In collaboration with

Report |Awareness Talk | Understanding Medical Device Regulations in the US Market.
|Online Mode| 29th October 2024 |

About The Talk

Awareness talk was organized with Akash Dhade in Online Mode, aimed to equip startups, and innovators with insights into the regulatory landscape of the US market. The talk covered key aspects of the US regulatory process. We received registrations from various audience including startups, academia, and industry professionals keen to gain a understanding of US regulatory requirements for medical devices.

Key Points Discussed in the Talk -

- Regulatory Pathways for US,
- FDA Classification,
- FDA Regulatory Pathway,
- 510(k) Process,
- Substantial Equivalence (SE)
- Case Study

Attendees - 22

Organized By

RIFC

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Overall Feedback: 6/7

"Would like to join the presentation and seminars in the future. Great presentation from Akash." - Rinoy Suvarnadas

"Thank you for the session. It was very informative... Looking forward to such sessions...' - Anuj