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Report | Symposium on Taking Biopharma Products to the Global Market | Friday | 22nd April 2022

About the Event

The program was organized with an intention to introduce Biopharma professionals working in biopharmaceutical development and manufacturing to FDA, EMEA and equivalent international regulatory requirements. It was useful for managers, researchers and for everyone working with Biologics to understand the importance of quality assurance and regulations for biopharmaceuticals.

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Session Description

The event began with a Welcome address & Introduction to Venture Center given by Narendra Chirmule & Dr. Smita Kale. The first talk on "Overview Biopharmaceutical Development Pathway" was conducted by Taruna Arora, followed by a talk on CMC Strategy for Biologic by Jyoti Iyer, & a talk on Key Regulatory Guidelines for the Development by Samir Sangitrao (Zydus Group). The event concluded with Sreekanth Rouduri giving a brief overview of Center for Biopharma Analysis (CBA at Venture Center).
The guest speakers were also a part of the panel discussion in the end on Global Biopharma Regulations.

Feedback Analysis

- No. of Participants- 100+
- Overall Feedback Score- 6.4/7
- Average Social Media Engagement- 3400+ impressions