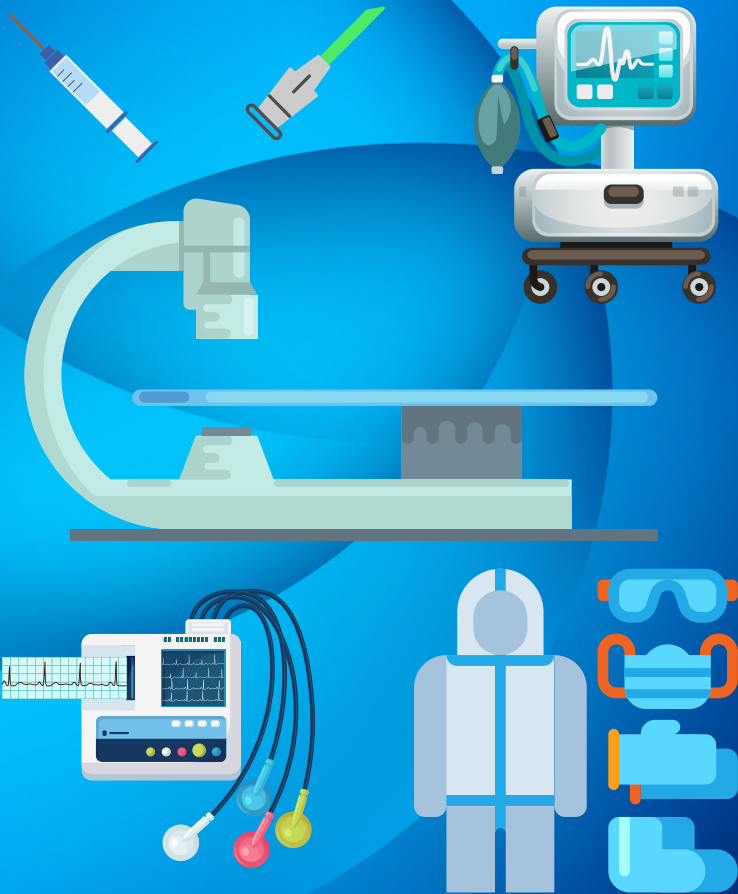


MDR 17



An Interactive Session with Regulator, State Licensing Authority, BIS & NABL

**REGISTER
NOW**



Session Topic

**Ease Of Doing Business
– Industry & Startups**

**Support Session on
Licensing Requirements
- MDR 17**

ORGANISED BY



Association of Indian Medical Industry



NHHID, Anna University



Bureau of Indian Standards
The National Standards Body of India

Date : 23rd July 2022, Sat
Time : 3 pm - 5pm
Meeting : Online

Register Here



FOR ENQUIRIES :

Ms Rama Venugopal , AIMED
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AIMED

AiMeD is an Umbrella Association of Indian Manufacturers of Medical Devices covering all types of Medical Devices including Consumables, Disposables, Equipments, Instruments, Electronics, Diagnostics and Implants.

NHHID, ANNA UNIVERSITY

National Hub for Healthcare Instrumentation Development (NHHID) is established by DST under Technology Development & Transfer in Anna University. This national facility has been formed with collaboration from reputed national institutes and R&D organizations.

BUREAU OF INDIAN STANDARDS

BIS is the National Standard Body of India established under the BIS Act 2016 for the harmonious development of the activities of standardization, marking and quality certification of goods.

NABL

National Accreditation Board for Testing and Calibration Laboratories (NABL) provides accreditation to Conformity Assessment Bodies (Laboratories) in India.



CDSCO

The Central Drugs Standard Control Organisation (CDSCO) under the Directorate General of Health Services, Ministry of Health & Family Welfare, Government of India is the National Regulatory Authority (NRA) of India.

CDSCO along with state regulators, is jointly responsible for grant of licenses of certain specialized categories of critical Drugs such as blood and blood products, I. V. Fluids, Vaccine and Sera.

TNDCA

Drugs Control Department, being statutory body, the organization performs a very important role in supporting healthcare service regulations and enhancing safety of community in Tamilnadu



AIMED
www.aimedindia.com



NHHID
www.nhhid.org



BIS
www.bis.gov.in



PROGRAM DETAILS

Session would be conducted as Online Meeting, an interactive engagement with Manufacturers, Innovators, Startups .

There is **NO REGISTRATION FEE**. However , prior registration is a must.

SESSION TOPIC

Ease of Doing Business
– Industry & Startups Support
Session on Licensing Requirements – MDR 17

DATE OF THE MEETING

Date – 23rd July 2022, Saturday
Time : 3 pm – 5 pm
Meeting Mode : Online

TARGET AUDIENCE

Medical Device Manufacturers, Innovators, Startups in Medical Device, MedTech space, Academia, Technology Business Incubators, Innovation Hubs, Medical Device Testing Laboratories, Healthcare services organisations – Hospitals, Diagnostics Centers , and other healthcare businesses, stakeholders to Conformity Assessment, Government organizations etc

PROGRAM OBJECTIVE

Medical Device Industry in India is brought under full regulatory purview and industry has to show compliance to the Regulatory announcements made to this effect, from time to time.

Operators dealing with Risk Category Class A and B devices come under the purview of State Licensing Authorities and those dealing with Risk Category Class C & D devices come under the purview of Central Licensing Authority.

Medical Device Manufacturers, Innovators, Startups of Tamilnadu have requested for interactive sessions with Central and State Licensing Authorities and other stakeholders like BIS etc to guide them to implement the regulatory requirements better.

AIMED which is the voice for Indian Medical Device Manufacturers in the country and NHHID – Anna University , a National Hub for Healthcare Instrumentation Development, and Bureau of Indian Standards have collectively agreed to organize periodical sessions for the benefit of the industry and startups, innovators of Tamilnadu.

REGISTER HERE



For Enquiries :

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PROGRAM AGENDA



Welcome Note 3 pm – 3 10 pm	Dr M Sasikala, Coordinator - NHHID; Professor, Dept of Electronics & Communication Engineering, Anna University
Contributing to Academia – Innovators – Industry Connect 3 10 pm – 3 20 pm	Mr. Subhash Vashisht Sr. Technical Advisor Medical Electronics, AiMeD
MDR 17 - Licensing Norms for Class A and B Medical Devices - Industry Readiness 3 20 pm - 3 30 pm	Representative of CDSCO
Guiding Industry with National Standards as per MDR 17 licensing requirement 3 30 pm – 3 40 pm	Mr Chandan Kumar, Scientist-C, Medical Equipment and Hospital Planning Department, BIS HQ, New Delhi.
Authentic Quality Certificates - Mandated by MDR 17 3 40 pm - 3 50 pm	Ms Poonam Gupta NABCB Assessor
Product Testing Labs - MDR 17 requirement 3 50 pm - 4 00 pm	Mr. Pankaj Johri, Director National Accreditation Board for Testing and Calibration Laboratories (NABL)
Guiding Industry for smooth transition of Licensing requirements for Class A and B devices – State Licensing Authority, TN 4 00 pm - 4 10 pm	Mr P U Karthigeyan Deputy Director of Drugs Control TN State Drug Control Department
Q & A Session - Moderator 4 10 pm - 5 00 pm	Ms Rama Venugopal Jt Coordinator - AIMED
Vote of Thanks	Dr Sasikala, NHHID

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