

BRBC

Regulatory
R Clinic

Report on R Clinic | 22 Feb 2019 | PSG Institute of Medical Sciences & Research, Coimbatore



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BRBC'S R Clinic on "Awareness of the Current Regulatory Requirements for Medical Products and Clinical Trials in India" was organized on 22 Jan 2019 at PSG-STEP, Coimbatore in collaboration with PST-STEP.

Mr. Navnath session was focused on the current medical devices rules from the point of view of the first generation scientific entrepreneur, salient features of medical devices rules and compliance strategies to the rules.

Dr. Ravindra highlighted the clinical investigation of medical devices and performance evaluation of in vitro diagnostics. He spoke on the prior requirements for the clinical studies in India. Pilot and pivotal clinical studies requirements and process flows as per the Medical Devices Rules 2017 and Schedule Y of D&C Act 1940.



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(22 Feb 2019)Awareness of the current Regulatory Requirements for medical products & Clinical trials in India					
Evaluation Results					
Category	Avg (Max-Min) Count				
Section 1 - Event		Rating Scale			
Overall satisfaction With Event	6.1(5,7)23	1	Bad	2	Well below average
Satisfaction with Content	6.0(5,7)23	3	Below average	4	Average
Satisfaction with speakers/mentors etc	6.0(5,7)23	5	Good	6	Very Good
Satisfaction with structure, design & pace of event	6.0(5,7)23	7	Excellent		
Satisfaction with networking opportunities	5.9(5,7)23				
Satisfaction with Event logistics, arrangements	5.9(4,7)23				
Please provide the following information You are a:					
Student	4				
Entrepreneur	4				
Employed	12				
Other	3				
You became know of this event via:					
Email	12				
Social Media	5				
Advt	1				
Other	4				

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Comments and Suggestions

The event was beneficial. It had given a good overview of regulatory bodies.
Very useful & informative. Got the opportunity to connect with an industry expert.
The program is beneficial to clarify regulatory doubts & conflicts.
It was valuable & wish to participate in future research programmer is suggestions to intimate us to mail too.
It would be a one-day event wherein the complete details about regulations for medical devices are explored.
Regulatory requirements shall helpful in detail.

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