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3rd Edition

MEDTECHPOLICY
MAHOTSAV

CONCLAVE

15TH May 2026

CONSTITUTION CLUB OF INDIA, NEW DELHI

Ease of Doing Business to Global Leadership:
Harmonisation, Quality Compliance & Export
Readiness for a Self-Reliant **MedTech India@2047**

SCAN TO REGISTER



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MEDTECH POLICY MAHOTSAV CONCLAVE 2026

15th May 2026, FRIDAY @ CONSTITUTION CLUB OF INDIA, NEW DELHI

3rd Edition of MedTech Policy Mahotsav 2026

Context

India's MedTech sector is entering a decisive decade, driven by expanding healthcare demand, policy push for domestic manufacturing, and global supply chain realignments. Despite strong intent, the sector continues to face regulatory fragmentation, high import dependency, limited alignment with global standards, and export constraints.

Purpose

Policy Mahotsav organized under a **non-profit organization GMAAF (Global Metech Advocacy & Advisory Forum)** is a high-level policy convergence platform bringing together government, regulators, and industry to define actionable reforms.

Key Objectives

- Regulatory simplification and harmonisation
- Strengthening quality compliance frameworks
- Boosting domestic manufacturing and innovation
- Enhancing export readiness
- Institutional convergence

Core Discussion Pillars

- Regulatory Harmonisation** – Streamlining approvals, risk-based frameworks
- Quality Compliance** – Global standards alignment
- Manufacturing & Innovation** – PLI, clusters, R&D
- Export Readiness** – Market access and global positioning

Format of Engagement

- Inaugural Session (Chief Guest Keynote)
- Leadership Roundtables
- Policy Dialogues
- Industry Showcases

Expected Outcomes

- Policy Recommendation Charter
- Priority reform roadmap
- Strengthened public-private coordination

Strategic Significance

The Mahotsav will act as a national policy accelerator, aligning stakeholders and enabling India's transition to a global MedTech leader.

Key Stakeholders Government:

Government: NITI Aayog, MoHFW, CDSCO, DPIIT
Industry: ADMI, AiMeD, CII, FICCI
Healthcare & Research: AIIMS, IITs, Hospitals

Conclusion

This initiative aims to catalyse a future-ready, globally competitive, and self-reliant MedTech ecosystem aligned with Viksit Bharat@2047.

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Session Initiator

Chair



Dr. Rajeev Singh Raghuvanshi
Drugs Controller General of India
CDSCO
Government of India



Dr. G. N. Singh
Former Drugs Controller General,
CDSCO



Jatin Mahajan
Managing Director,
J Mitra & Co Ltd



Shri Aseem Sahu
Deputy Drugs Controller
(India), CDSCO(Hq)



Himanshu Baid
Managing Director
Poly Medicure Ltd.



Rajiv Nath
Forum Coordinator,
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Md. Afzal Kamal
Patron & Mentor
GMAAF



Dr. Jitendra Sharma
MD & Founder CEO
Andhra Pradesh
MedTech Zone Limited,



Dr. Ashoo Grover
Deputy Director General,
Indian Council of Medical
Research



Ganesh Sabat
Deputy Director General,
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Managing Partner



Abhijit Ghosh
Asst. Director
Central Drugs Standard
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Praveen Kumar Mittal
Executive Director
Medical Devices Export
Promotion Council



Dr. Rajiv Chhibber
Communications Specialist &
Joint Forum Coordinator
Association of Indian Medical
Devices



Pavan Choudary
Chairman, MTAL



Dr. Harsh Mahajan
Founder & Chief Radiologist
Mahajan Imaging



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Scientist, Indian Council of
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Prof. Bejon Misra
Founder Director Patient Safety &
Access Initiative of India



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Lifestyle, India & South Asia. Advisor/ Mentor/ Guide



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Member Executive - APEX Chamber of Commerce
& Trade of NCT Delhi.



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Tentative Program

Time	Session Type & Topic
10:00 AM – 10:20 AM	Keynote Address India@2047 & the MedTech Imperative: From Regulatory Reform to Global Manufacturing Leadership
10:20 AM – 11:15 AM	Inaugural Power Panel : MedTech Policy: Aligning Regulation, Industrial Strategy & Export Ambition for Viksit Bharat
11:15 AM – 12:00 PM	Panel Discussion 1 : Regulatory Harmonisation & Ease of Doing Business: Simplifying Licensing, QMS & Compliance under India's MDR Framework
12:00 PM – 12:45 PM	Panel Discussion 2 : Quality Compliance to Global Acceptance: Bridging Indian Standards with USFDA, EU MDR & International Benchmarks
12:45 PM – 01:30 PM	Networking Lunch Break
01:30 PM – 02:15 PM	Panel Discussion 3 : Pricing, Procurement & Ethical Market Access: Balancing Affordability with Innovation in the Medical Device Sector
02:15 PM – 03:00 PM	Panel Discussion 4 : Diagnostics, AI & Emerging Technologies: Regulatory Pathways for IVDs, SaMD & Digital Health Devices
03:00 PM – 03:45 PM	Panel Discussion 5 : Reimagining Price Regulation in Healthcare: From MRP Arbitrage to Transparent, Evidence-Based Value Frameworks
03:45 PM – 04:30 PM	Grand Concluding Policy Dialogue (Panel Discussion 6) : Export Readiness & Global Competitiveness: Policy, Trade Facilitation & Regulatory Reforms for a Self-Reliant MedTech India@2047
5 PM Onwards	CUSTODIAN OF HUMANITY AWRADS

4th Edition
**CUSTODIAN
HUMANITY
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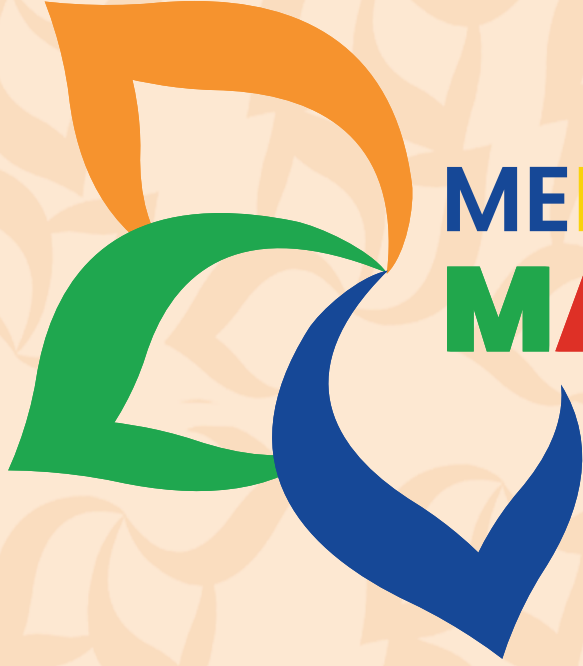
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Participants Profile

- Healthcare Policymakers
- Medical Device Manufacturers
- Hospital Chains
- Startups & Innovators
- Doctors & Medical Associations
- Global Think Tanks
- Trade Bodies & Industry Chambers

Stay Connected | Collaborate | Transform Healthcare

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