# Procalcitonin-guided antibiotic use in Acute Respiratory Tract Infections (ARTIs) in primary care

Submission date

18/01/2005

Registration date
15/02/2005

Completed

Condition category

Recruitment status

[] Prospectively registered

[X] Protocol

Statistical analysis plan

[X] Results

[X] Individual participant data

Plain English summary of protocol

Not provided at time of registration

Respiratory

## Contact information

Type(s)

Scientific

20/09/2017

Contact name

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# Additional identifiers

ClinicalTrials.gov (NCT) NCT00099840

Protocol serial number EKBB 254/04

# Study information

#### Scientific Title

Procalcitonin-guided antibiotic use in Acute Respiratory Tract Infections (ARTIs) in primary care

#### Acronym

**PARTI-Study** 

#### Study objectives

Acute respiratory tract infections (ARTI) are among the most frequent reasons for seeking medical attention in primary care. Although from predominantly viral origin, ARTIs are the most important condition for the prescription of antibiotics (AB), mainly due to the difficulty in primary care to differentiate between viral and bacterial etiology. Unnecessary AB use increases drug expenditures, side effects and AB resistance. A novel approach is to guide AB use by procalcitonin (ProCT), since serum levels are elevated in bacterial infections but remain lower in viral infections and inflammatory diseases. We aim to compare a strategy based on evidence-based guidelines with ProCT guided AB therapy in ARTIs with respect to outcome (days with restriction) and AB use.

#### 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

Acute respiratory tract infections

#### **Interventions**

All participating physicians will receive evidence-based guidelines for the management of patients with ARTIs. Patients with ARTI and in need of antibiotics by physicians' clinical judgment and with informed consent will be randomized to procalcitonin (ProCT) guided antibiotic prescription ("ProCT group") versus guidelines guided antibiotic prescription ("control group").

#### Intervention Type

Other

#### Phase

**Not Specified** 

#### Primary outcome(s)

Days with restrictions from ARTI

#### Key secondary outcome(s))

- 1. Rate of AB prescriptions; days with AB use
- 2. Symptoms from ARTI
- 3. Relapse rate from ARTI within 28 days
- 4. Days with side effects from ABs and off work
- 5. Cost-effectiveness

#### Completion date

31/03/2006

# **Eligibility**

#### Key inclusion criteria

18 years or older, with ARTI of >1 and <28 days duration and in need of antibiotics based on the clinical judgment of the primary care physician

#### Participant type(s)

Patient

#### Healthy volunteers allowed

No

#### Age group

Adult

#### Lower age limit

18 years

#### Sex

All

## Key exclusion criteria

- 1. Patients without informed consent
- 2. Not fluent in German
- 3. Antibiotic pretreatment in previous 28 days
- 4. Severe immune-suppression

#### Date of first enrolment

01/12/2004

#### Date of final enrolment

31/03/2006

## Locations

#### Countries of recruitment

Switzerland

# Study participating centre University Hospital Basel

Basel Switzerland 4031

# Sponsor information

#### Organisation

University Hospital Basel (Switzerland)

#### **ROR**

https://ror.org/04k51q396

# Funder(s)

#### Funder type

University/education

#### **Funder Name**

University Hospital Basel - Clinic of Endocrinology, Basel Institute of Clinical Epidemiology (BICE), Dept. of Internal Medicine, Dept. of Central Laboratories (infrastructure)

#### **Funder Name**

BRAHMS AG, Hennigsdorf, Germany (assay material)

#### **Funder Name**

Schweizerischer Nationalfonds zur Förderung der Wissenschaftlichen Forschung

#### 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	13/10/2008	Yes	No
Protocol article	protocol	18/08/2005	Yes	No
Other publications	secondary analysis	24/03/2016	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/2025	No	Yes