Department of Health Research
Ministry of Health and Family Welfare
Government of India

Central Drugs Standard Control Organization
Directorate General of Health Services
Ministry of Health & Family Welfare
Government of India

Regulatory Awareness & Experiential Learning Workshop for Medical Device/Diagnostic Innovators/Start-ups in India

A Joint Initiative of ICMR and CDSCO

Dr. Rajiv Bahl, Secretary,
DHR and DG, ICMR

Dr. Rajeev Singh Raghuvanshi,
DCGI, CDSCO

Sh. Aseem Sahu
Deputy Drug Controller, CDSCO
(Medical Devices)

Dr Nivedita Gupta
Sc-F & Head, Communicable
Diseases (CD) Division, ICMR

Dr. Suchita Markan
Sc-E, ICMR,
Mission In-charge, MDMS

Dr. Aparna Mukherjee
Sc-E, ICMR-
Clinical Studies and Trial Unit
(CSTU)

Topics:

- An overview of medical device regulations
  - Risk-based classification
  - Essential principles for safety & performance of medical devices
- Role of ICMR in assuring quality of medical devices/diagnostics
- Clinical investigation/clinical performance evaluation of medical device/diagnostics as per MDR, 2017
- Ethical guidelines for conducting clinical studies in India
- Experience sharing on CDSCO approved technologies – Case studies
- Q & A Session

12th Oct. 2023
10:00 AM – 5:00 PM
Venue: Conference Hall, ICMR Hqrs (Hybrid Mode)

Registration Link: https://forms.gle/eE8nsYQ15pyiwKh8A

Last Date for Registration: 30th Sep, 2023

Who Should Attend:

- Medical device/diagnostics manufacturers/Start-ups
- Biomedical engineers/medical device & diagnostic professionals
- Scientists/Clinicians/Healthcare professionals in the Medtech Innovation space