

## ORGANISATION

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### Workshop Fee

400 € Non-Member

300 € Member of AGAH | EUFEMED and DGKliPha  
or young scientists until the age of 30 years

100 € Member of Regulatory Agencies and  
ethics commission

Delegate number is limited to 70 persons.

**Final Registration: November 1, 2019**

## VENUE

### GSI - Gustav Stresemann Institut Bonn

Langer Grabenweg 68

53175 Bonn

Web: [www.gsi-bonn.de](http://www.gsi-bonn.de)

## REGISTRATION

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# AGAH

Arbeitsgemeinschaft  
für angewandte  
Humanpharmakologie e.V.

Association for Applied  
Human Pharmacology

## AGAH DISCUSSION FORUM



## HOW TO INTERPRETE AN INVESTIGATOR'S BROCHURE FOR MEANINGFUL RISK ASSESSMENT

**NOVEMBER 11, 2019  
BONN**

## CONTENT

The Investigator's Brochure is the key document for the Sponsor to compile the pertinent knowledge on the pharmaceutical, pharmacological, and toxicological characteristics of and –if any- the clinical experience with the investigational medicinal product (IMP). Critical appraisal of potential safety risks and appropriate precautions on how to avoid putting healthy subjects or patients receiving the IMP at risk is required to provide guidance for the investigator.

In first-in-human and early phase clinical trials, clinical experience with the IMP is lacking or sparse and non-clinical safety assessments must enable an adequate risk assessment to support the transition from animal models to man.

This AGAH discussion forum will debate what is required from an IB to provide proper guidance for the investigator about an IMP in early clinical trials.

The discussion forum addresses investigators and trial teams with some hands-on experience in early phase clinical trials. Clinicians performing first in human trials / early clinical trials in patients may equally profit from this workshop.

Hildegard Sourgens  
Past President EUFEMED\*

Kerstin Breithaupt-Grögler  
Consultant to the AGAH Board

AGAH e. V. is a founding member of the European Federation for Early Medicines Development, Brussels

## MORNING SESSION

- 08:00 Registration
- 09:00 Welcome and Moderation  
*K. Breithaupt, Frankfurt/M.  
H. Sourgens, München*  
**Online Survey** (Who attends this workshop?)
- 09:15 **The Investigator's Brochure - Key information to support early clinical trials**  
*J. van Gerven, Leiden/NL*
- 10:00 **Deficiencies / Uncertainties in IBs - The Regulatory Authority's experience**  
*E. Stahl, Bonn; T. Sudhop, Bonn*  
**Online Survey** (Experience of the audience: Quality of the IB regarding Sections 1 to 6)  
*K. Erb-Zohar, Schotten; J. Rengelshausen, Aachen*
- 11:00 Break
- 11:30 **Guidance for the Investigator**  
Chair: *G. Mikus, Heidelberg*  
View point of Sponsor; Expectations from Regulatory Agency, Ethics Committee, and Investigator  
*F. Donath, Erfurt  
J. Hasford, München  
K. Francke, Berlin  
T. Sudhop, Bonn  
G. Wensing, Wuppertal*  
**Online Survey** (Experience of the audience: Quality of the IB regarding Section 7)  
*K. Erb-Zohar, Schotten; J. Rengelshausen, Aachen*
- 12:30 **Questions raised by the Audience - Open-Forum Discussion**  
*F. Donath, Erfurt  
J. Hasford, Munich  
S. Heil, Bonn  
G. Mikus, Heidelberg  
C. Riedel, Bonn  
T. Sudhop, Bonn  
G. Wensing, Wuppertal*

## AFTERNOON SESSION

- 13:00 Break
- 13:45 **Reference safety information – a key element of the IB**  
*K. Erb-Zohar, Schotten*  
**Online Survey** (Experience of the audience: Reference safety information in the IB)  
*K. Erb-Zohar, Schotten; J. Rengelshausen, Aachen*
- 14:15 **Discussion**  
*F. Donath, Erfurt  
K. Erb-Zohar, Schotten  
D. Kubitzka, Wuppertal  
G. Mikus, Heidelberg  
E. Stahl, Bonn*
- 14:30 **Potential risks for human subjects associated with inadequate non-clinical safety assessment**  
*S. Plassmann, Basel/CH; L. Wiesner, Bonn*
- 15:00 Break
- 15:15 Case study presented by *J. van Gerven, Leiden*:  
**Integrating data from the IB – a new tool for translational integration of preclinical effects**  
**Online Survey** (Experience of the audience: Room for improvement in the IB)  
*K. Erb-Zohar, Schotten; J. Rengelshausen, Aachen*
- 16:30 **Discussion**  
*S. Plassmann, Basel/CH  
J. van Gerven, Leiden/NL  
L. Wiesner, Bonn*
- 16:50 **Wrap-up**
- 17:00 End of Workshop