

# Granulocyte Colony-Stimulating Factor (G-CSF) and Interleukin-8 (IL-8) for early diagnosis of sepsis in neonates and critically ill children: safety and cost effectiveness of a new laboratory prediction model

<b>Submission date</b> 05/08/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 28/10/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 10/10/2014	<b>Condition category</b> Infections and Infestations	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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CH-8032

## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

## Secondary identifying numbers

N/A

# Study information

## Scientific Title

Not provided at time of registration

## 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)

Hospital

## Study type(s)

Diagnostic

## Participant information sheet

## Health condition(s) or problem(s) studied

Sepsis in neonates and critically ill children

## Interventions

New laboratory markers for infection given to physicians in charge of the patients in order to improve diagnostic work-up.

## Intervention Type

Other

## Phase

Not Applicable

## 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**

Newborns and critically ill paediatric patients of a tertiary University Children's Hospital

**Participant type(s)**

Patient

**Age group**

Other

**Sex**

Both

**Target number of participants**

Not provided at time of registration

**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**

Switzerland

**Study participating centre**

University Children's Hospital

Zurich

Switzerland

CH-8032

# Sponsor information

## Organisation

Chance for the Critically Ill Child Foundation (Stiftung Chance fuer das kritisch kranke Kind)  
(Switzerland)

## Sponsor details

Steinwiesstrasse 75  
Zurich  
Switzerland  
CH-8032

## Sponsor type

Charity

## ROR

<https://ror.org/02htkw777>

# Funder(s)

## Funder type

Charity

## Funder Name

Chance for the Critically Ill Child Foundation (Stiftung Chance fuer das kritisch kranke Kind)  
(Switzerland)

# 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?
<a href="#">Protocol article</a>	protocol	01/12/2004		Yes	No

