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MedTech mitra
Handholding MedTech innovators for clinical evaluation, regulatory facilitation and uptake of new products
A NITI Aayog-ICMR-CDSCO initiative



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



Handholding Support to MedTech Innovators: Essentials of Regulation Compliant Clinical Investigation

Under MedTech Mitra: A NITI Aayog-ICMR-CDSCO Initiative

August 08th, 2024



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



Dr. V. K. Paul
Member, NITI Aayog



Dr. Rajeev Singh Raghuvanshi,
DCGI, CDSCO

Topics to be covered:

- 1 Regulatory requirements of conducting Clinical Investigation for Investigational Medical Devices.
- 2 Regulatory requirements of Clinical Performance Evaluation of New In Vitro Diagnostics.
- 3 Basics of Clinical Study Design : Observational and Experimental.
- 4 Overview of Statistical Applications in Biomedical Research & sample size determination.
- 5 Experience sharing regarding clinical studies by Innovators on their CDSCO approved Technologies.

Registration Link: <https://forms.gle/k8EHsoBAsMK8aTAd9>

08th August 2024 | 10:00 AM – 5:30 PM | HYBRID MODE

VENUE: Conference Hall, ICMR Hqrs.

Last Date for Registration:
31st July, 2024