

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

<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

### Contact name

Dr - -

### Contact details

UKCCCR Register Co-ordinator  
MRC Clinical Trials Unit  
222 Euston Road  
London  
United Kingdom  
NW1 2DA

## 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?
<a href="#">Results article</a>	results	01/11/2000		Yes	No
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes