



*25 Years of Academic Excellence*

**e- Faculty Development Programme**  
**“Research updates in Pharmaceutical Formulation and Analytical Method Development”**  
**8<sup>th</sup> to 13<sup>th</sup> July 2020**

**Focus area**

- Quality by design approach in Formulation development
- USP RS development and characterization
- New approaches in analytical method development
- Bio analysis in drug discovery and development

**Key features**

- Free registration
- Faculty and Industrial persons can register
- E-Certificate for all registered participants
- Registration deadline: **6<sup>th</sup> July 2020**
- Participate through GPRCP Hyderabad YouTube channel

**For Registration and further details**  
 Visit: [www.gprcp.ac.in](http://www.gprcp.ac.in)

**Contact: 9849044641**  
**9959546008**

<b>Organising Secretaries</b> Prof.Dr.Y.Padmavathi Dept. of Pharmaceutical Analysis Prof.Dr.K.Latha Dept. of Pharmaceutics	<b>Conveners</b> Prof.Dr.B.Madhava Reddy Principal, GPRCP Dr.B.Prabhashankar President, IPA,TS	<b>Organising Committee</b> Dr. D. Prasanthi Mr. P. Ravi Kumar Mr. Shaik Naseeb basha Mrs. K. Pallavi Mr. N. Raghavendra Babu
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Organized by  
**Department of Pharmaceutics & Pharmaceutical Analysis**  
**G. PULLA REDDY COLLEGE OF PHARMACY**  
 In Association with  
**INDIAN PHARMACEUTICAL ASSOCIATION**  
 Telangana state branch




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**Scientific Sessions Schedule**

	<b>Dr. Partha Mukherjee</b> Senior Manager, RSE, (India) USP.	<b>8<sup>th</sup> July 2020 - 10:30 AM to 12:00 PM</b> Topic: <b>Introduction to USP, Reference standard Development and Characterization</b>
	<b>Mrs. A. Swetha sree</b> Analytical scientist Dr.Reddy's Laboratories Ltd, Hyderabad.	<b>9<sup>th</sup> July 2020 - 10:30 AM to 12:00 PM</b> Topic: <b>Role of Analyst in Pharma Industry and New Method Development Approaches</b>
	<b>Dr. N. Badri Vishwanadan</b> Cofounder and CEO M/s Transform SciTech Pvt. Ltd., Hyderabad.	<b>10<sup>th</sup> July 2020 - 10:30 AM to 12:00 PM</b> Topic: <b>Simplifying Qbd</b>
	<b>Dr. V S Kanthikiran</b> Varanasi Head -Preclinical & Bioassay, Global Clinical Management, Dr. Reddy's Laboratories Ltd., Hyderabad.	<b>11<sup>th</sup> July 2020 - 10:30 AM to 12:00 PM</b> Topic: <b>Rapid Bio Analysis in Drug Discovery and Development</b>
	<b>Dr. Vijaykumar Agabandi</b> Deputy Manager, Neuheit Pharma Technologies Pvt. Ltd., Hyderabad.	<b>12<sup>th</sup> July 2020 - 10:30 AM to 12:00 PM</b> Topic: <b>Role of In Vitro-In vivo correlation (IVIVC) in Bioequivalence studies and Bio waiver</b>
	<b>Dr. Santosh Kumar Narla,</b> NAG Manager US Regulatory Affairs, Dr. Reddy's Laboratories Ltd., Hyderabad.	<b>13<sup>th</sup> July 2020 -10:30AM to 12:00 PM</b> Topic: <b>Overview and scope of drug regulatory affairs</b>
	<b>Dr. Venkateswari</b> Muthukrishnan Associate Director, Nuventra Pharma Sciences Durham, North Carolina, USA.	<b>13<sup>th</sup> July 2020 - 03:30 to 05:00 PM</b> Topic: <b>Which Abbreviated Approval Pathway to Choose for A Drug Product? - 505b (2) or ANDA</b>

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**G. PULLA REDDY COLLEGE OF PHARMACY**  
**Mehdipatnam, Hyderabad**  
**ISO 9001 – 2015 Certified College**  
**(Affiliated to Osmania University and Approved by AICTE & PCI)**

