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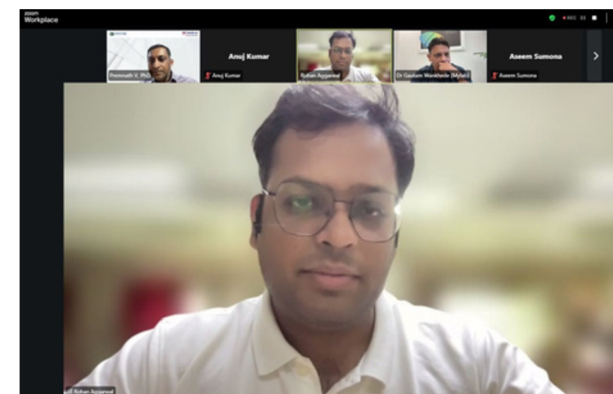
In collaboration with



Report |Mentor Mixer on Best Practices in navigating diagnostics regulatory approvals in India: Insights from Mylab with Dr. Gautam Wankhede | Tuesday | 7th May 2024

About the Event

BRBC's Mentor Mixer was organized on 7th May 2024. Dr. Wankhede's talk that explained Mylab's regulatory journey as a case study was followed by a moderated discussion to highlight some real-time scenarios faced by diagnostics startups (Module Innovations and Vidcare) while securing regulatory approvals for diagnostic products in India.



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Feedback Compilation

7 May 2024 Best Practices in navigating diagnostics regulatory approvals in India : Insights from Mylab			
Evaluation Results			
Category	Avg (Max-Min) Count	Rating Scale	
Section 1 - Event		Rating Scale	
Overall Satisfaction with Event	6.4(5,7)15	1	Bad
Satisfaction with Content	6.4(5,7)15	2	Well Below average
Satisfaction with Speakers/mentors etc	6.6(6,7)15	3	Below Average
Satisfaction with structure,design &pace of event	6.3(5,7)15	4	Average
		5	Good
		6	Very Good
		7	Excellent
Please provide your feedback/suggestions for the workshop			
I was a bit late to join the session due to academic commitment, but the session was arranged so beautifully and gave more insights about various aspects of the Diagnostics regulatory approvals			
For people who are new to this field of startups like me, the session was extremely helpful.			
Informative			
It was a good and insightful session			
Having inputs from a practitioner was very helpful.			
It was a well organized event and the moderator did well in the round table discussion.			
Overall very insightful			
Great Initiative			
Nice			
Excellent session			
aware of the technology during the session.			
Overall good			
Good			
If you have any other suggestions, comments, etc., please share them below.			
It will be beneficial to postgrad students like us if many more sessions about medical devices, IVD's and biologics will be available			
We can also have a session regarding compliances to be done once your product is in the market. 2) Regarding import of certain raw material or components of medical devices			
We should have more sessions where actual practitioners/ entrepreneurs in the field talk about these topics and share personal insights.			
It would be excellent to have a similar event discussing the importance of essential diagnostic list in India that was published in 2019. I am curious to understand why the diagnostic tests are listed as "essential" when they are highly standardized and widely available through vendors. Or is my assumption wrong? It would be good to understand how entrepreneurs could target the essential diagnostic list and develop products that closer to India's national health plan and therefore market needs.			
Want to connect to discuss about funding for a project.			