

Trial of PACEBOM versus CHOP in histologically aggressive non-Hodgkins lymphoma

Submission date 01/07/2001	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 01/07/2001	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 01/02/2012	Condition category Cancer	<input type="checkbox"/> Individual participant data

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
LY3

Study information

Scientific Title

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-Hodgkins)

Interventions

Patients were randomised to one of two regimens:

1. CHOP Regimen: Multi-drug chemotherapy with cyclophosphamide, hydroxydaunorubicin, vincristine and prednisolone (CHOP) repeated every 28 days. A minimum of six courses to be given with two courses beyond the attainment of complete response
2. PACEBOM Regimen: Multi-drug chemotherapy with prednisolone, adriamycin, cyclophosphamide and etoposide (PACE) alternating every 7 days with bleomycin, vincristine and methotrexate (BOM). Six cycles of PACE and five of BOM to be given

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/11/1987

Completion date

31/10/1992

Eligibility

Key inclusion criteria

1. Aged 16 to 69 years
2. Previously untreated histologically aggressive lymphoma with a large cell component: Diffuse large cell, Diffuse immunoblastic, Diffuse mixed cell
3. Stages II-IV
4. No contraindications to treatment protocols

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Patients with Burkitt's and lymphoblastic lymphoma to be excluded

Date of first enrolment

01/11/1987

Date of final enrolment

31/10/1992

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
UKCCCR Register Co-ordinator
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

Funder Name

Lymphoma Research Trust

Funder Name

Lisa Lear Fund

Funder Name

Isle of Man Anti-Cancer Association

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/07/1996		Yes	No