ISRCTN85109383 https://doi.org/10.1186/ISRCTN85109383

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

Submission date
08/03/2004

Recruitment status
No longer recruiting

Registration date
09/03/2004

Overall study status
Completed

Last Edited
24/08/2007

Condition category
Haematological Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] 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

Study design
Randomised controlled trial

Primary study design
Interventional

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