

UKALL R2 - Acute Lymphoblastic Leukaemia trial

Submission date 06/04/2000	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 06/04/2000	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 04/10/2007	Condition category Cancer	☐ Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
G8223452

Study information

Scientific Title

Acronym
UKALL R2

Study objectives

To compare continuing chemotherapy with unrelated donor BMT in relapsed childhood ALL following a common re-induction and consolidation regimen.

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

Acute Lymphoblastic Leukaemia

Interventions

Continuing chemotherapy/unrelated donor BMT

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Survival, disease free survival

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/09/2008

Eligibility

Key inclusion criteria

They are children under 15 years at original diagnosis and treated on UKALL trials or pilot studies with first relapse less than 4 years from the start of treatment of ALL at any site and no suitable related bone marrow treatment (BMT) donor is available

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

1. They are multiply relapsed patients
2. They have a prior history of toxicity or organ damage such that completion of the protocol is felt unlikely at the outset
3. They have had an isolated CNS or testicular relapse more than 6 months off treatment.

Date of first enrolment

01/02/1995

Date of final enrolment

30/09/2008

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

The Children's Hospital

Birmingham

United Kingdom

B16 8ET

Sponsor information**Organisation**

Medical Research Council (MRC) (UK)

Funder(s)**Funder type**

Research council

Funder Name

UK Medical Research Council

Results and Publications

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?
Results article	Results	01/07/2005		Yes	No