

Epidemiology and treatment of soil-transmitted helminthiasis, with particular consideration to strongyloidiasis, in Yunnan province, China

Submission date 16/08/2008	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 21/08/2008	Overall study status Completed	☐ Protocol
Last Edited 13/11/2008	Condition category Infections and Infestations	☐ Statistical analysis plan
		☒ 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

Study information

Scientific Title

Study objectives

1. Single-dose albendazole and single-dose tribendimidine (both drugs will be administered orally at 200 mg for children aged 5-14 years, and 400 mg for individuals aged ≥ 15 years) are safe and efficacious against soil-transmitted helminth infections (*Ascaris lumbricoides*, hookworm and *Trichuris trichuria*)
2. Single-dose tribendimidine has an effect on *Strongyloides stercoralis* and *Taenia* spp

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics Committee of the University and the State of Basel (Ethikkommission beider Basel) (EKBB) Date of approval: 12/06/2007 (ref: 149/07)

Primary study design

Interventional

Study design

Open-label, randomised controlled trial

Study type(s)

Health condition(s) or problem(s) studied

Helminth infections

Interventions

The participants who were diagnosed with helminthiases at the baseline parasitological survey were randomly allocated to the following arms:

Arm 1: Single-dose albendazole (oral), 200 mg for children aged 5-14 years, and 400 mg for individuals aged ≥ 15 years

Arm 2: Single-dose tribendimidine (oral), 200 mg for children aged 5-14 years, and 400 mg for individuals aged ≥ 15 years

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Albendazole, tribendimidine

Primary outcome(s)

1. Reduction of infection prevalence of intestinal helminths, assessed by examination of stool samples 2-4 weeks after drug administration
2. Frequency and severity of adverse events were recorded within 24 hours after drug administration

Key secondary outcome(s)

Kato-Katz-derived egg count reduction of common soil-transmitted helminths, assessed 2-4 weeks after drug administration.

Completion date

31/07/2007

Eligibility

Key inclusion criteria

1. Both males and females, age ≥ 5 years
2. Submission of ≥ 1 stool sample for the baseline parasitological survey
3. For females, not pregnant, as verbally assessed by medical personnel on the day of treatment
4. Absence of major systemic illnesses, as assessed by medical personnel on the day of treatment
5. Written informed consent by the head of the household on behalf of the whole family

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Other

Sex

All

Key exclusion criteria

1. Presence of any abnormal medical condition, as judged by the medical personnel
2. No stool sample submitted for baseline parasitological survey
3. Enrolled in any other clinical investigation during the study
4. For females: pregnancy

Date of first enrolment

01/05/2007

Date of final enrolment

31/07/2007

Locations

Countries of recruitment

China

Switzerland

Study participating centre

Department of Public Health and Epidemiology
Basel
Switzerland
4002

Sponsor information

Organisation

Swiss National Science Foundation (SNSF) (Switzerland)

ROR

<https://ror.org/00yjd3n13>

Funder(s)

Funder type

Research organisation

Funder Name

Main funder:

Funder Name

Commission for Research Partnership with Developing Countries (KFPE) through the Swiss Agency for Development and Cooperation (SDC) (Switzerland)

Funder Name

Other funders:

Funder Name

Voluntary Academic Society (Freiwillige Akademische Gesellschaft) (Switzerland)

Funder Name

Janggen-Pöhn Foundation (Janggen-Pöhn-Stiftung) (Switzerland)

Funder Name

Chinese Ministry of Science and Technology, through their support to the Key Laboratory of Parasite and Vector Biology of the Chinese Ministry of Health (China) (grant ref: 2005DKA21104)

Funder Name

Swiss National Science Foundation (Switzerland) (project numbers PPOOB-102883 and PPOOB-119129)

Alternative Name(s)

Schweizerischer Nationalfonds, Swiss National Science Foundation, Fonds National Suisse de la Recherche Scientifique, Fondo Nazionale Svizzero per la Ricerca Scientifica, Fonds National Suisse, Fondo Nazionale Svizzero, Schweizerische Nationalfonds, The Swiss National Science Foundation (SNSF), SNF, SNSF, FNS

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

Switzerland

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	01/08/2008		Yes	No