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Individual Account Application

Each entity submitting data to ClinicalTrials.gov must adhere to the following terms and conditions, which are intended to ensure the accuracy, currency and validity of the data.

1. Only data on trials approved by the appropriate regulatory authority may be submitted to ClinicalTrials.gov.
2. Notice of recruiting status changes must be done immediately, and all submitted data must be reviewed, verified, and updated every six months.
3. The submitting organization is responsible for the completeness and accuracy of the data submitted to ClinicalTrials.gov.
4. Trial data must be submitted in English.

Sponsor Information The sponsor organization is the entity with primary responsibility for planning and conducting the trial(s) to be registered.

Register STUDY: Trial1

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<th>Type of Organization</th>
<th>City</th>
<th>State</th>
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<td>Frankfurt</td>
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Organization Name: Johann Wolfgang Goethe University Hospitals

Organization Address: Theodor-Stern-Kai 7 60590 Frankfurt, Germany

Organization Affiliations and Accessors

Parent Organizations

Official Investigator: Prof. Dr. R. Kaufmann

Phone: +49 69 6301 7777

Email: rkaufmann@med.uni-frankfurt.de

Organization URL (optional)

Paying Organizations

Investigator Information

Investigator Name

Affiliation (for the sponsor)

Investigator Phone

Investigator Email

Regulatory Information

The regulatory authority must be a national or international health authority, an institutional review board, or an ethics committee.

Regulatory Authority

Regulatory Authority Address

To the best of my knowledge, the above information is true and current. Questions about this form and the Protocol Registration System (PRS) may be sent to

submit@ClinicalTrials.gov

Submit Application  Reset