Training on Good Regulatory Practices and their Implementation in the Medical Devices Sector
Mexico
Session III

Date: November 17th, 2021
Time: 12:00 – 14:00 (Mexico City Time)
Language: Spanish
Platform: Zoom

TIME	AGENDA
12:00 – 12:05	**Housekeeping Message** *Sandra Ligia González, IACRC*
12:05 – 12:10	**Welcome Message** *Miriam Loera, COFEPRIS*
12:10 – 13:00	**The Process of Development of Technical Regulations Applicable to Medical Devices and In Vitro Diagnostic Reagents and the Guarantee of Compliance with International Obligations – Part I (40 min.)** *Moderator: Berenice Terrazas, CGJC, COFEPRIS* - **Regulation of Health Supplies** – *Jonathan R. Flores, CAS, COFEPRIS.* - **Official Mexican Standards** - *Victor Torres, DGN.* Q&A: 10 min.
13:00 – 14:00	**The Process of Development of Technical Regulations Applicable to Medical Devices and In Vitro Diagnostic Reagents and the Guarantee of Compliance with International Obligations – Part II (40 min.)** *Moderator: Jonathan R. Flores, CAS, COFEPRIS.* - **The Regulatory Process in COFEPRIS,** *Berenice Terrazas, CGJC, COFEPRIS.* ○ **Regulatory Impact Analysis, Public Consultations, TBT Notifications** Q&A: 10 min. - ***Pharmacopeia* of the United States of Mexico - Medical Device Supplement.** *Rafael Hernández, FEUM.* Q&A: 10 min.
13:50 – 14:00	**Closing Remarks III** *Miriam Loera, COFEPRIS; Daniel Vázquez, USAID; Sandra Ligia González, IACRC*