**ONE WEEK E-FACULTY DEVELOPMENT PROGRAM ON “RESEARCH UPDATES IN PHARMACEUTICAL FORMULATION AND ANALYTICAL METHOD DEVELOPMENT” IN ASSOCIATION WITH INDIAN PHARMACEUTICAL ASSOCIATION TELANGANA STATE BRANCH**

**(8th – 13th July, 2020)**

One week e- Faculty Development Program (FDP) on “Research updates in Pharmaceutical Formulation and Analytical method development” was jointly organized by Department of Pharmaceutics and Pharmaceutical Analysis, G Pulla Reddy College of Pharmacy, Hyderabad, in

Association with Indian Pharmaceutical Association Telangana state branch from 8th – 13th July 2020.

700 Faculty members across the country, from various institutions and organizations attended the programme. The program was inaugurated with invocation song by Ms. K. Archana Reddy, Asst. Professor, Dept.of pharmacy practice. Organizing secretaries of FDP, Dr. Y. Padmavathi, Professor, Dept.of Pharmaceutical Analysis and Dr.K.Latha Professor, Dept.of Pharmaceutics and Principal Dr. B. Madhava Reddy convenor of the program gave welcome address.

Day1; 8 th July 2020; Dr. Partha Mukherjee, Senior Manager- RSE, USP(India) Pvt ltd, delivered talk on the Topic “Introduction to USP, Reference Standard Development and Characterization”,The session was coordinated by Dr.Y.Padmavathi, Professor, Dept.of Pharmaceutical Analysis.

Day2; 9<sup>th</sup> July 2020; Mrs. A. Swetha Sree, Analytical scientist, Dr.Reddy’s Laboratories Ltd., Hyderabad. spoke on the topic “Role of Analyst in Pharma Industry and New Method Development Approaches”. Mr.P.Ravi Kumar, Asst.Professor, Dept.of Pharmaceutical Analysis, coordinated the program.

Day3; 10<sup>th</sup> July 2020; Dr.N.Badri Vishwanathan, Cofounder and CEO- M/s Transform SciTech Pvt. Ltd., Hyderabad, delivered a talk on the title “Simplifying QbD”.The Session was coordinated by Dr.K. Latha, Professor, Dept.of Pharmaceutics.

Day4;11<sup>th</sup> July 2020; Dr V S Kanthikiran Varanasi, Head -Preclinical & Bioassay, Global Clinical Management, Dr. Reddy’s Laboratories Ltd., Hyderabad, delivered talk on “Rapid Bio Analysis in Drug Discovery and Development”. Dr.Y.Padmavathi, Professor, Dept.of Pharmaceutical Analysis, coordinated the program.

Day5;12<sup>th</sup>July 2020; Dr.Vijaykumar Nagabandi, Deputy Manager-Neuheit Pharma Technologies Pvt. Ltd., Hyderabad. spoke on the topic “Role of In Vitro-In vivo correlation (IVIVC) in Bioequivalence Studies and Biowaiver”. Mr.SK. Nasseb Basha, Asst.Professor, Dept.of Pharmaceutics, coordinated the session.

Day6; 13<sup>th</sup> July 2020; Morning session- 10:30 to 11:30AM; Dr. Santosh Kumar Narla,Deputy Manager, US Regulatory Affairs, Dr. Reddy’s Laboratories Ltd., Hyderabad. delivered a talk on “Overview and Scope of Drug Regulatory Affairs “. The session was coordinated by Dr.D.Prasanthi, Assoc.Professor, Dept.of Pharmaceutics

Day6; 13<sup>th</sup> July 2020, Afternoon Session- 3.30-5.00 PM; Dr. Venkateswari Muthukrishnan Associate Director, Nuventra Pharma Sciences, Durham, North Carolina, USA. delivered a talk on the title “Which Abbreviated Approval Pathway to Choose for A Drug Product? - 505b (2) or ANDA. Dr.K. Latha, Professor, Dept.of Pharmaceutics, coordinated the session.

The six days e-FDP was ended up by valedictory program with vote of thanks proposed by Mrs. K. Pallavi, Asst. Professor, Dept.of Pharmaceutics.

E- Certificated were sent to all the participants.