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Randomised Controlled Trial of CID versus CD for Induction of Remission in Low Grade Non-Hodgkin's Lymphoma and Randomised Controlled Assessment of Interferon as Maintenance Treatment after Remission Induction

Submission date	Recruitment status	[_] Prospectively registered
19/08/2002	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
19/08/2002	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
30/11/2015	Cancer	

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
SNLG NHLVIII

Study information

Scientific Title
Randomised Controlled Trial of CID versus CD for Induction of Remission in Low Grade Non-Hodgkin's Lymphoma and Randomised Controlled Assessment of Interferon as Maintenance Treatment after Remission Induction

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)
Not Specified

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

Interventions
INDUCTION / CONSOLIDATION:
Patients are randomised to one of two treatment arms:
1. Arm A: Chemotherapy with Chlorambucil, idarubicin and dexamethasone (CID), cycle to be repeated every 21 days. Patients showing a response following three courses will receive three further consolidation courses.
2. Arm B: Chemotherapy with Chlorambucil and dexamethasone (CD), cycle to be repeated every 21 days. Patients showing a response following three courses will receive three further consolidation courses.

RADIOTHERAPY: Patients may be given localised radiotherapy to initial areas of bulky disease if a complete response in that particular site has not been obtained following induction.

MAINTENANCE: Patients who have shown a complete response or good partial response following six cycles of induction therapy are eligible for the second randomisation. Patients are randomised to one of three groups:
1. Group A: No further therapy.
2. Group B: Low dose interferon, 1MU subcutaneously three times a week until progression or maximum of 36 months.
3. Group C: High dose interferon, 3MU subcutaneously three times a week until progression or maximum of 36 months.

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
30/06/2000

Eligibility

Key inclusion criteria
1. Kiel classification of low grade non-Hodgkin's lymphoma
2. Stages II to IV
3. Age 15 to 70 years
4. Measurable disease
5. No prior chemotherapy
6. No central nervous system (CNS) involvement
7. Eastern Cooperative Oncology Group (ECOG) performance status of 0-2
8. Adequate bone marrow, renal and hepatic function
9. No medical contraindications to protocol treatments

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Sex
Not Specified

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/1995

Date of final enrolment
30/06/2000

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
UKCCCR Register Co-ordinator
London
United Kingdom
NW1 2DA

Sponsor information

Organisation
Scotland & Newcastle Lymphoma Group (UK)

Funder(s)

Funder type
Research organisation

Funder Name
Scottish & Newcastle Lymphoma Group (UK)

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	01/11/2006		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes