

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	☐ Prospectively registered
		☐ Protocol
Registration date 12/09/2003	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 14/02/2018	Condition category Cancer	☐ 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

Primary study design

Interventional

Study design

Randomised controlled trial

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