MedTech mitra

Handholding MedTech innovators for clinical evaluation, regulatory facilitation and uptake of new products

A NITI Aayog-ICMR-CDSCO initiative

UNDER THE GUIDANCE OF

NITI Aayog

स्वास्थ्य एवं परिवार कल्याण मंत्रालय
MINISTRY OF HEALTH AND FAMILY WELFARE

औषध विभाग
Department of Pharmaceuticals

About MedTech Mitra

Under the guidance of NITI Aayog, Indian Council of Medical Research in partnership with Central Drugs Standard Control Organization (CDSCO) aim to foster development of affordable and accessible indigenous Medical devices/ In-vitro diagnostics by providing strategic handholding support to MedTech innovators for clinical evaluation, regulatory facilitation and uptake of new products.

Scope of Facilitation by MedTech Mitra

Fostering product development through Engineers-Clinicians partnership

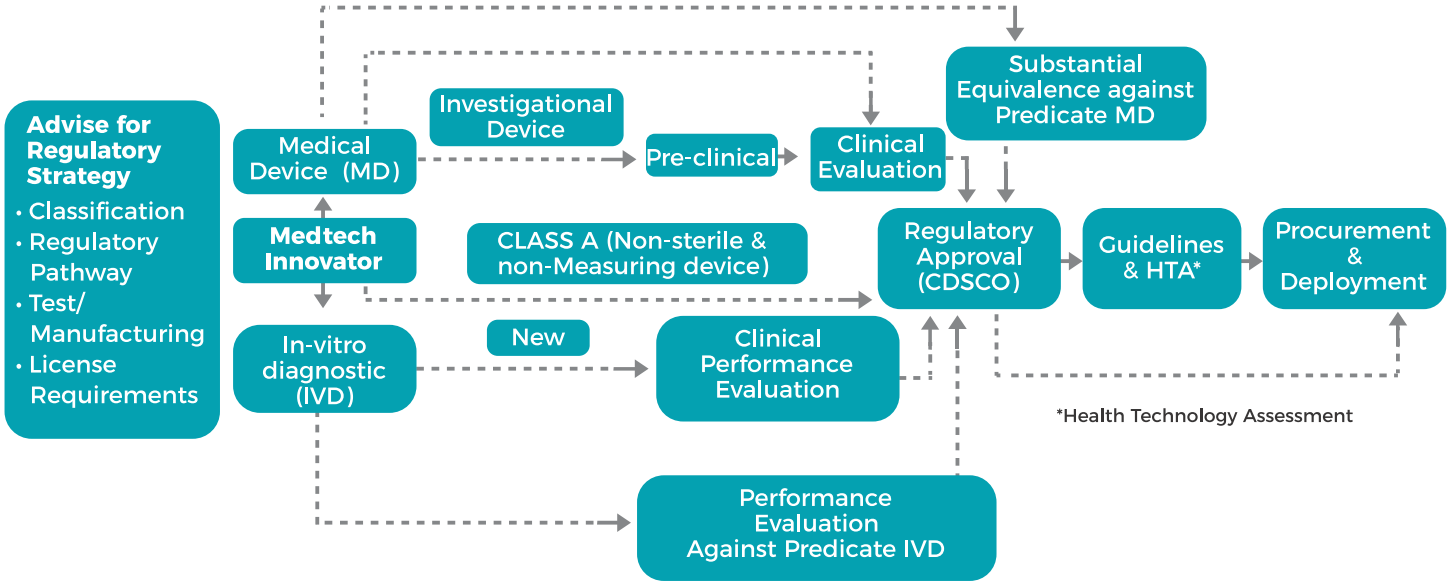
Handholding Support

- Regulatory facilitation
- Pre-clinical/ Clinical evaluation
- Guidelines & HTA (Health Technology Assessment)
- Uptake of New Products

Information cell

- Applicable Standards
- Testing and/or Manufacturing facilities
- Funding Opportunities
- I.P. Advisory

