

# Randomised comparison of cyclical anthracycline-based chemotherapy [PA(B1)OE] with alternating chemotherapy [Ch1VPP/PABLOE] in advanced Hodgkin's disease

<b>Submission date</b> 01/07/2001	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 01/07/2001	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
<b>Last Edited</b> 01/02/2012	<b>Condition category</b> Cancer	<input type="checkbox"/> 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**  
HO3001

## 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

**Study type(s)**

Treatment

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

Lymphoma (Hodgkin's)

**Interventions**

FIRST RANDOMISATION: Patients are randomised to one of two chemotherapy regimens:

1. PA(B1)OE Regimen: Multi-drug chemotherapy with adriamycin, vincristine, prednisolone, etoposide and bleomycin (PA(B1)OE) repeated every 21 days for six to eight courses. Bleomycin is given for the first four course only.
2. Ch1VPP/PABLOE Regimen: Multi-drug chemotherapy with chlorambucil, procarbazine, prednisolone and vinblastine(CH1VPP) alternating with PA(B1)OE. The total cycle Ch1VPP/PA (B1)OE takes 7 weeks. A minimum of six, three each of Ch1VPP and PA(B1)OE, and a maximum of eight courses of chemotherapy to be given.

SECOND RANDOMISATION: Patients in complete remission following chemotherapy whose original presentation was with bulky (>5 cm) nodal disease are eligible for the second randomisation. Patients are randomised to one of two groups:

1. Group A: Radiotherapy 35-40 Gy given over 4 weeks.
2. Group B: No radiotherapy.

**Intervention Type**

Drug

**Phase**

Not Specified

**Drug/device/biological/vaccine name(s)**

Cancer drugs

**Primary outcome(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

01/04/1996

**Eligibility****Key inclusion criteria**

1. Previously untreated and properly stage patients with Hodgkin's disease for whom chemotherapy is indicated, ie stage I and IIA (poor prognosis), IB, IIB, III and IV
2. Patients must be free from any irreversible medical condition that would drastically limit their life span or prohibit use of combination chemotherapy
3. Aged 15 to 69 years inclusive

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex****Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/01/1990

**Date of final enrolment**

01/04/1996

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre**

UKCCCR Register Co-ordinator

London

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# 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

British National Lymphoma Investigation (BNLI)

# Results and Publications

## Individual participant data (IPD) sharing plan

## IPD sharing plan summary

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
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[Results article](#)

Results

18/05/2001

Yes

No