

A randomised, placebo controlled trial of prednisone in early Henoch Schonlein Purpura

Submission date
08/03/2004

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
09/03/2004

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
24/08/2007

Condition category
Haematological Disorders

☐ 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
95/25S(E)

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

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Henoch Schonlein Purpura

Interventions

Intervention:

Prednisone 2 mg/kg for 7 days, 75% on days 8 and 9, 50% on days 10 and 11 and 25% on days 12, 13, 14, then stopped.

Control:

Comparison group was an identical appearing placebo.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Prednisone

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/01/2000

Eligibility

Key inclusion criteria

Eligible participants were children between the ages of 2 and 15, presenting to the emergency room within 7 days of onset of Henoch Schonlein Purpura.

Children were excluded if:

1. Another explanation for purpura was present
2. They had a known underlying systemic vasculitis
3. They had been treated with any form of corticosteroids in the preceding month
4. They had a known chronic illness affecting the renal, gastrointestinal or immune systems
5. They had an active infection
6. They were experiencing a life-threatening complication of Henoch Schonlein Purpura

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/09/1996

Date of final enrolment

31/01/2000

Locations

Countries of recruitment

Canada

Study participating centre

IWK Health Centre

Halifax

Canada

B3J 3G9

Sponsor information

Organisation

Children's Hospital of Eastern Ontario Research Institute (CHEORI) (Canada)

ROR

<https://ror.org/05nsbhw27>

Funder(s)

Funder type

Research organisation

Funder Name

Children's Hospital of Eastern Ontario Research Institute (CHEORI) (Canada) (ref: 95/25S[E])

Results and Publications

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	02/04/2004		Yes	No