

MEDICAL DEVICE REGULATIONS

Legal Metrology (Packaged Commodities) Rules , 2011 on Medical Devices



Interactive Session with Regulators on Legal Metrology Rules for Medical Device Sector



Jointly Organised by



July 26, 2023 - Wednesday
4:00 PM - 5:45 PM IST
Mode - Webinar

For Program Enquiries

CCC Secretariat :

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AiMeD - Association of Indian Medical Device Industry

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 that encourages Responsible Manufacturing. With a Primary Membership of over 400 Manufacturers and additionally of over 300 Associate Members representing the interest of over 1500 Manufacturers of Medical Devices to address the manufacturer's problems.

The aim behind forming AiMeD is allow the Indian Government to access a single point of contact and provide various services to the manufacturers like Advocacy on policy issues, Information services, Regulations for Medical Devices, Education and Training, services, Testing Assistance and guidance for Quality Certification (ISO, CE, GMP), Lobbying for funding for R&D from the Government, encourage innovations from member units, Improve clinician and patient access to the modern, innovative and reliable Medical Device technologies through organizing and supporting Meeting, Seminars, Symposia, Exhibitions and Demonstrations and also, to promote global harmonization and respect for the Indian Device Industry.

AiMeD ensures convergence at a one common platform, when needed, for ensuring overall growth of Indian Medical Devices Industry and further address the needs of all members of the Medical Device Industry irrespective of their affiliation or their company size. It also helps in wider dissemination of information to each player of the industry. AiMeD has been horizontally divided into common groupings like – R&D and Product Development, Legal & Tax Issues, Export Related Issues, Medical Device Regulatory Issues common to all vertical groups of different families of product lines with Zonal Representations.

Association Details

Forum Coordinator

Association of Indian Medical Device Industry (AIMED)

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Consultants Consortium of Chennai

Consultants Consortium of Chennai (CCC) is an Association of Management Systems Consultants, registered under the Tamil nadu Societies Registration Act, 1975.

The Key Objective of CCC is to work in alignment with the National Quality Mission launched by the Government of India by educating the industry about the role and importance of Standards, Certifications and Regulations in the country as well as in the global markets.

CCC strives to encourage Responsible Consulting through Members of its Association and focus is to Standardize Consulting Practices in alignment with changing market and industry needs in a dynamic manner.

The focus is also to identify areas and industries in which the Consultants in the Profession of Management Systems Consulting can contribute to the betterment of the Society and the Country and to promote Management Systems Consulting in such areas or industries.

One of the core objectives is to collaborate with various Industry platforms, Trade Associations, Chambers of Commerce, Management Associations, Apex Bodies of Government or Quasi Government Organization to contribute to the betterment of the industry initiatives, which will in turn contribute to the betterment of the society.

CCC is on a Mission to drive Responsible Consulting in the domain of Management Systems Consulting space.

Association Details

Consultants Consortium of Chennai

“VANITHA”, No 5, Third Avenue, Besant Nagar, Chennai – 600090

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PROGRAM OBJECTIVE



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The Department of Consumer Affairs has recently announced that all Medical Devices must contain information about the Retail Price, the Place of Origin, Size and Dimensions, and Consumer Care Information on the Product Package.

As per the announcement, it was required by the Legal Metrology (Packaged Commodities) Rules, 2011, that every packaged good, including Medical Devices, bear such information on them or on a Label firmly attached to the Device.

The provisions requires declaration of mandatory information including Retail Sales Price in the form of MRP, Country of Origin, Size and Dimensions of the Commodity where the sizes of the commodity in the package are relevant, Unit Sale Price and Consumer Care details, on the pre-packaged commodities based on which the Consumer can take an informed decision.

For various offenses, such as selling above MRP in Hospitals, Retail Locations, or on e-commerce platforms, altering MRP or other declarations/without declarations, etc., enforcement provisions are made under the Legal Metrology Act/Packaged Commodities Rules in order to protect Consumer Rights.

The provisions of the Legal Metrology Act are also applicable on sale of products in loose or without packaging and sale through E-commerce websites in the interest of Consumers.

Hence a session is organised to create an awareness to industry members and stakeholders connected to Medical Device sector about Legal Metrology Rules as well as Compliance to Labeling requirement of similar nature from the Medical Device Rule 2017.

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Contact Co-ordinate

contact@ccc-consultants.org

Opening Remarks

4 00 PM - 4 10 PM

Dr Sanjeev Kumar Gupta
MDQMS Consultant & Regulatory
Affairs Professional - Intrust
Consulting & Forum Member -
Consultants Consortium of Chennai

Legal Metrology (Packaged
Commodities) Rules of 2011
on Medical DevicesImplementation Plan -
Industry Role

4 10 PM - 4 20 PM

Mr Rajiv Nath
Forum Coordinator
AIMED -Association of Indian
Medical Device Industry

Legal Metrology (Packaged
Commodities) Rules of 2011 on
Medical DevicesOverview of the Rules for
Medical Device Sector and
Action Plan by DOCA for
Consumer Protection

4 20 PM - 4 50 PM

Mr Ashutosh Agarwal
Dy Director - Legal Metrology
Department of Consumer Affairs
(DOCA), Ministry of Consumer
Affairs, Govt of India



Mr Devendra Kumar Shakya
Assistant Director, Legal Metrology
RRSL, Ahmedabad

Medical Device Regulations
Regulators Role in Enforcement
of Legal Metrology Rules , to
protect Consumers Interest

4 50 PM - 5 05 PM

Mr Aseem Sahu
Deputy Drugs Controller (I)
CDSCO, North Zone
Ghaziabad



Consumer Protection under LMR

Consumer Viewpoint on the
inclusion of information on the
Medical Device Package

5 05 PM - 5 20 PM

Ms Simi T B
Policy Analyst
CUTS International -
CONSUMER UNITY & TRUST
SOCIETY (CUTS)



Q & A

Session Sum up

Vote of Thanks

5 20 PM - 5 45 PM

Dr Sanjeev Kumar Gupta
CCC Forum

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REGISTRATION DETAILS



Date - July 26, 2023 - Wednesday
Time- 4:00 PM - 5:45 PM IST

Fee :

No Fee. Session open for all.

Who can Register ?

Individuals who use any weight or measure for a transaction or industrial production

Manufacturers , Importers & Retailers, Dealers, Distributors, resellers, stockiest, exhibitors etc. who work with weight, measure or number-related items in Medical Device Sector.

Management Systems Consultants, Regulatory Affairs Professionals , EXIM Consultants, Legal Professionals

Certification Bodies, Inspection Bodies , Testing Labs

Academia – Research & Innovation Institutions in Medical Device space

Startups, Innovators in Medical Device Sector

Registration Link

Click the link below to register as Delegate

https://us02web.zoom.us/webinar/register/WN_bVxiR3NuSBKJWG6WAxie8w#/registration

Contact Coordinate - CCC Secretariat :

Mrs Rama Venugopal

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Share your queries in advance related to Legal Metrology Rules to :

Request Industry Members to share their questions (if any), related to Implementation of Legal Metrology and Packaging Rules to following co-ordinate at AIMED Secretariat :

AIMED Secretariat :

Mr. Chitta Ranjan Biswal, Technical Officer

✉ forumcoordinator@aimedindia.com