ISRCTN18173987 https://doi.org/10.1186/ISRCTN18173987

A trial to compare radiotherapy alone with radiotherapy followed by chemotherapy in the treatment of non-Hodgkin's lymphoma

Submission date 01/07/2001	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 01/07/2001	**Overall study status** Completed	
Last Edited 15/01/2019	**Condition category** Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr - -

Contact details
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
NHD1

Study information

Scientific Title
A trial to compare radiotherapy alone with radiotherapy followed by chemotherapy in the treatment of non-Hodgkin's lymphoma

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

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Lymphoma (non-Hodgkin's)

Interventions
1. Group A: Radical radiotherapy, 30 Gy in 15 fractions over 3 weeks
2. Group B: Radical radiotherapy, 30 Gy in 15 fractions over 3 weeks followed 1 month later by chemotherapy. Chemotherapy with chlorambucil for 14 consecutive days repeated every 28 days for 2 years.

Intervention Type
Mixed

Primary outcome measure
Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date
01/01/1990

Completion date
03/11/1992

Eligibility

Key inclusion criteria
1. Aged 15-65 years
2. Proven non-Hodgkin's disease with good pathology involving nodes, Waldeyer's ring is also included provided that nodes are also involved
3. Stage I or II

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
Patients with abdominal nodes only are excluded

Date of first enrolment
01/01/1990

Date of final enrolment
03/11/1992

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

MRC Clinical Trials Unit
London
United Kingdom
NW1 2DA

Sponsor information

Organisation
Cancer Research UK (CRUK) (UK)

Sponsor details
PO Box 123
Lincoln's Inn Fields
London
United Kingdom
WC2A 3PX
+44 (0)207 317 5186
kate.law@cancer.org.uk

Sponsor type
Charity

Website
http://www.cancer.org.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, CRUK

Funding Body Type
Private sector organisation

Funding Body Subtype
Other non-profit organizations

Location
United Kingdom

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration