Online
SPARK-BIH Educational/
Entrepreneurial Forum

BIH Berlin Institute of Health Charité & MDC

SPARK BIH

Regulatory Requirements for Medical Devices in Europe

27 October 2020 | 5-6 pm (CET) | SPARK-BIH Online Lecture

5 pm **Regulatory Requirements for Medical Devices in Europe – How to achieve market approval under the new Medical Device Regulations**
Dr. Sebastian Grömminger
Consultant for the approval of in-vitro diagnostics at
Johner Institut GmbH

What does a medical or diagnostic device need to fulfill to be commercialized in Europe?
Medical devices (MD) as well as in-vitro diagnostic devices (IVD) cannot be placed on the European market without conforming to the strict safety requirements of the European Union. These are regulated by the Medical device regulation (MDR) and the in-vitro diagnostic regulation (IVDR). In this Online Educational Forum Lecture Dr. Sebastian Grömminger, will help you to understand today's EU regulation requirements and its basic implementation lines.

Online via GoToMeeting | Please register here.

>> spark-bih.de
>> bihealth.org

SPARK-BIH is an initiative created at Stanford University, to overcome challenges associated with translation of academic discoveries.

BIH Innovations The Technology Transfer Office of BIH and Charité
CHARITÉ UNIVERSITÄTSMEDIZIN BERLIN
MDC MAX-DELBRÜCK-CENTRUM FÜR MOLEKULARE MEDIZIN IN DER HELMHOLTZ-GEMEINSCHAFT
STIFTUNG CHARITÉ