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VENTURE CENTER | Social Innovations at Venture Center

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Startup Launch Acceleration Program

Planning the Regulatory Pathway for Medical Devices & Diagnostics

Wednesday | 11 Dec 2024 |Time: 0930 – 1300

Speaker

Chetna Dharmavat
Assistant Manager-Regulatory Services, Venture Center

Speaker

Akash Dhade
Associate Manager-Regulatory Services, Venture Center

What to expect:

- Understanding of USFDA and Indian Regulations to identify the most appropriate regulatory pathway for the medical devices
- Understanding the documentation requirements as per the applicable regulations
- Understanding the ISO 13485 (Quality Management System standard) compliance requirements for medical device manufacturers
- Understanding of the clinical investigation and clinical performance requirements for medical devices

For Whom:

Budding / aspiring entrepreneurs (Researchers, students, engineers, clinicians etc) & Early stage inventive enterprises and science-based startups

Registration Fees: Rs 500/-

REGISTER NOW!

Hybrid Mode (Online via Zoom & In-person at Venture Center)

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