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**Investigator agreement**

**Eurobact II: Epidemiology and determinants of outcomes of Hospital Acquired Blood Stream Infections in the Intensive Care.**

As an Investigator of the epidemiological research study Eurobact II, endorsed by ESICM Trials group and Infection section and ESMID ESCGIP and sponsored by OUTCOMEREA 51 rue des poilus, Drancy 93700 France

I agree to the following responsibilities:

* Comply with ICH Guidelines for Good Clinical Practice (GCP).
* Ensure that patient privacy is respected in compliance with all applicable local regulations.
* Comply with local regulations to conduct research, and as applicable:
* Apply for ethical approval and/or local site approvals and ensure they are in place prior to the initiation of the study.
* Where required inform and/or obtain consent from the patient or person responsible.
* Collaborate with the National coordinator (NC) for all ethical and regulatory approvals as required in my country and provide them with scanned copies of relevant documents.
* Conduct the study according to the protocol.
* Maintain effective communication with the NC and the sponsor, reply promptly to data queries from study data manager.
* Archive the required documents in the Investigator's Site File (ISF) and maintain for a period of at least 5 years. Make available all required patient records and all documents related to the study for use by potential auditors.
* I have received the current version of study documents, containing protocol, summary, CRF, log of patients included, Confidential Disclosure Agreement and template CV. I will review and sign study-related documents in a timely manner.
* I understand that the EUROBACT study has no funding that can be allocated to local investigators. All potential administrative fees should be paid by the participating site or investigator.



Investigator (Printed / typed name)



Investigator Signature Date

Version 1.0 dated 9 of July 2019.