

# Non-Hodgkins lymphoma T-cell protocol

<b>Submission date</b> 01/07/2001	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 01/07/2001	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 23/02/2015	<b>Condition category</b> Cancer	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

**Contact name**  
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**Contact details**  
UKCCCR Register Co-ordinator  
MRC Clinical Trials Unit  
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London  
United Kingdom  
NW1 2DA

## Additional identifiers

**Protocol serial number**  
NHL9503

## Study information

**Scientific Title**  
Non-Hodgkins lymphoma T-cell protocol

**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)**

Treatment

**Health condition(s) or problem(s) studied**

Lymphoma (non-Hodgkin's)

**Interventions**

1. Group A: Chemotherapy including third intensification block at 35 weeks
2. Group B: Chemotherapy without third intensification block at 35 weeks

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

28/02/2000

**Eligibility**

**Key inclusion criteria**

1. T-cell non-Hodgkin's lymphoma
2. Stages I-IV
3. Confirmed remission achieved
4. Less than 25% blasts in bone marrow
5. No circulating blasts in blood
6. Aged <18 years

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Child

**Upper age limit**

18 years

**Sex**

All

**Key exclusion criteria**

Patients with T-cell anaplastic lymphoma or peripheral T-cell lymphoma are excluded.

**Date of first enrolment**

01/01/1997

**Date of final enrolment**

28/02/2000

## Locations

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

**MRC Clinical Trials Unit**

London

United Kingdom

NW1 2DA

## Sponsor information

**Organisation**

Cancer Research UK (CRUK) (UK)

**ROR**

<https://ror.org/054225q67>

## Funder(s)

**Funder type**

Charity

**Funder Name**

Cancer Research UK

**Alternative Name(s)**

CR\_UK, Cancer Research UK - London, Cancer Research UK (CRUK), CRUK

**Funding Body Type**

Private sector organisation

**Funding Body Subtype**

Other non-profit organizations

**Location**

United Kingdom

**Funder Name**

United Kingdom Children's Cancer Study Group (UKCCSG)

## Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary