

Comparison of intermediate and high dose valaciclovir for the prophylaxis of CMV viraemia in allogeneic stem cell transplant recipients

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 30/11/2015	Condition category Infections and Infestations	<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

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Additional identifiers

Protocol serial number
N0256114044

Study information

Scientific Title

Comparison of intermediate and high dose valaciclovir for the prophylaxis of CMV viraemia in allogeneic stem cell transplant recipients

Study objectives

Does high dose valaciclovir four times daily have greater efficacy in reducing CMV viraemia than valaciclovir in allogeneic stem cell recipients?

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

Infections and Infestations: Cytomegalovirus (CMV) infection

Interventions

Randomised controlled trial.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

1. Rate of CMV viraemia
2. Time to CMV viraemia
3. Incidence of CMV disease
4. Survival

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/07/2004

Eligibility

Key inclusion criteria

110 patients, 55 in each treatment arm.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

03/07/2002

Date of final enrolment

01/07/2004

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

The Royal Free & University Medical School

London

United Kingdom

NW3 2QG

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

The Royal Free Hampstead NHS Trust (UK)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration