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A randomised trial of CF (infusional 5-fluorouracil and cisplatin) alone versus CF plus concurrent radiotherapy in patients with locally advanced pancreatic carcinoma

Submission date 19/08/2002	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 19/08/2002	**Overall study status** Completed	
Last Edited 24/10/2019	**Condition category** Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

Protocol serial number
PANCRAD

Study information

Scientific Title

A randomised trial of CF (infusional 5-fluorouracil and cisplatin) alone versus CF plus concurrent radiotherapy in patients with locally advanced pancreatic carcinoma

Study objectives
Not provided at time of registration

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)
Not Specified

Health condition(s) or problem(s) studied
Pancreas cancer

Interventions
1. Regimen A: 5-fluorouracil, continuous infusion for 18 weeks, plus cisplatin repeated every 3 weeks for six cycles.

2. Regimen B: Cisplatin repeated every 3 weeks for four cycles plus 5-fluorouracil, continuous infusion for 12 weeks followed by continuous infusion for a further 6 weeks at a reduced dose. Radiotherapy 50 Gy in twenty-five fractions given over 5 weeks. Radiotherapy to commence on week 13 of chemotherapy.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Cancer drugs

Primary outcome(s)
Not provided at time of registration

Key secondary outcome(s))
Not provided at time of registration

Completion date
11/11/1994

Eligibility

Key inclusion criteria
1. Histological evidence of locally advanced or unresectable pancreatic adenocarcinoma
2. Patients evaluable for response must have bidmensionally measurable disease as assessed by Computed Tomography (CT) scans
3. No prior chemotherapy or radiotherapy
4. Life expectancy of >3 months
5. Adequate bone marrow and renal function
6. World Health Organisation (WHO) performance status 0-2 at randomisation
7. No medical contraindications to treatment protocols

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
01/01/1989

Date of final enrolment
11/11/1994

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
UKCCCR Register Co-ordinator
London
United Kingdom
NW1 2DA

Sponsor information

Organisation
The Royal Marsden NHS Foundation Trust (UK)

ROR
https://ror.org/0008wzh48

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Royal Marsden Hospital (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration