

An investigation of the predictability of carboplatin plasma concentrations in children following dosing on the basis of surface area of renal function

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

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr --

Contact details

UKCCCR Register Co-ordinator

MRC Clinical Trials Unit

222 Euston Road

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Additional identifiers

Protocol serial number

NAG9402

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

Cancer of brain and nervous system

Interventions

Patients receiving two courses of carboplatin are randomised to one of two groups:

1. Group A: Surface area based (course I) followed by renal function based (course II) measurement.
2. Group B: Renal function based (course I) followed by surface area based (course II) measurement.

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

28/02/1997

Eligibility

Key inclusion criteria

1. Aged under 18 years
2. Receiving carboplatin on at least two occasions

3. Being treated according to listed UKCCSG protocols (NB9001, NB9301, NB8702, GC8901, CNS9204, CNS9102, WAG8702)

4. Tumour site: Neuroblastoma; Extra-cranial germ cell; Brain

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Upper age limit

18 years

Sex

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/1993

Date of final enrolment

28/02/1997

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

UKCCCR Register Co-ordinator

London

United Kingdom

NW1 2DA

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

United Kingdom Children's Cancer Study Group (UKCCSG)

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?
Results article	results	01/11/2000		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes