ISRCTN59778423 https://doi.org/10.1186/ISRCTN59778423

A pilot study to compare CAMPATH-1H produced from YO or CHO cells in patients with B-cell chronic lymphocytic leukaemias (Protocol CLL-TF57)

Submission date
12/09/2003

Recruitment status
No longer recruiting

Registration date
12/09/2003

Overall study status
Completed

Last Edited
14/02/2018

Condition category
Cancer

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Martin Dyer

Contact details
University Hospitals of Leicester
c/o Research and Development Office
Leicester General Hospital NHS Trust
Leicester
United Kingdom
LE1 4PW
+44 (0)116 258 4109
nicola.turner@uhl-tr.nhs.uk

Additional identifiers

Protocol serial number
N0123119552

Study information

Scientific Title
A pilot study to compare CAMPATH-1H produced from YO or CHO cells in patients with B-cell chronic lymphocytic leukaemias (Protocol CLL-TF57)

Study objectives
To determine and compare the rates of lymphocyte clearance with YO-CAMPATH-1H and CHO-CAMPATH-1H in patients with advanced CLL or T-cell prolymphocytic leukemia (T-PLL) who have failed prior chemotherapy to test whether YO-CAMPATH is more potent.

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)
Treatment

Health condition(s) or problem(s) studied
B-cell chronic lymphocytic leukaemia

Interventions
Randomised controlled trial

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
Rates of clearance of malignant lymphocytes from peripheral blood with the two antibody preparations to allow direct comparison of the efficacy of both.

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

Completion date
01/07/2003

Eligibility

Key inclusion criteria
Haematology patients

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
01/07/2002

Date of final enrolment
01/07/2003

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
University Hospitals of Leicester
Leicester
United Kingdom
LE1 4PW

Sponsor information

Organisation
Department of Health (UK)

Funder(s)

Funder type

Hospital/treatment centre

Funder Name
University Hospitals of Leicester NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration