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Efficacy of EPs 7630 in children with acute non-streptococcal tonsillopharyngitis

Submission date 26/03/2003	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 26/03/2003	**Overall study status** Completed	
Last Edited 07/10/2021	**Condition category** Respiratory	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
UM007.2

Study information

Scientific Title

Efficacy of EPs 7630 in children with acute non-streptococcal tonsillopharyngitis

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

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Acute tonsillopharyngitis

Interventions
78 Children were randomised to receive either:
1. Herbal remedy EPs 7630, 20 drops per hour during the first 1 - 2 days, followed by 20 drops three times daily
2. Placebo, 20 drops per hour during the first 1 - 2 days, followed by 20 drops three times daily

The duration of individual treatment lasted over a maximum of 6 days.

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
EPs 7630

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/2003

Completion date
31/12/2003

Eligibility

Key inclusion criteria
Patients who met the following inclusion criteria were suitable for the trial:
1. Age 6¿10 years, acute tonsillopharyngitis, duration of complaints less than 48 h, negative dip-and-react-test test for ß-hemolytic streptococcus and severity of symptoms ≥6 points
2. In addition legal guardians had to sign an informed consent

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
78

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/2003

Date of final enrolment
31/12/2003

Locations

Countries of recruitment
Germany

Study participating centre
Director Research Center HomInt
Karlsruhe
Germany
76202

Sponsor information

Organisation
ISO Arzneimittel GmbH & Co KG (Germany)

Sponsor details
Bunsenstrasse 6-10
Ettlingen
Germany
76275

Sponsor type
Industry

Website
http://www.iso-arznei.de

ROR
https://ror.org/045xrc244

Funder(s)

Funder type
Industry

Funder Name
ISO Arzneimittel GmbH & Co KG (Germany)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

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			07/10/2021	Yes	No