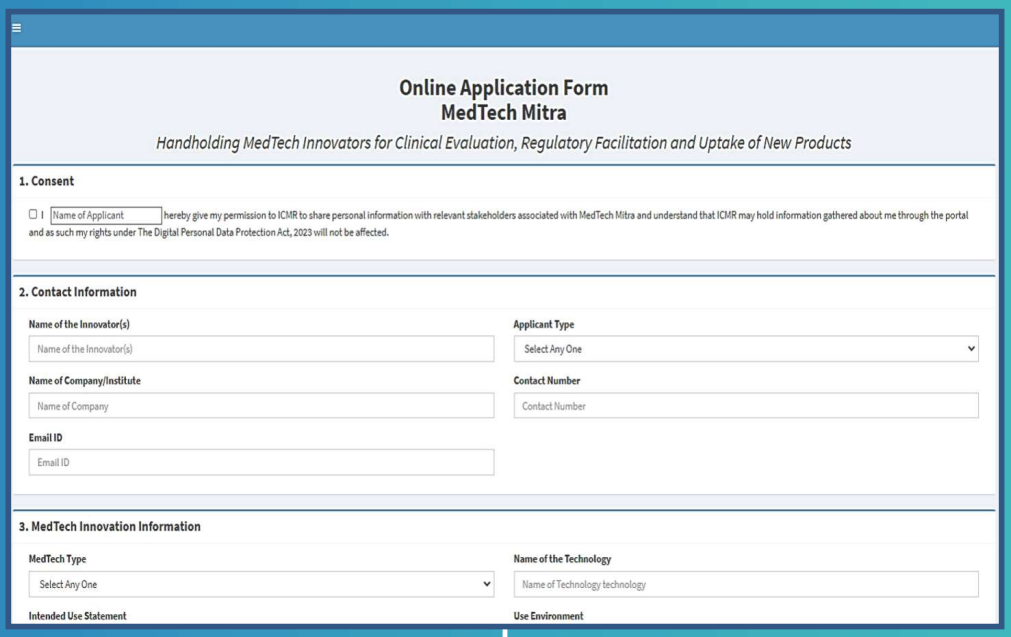
Handholding for Pre-clinical/ Clinical evaluation/ HTA/ Guidelines (GL)/ Public Access



How to reach us?

Innovator visits ICMR website (https://main.icmr.nic.in/)

Click on MedTech Mitra portal (medtechmitra. icmr. org.in) which displays the Application form



Online Application Form
MedTech Mitra

Handholding MedTech Innovators for Clinical Evaluation, Regulatory Facilitation and Uptake of New Products

1. Consent

☐ I Name of Applicant hereby give my permission to ICMR to share personal information with relevant stakeholders associated with MedTech Mitra and understand that ICMR may hold information gathered about me through the portal and as such my rights under The Digital Personal Data Protection Act, 2023 will not be affected.

2. Contact Information

Name of the Innovator(s) — Name of the Innovator(s)
Applicant Type — Select Any One
Name of Company/Institute — Name of Company
Contact Number — Contact Number
Email ID — Email ID

3. MedTech Innovation Information

MedTech Type — Select Any One
Name of the Technology — Name of Technology technology
Intended Use Statement
Use Environment

Innovator submits the query by filling the MedTech Mitra application form

An email acknowledging the submission is sent to the Innovator

Evaluation of adequacy of information for the query

Additional info to be submitted by innovator, if required

MedTech Mitra and Innovator meeting scheduled to understand and address the query

Technical & Regulatory Advice to all the Innovators

Requisite information sought provided to all the innovators

Render handholding support and/or financial assistance to selected Innovators as recommended

ABOUT ICMR

The Indian Council of Medical Research (ICMR), is an autonomous organization under the Department of Health Research (DHR) for the planning, promoting, coordinating and conducting biomedical research in India. The ICMR (established in 1911) is one of the oldest medical research organizations in the world with a broad mandate to generate new knowledge through the conduct and support of biomedical research in all areas that would have a bearing on improving the health of Indian people.

To cater to the huge import dependency in medical device and diagnostic sector and nations clarion call for self-reliance, Indian Council of Medical Research (ICMR) has initiated the Medical Device and Diagnostics Mission Secretariat (MDMS) under Division of Development Research to foster indigenous manufacturing of medical device and diagnostics for an AatmaNirbhar Bharat.

About CDSCO

The Central Drugs Standard Control Organisation (CDSCO) under Directorate General of Health Services, Ministry of Health & Family Welfare, Government of India is the National Regulatory Authority (NRA) of India. CDSCO is constantly thriving upon to bring out transparency, accountability and uniformity in its services in order to ensure safety, efficacy and quality of the medical product manufactured, importe and distributed in the country, through enforcement and implementation of the Medical Devices Rules, 2017.

Knowledge Partners

Key Coordinators