteria

- 1. Aged over 18 years
- 2. Histologically or cytologically proven pancreatic ductal adenocarcinoma carcinoma
- 3. Locally advanced or metastatic disease precluding curative surgical resection
- 4. Contrast enhanced Computed Tomography (CT) scan of the thorax, abdomen and pelvis within 28 days of randomisation
- 5. Unidimensionally measurable disease (CT) in accordance with the Response Evaluation Criteria in Solid Tumors (RECIST) guidelines
- 6. World Health Organisation (WHO) performance status zero, one or two
- 7. Platelets more than $100 \times 10^9 / l$; white blood cell count (WBC) more than $3 \times 10^9 / l$; neutrophils more than $1.5 \times 10^9 / l$ at entry
- 8. Serum bilirubin less than 35 µmol/l
- 9. Calculated creatinine clearance over 50 ml/min
- 10. No concurrent uncontrolled medical condition
- 11. No previous malignant disease other than non-melanotic skin cancer or carcinoma in situ of the uterine cervix
- 12. Life expectancy more than three months
- 13. Adequate contraceptive precautions if relevant
- 14. Informed written consent

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Not Specified

Target number of participants

1100

Total final enrolment

1062

Kev exclusion criteria

- 1. Medical or psychiatric conditions compromising informed consent
- 2. Intracerebral metastases or meningeal carcinomatosis
- 3. Clinically significant serious disease or organ system disease not currently controlled on present therapy
- 4. Uncontrolled angina pectoris
- 5. Pregnancy or breast feeding

- 6. Treatment with chemotherapy, radiotherapy or other investigational drug within the last four weeks prior to inclusion
- 7. Known malabsorption syndromes
- 8. Patients with a known hypersensitivity to Fluorouracil (5-FU) or with a Dihydropyrimidine Dehydrogenase (DPD) deficiency
- 9. Immunosuppressive therapy less than four weeks prior to the start of treatment
- 10. People of child-bearing potential unless effective methods of contraception are used

Date of first enrolment

01/04/2006

Date of final enrolment

15/03/2013

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Royal Surrey County Hospital Guildford United Kingdom GU2 7XX

Sponsor information

Organisation

The University of Liverpool (UK)

Sponsor details

Research and Business Services
The Foresight Centre
1 Brownlow Street
Liverpool
England
United Kingdom
L69 3GL

Sponsor type

University/education

Website

http://www.liv.ac.uk

ROR

https://ror.org/04xs57h96

Funder(s)

Funder type

Charity

Funder Name

Cancer Research UK (C11497/A5690)

Alternative Name(s)

CR_UK, Cancer Research UK - London, CRUK

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

United Kingdom

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

Not provided at time of registration

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	01/07/2014		Yes	No
Plain English results			27/07/2022	No	Yes