



## L. M. College of Pharmacy

### Event Report of 'International Conference on Global Landscape of Pharmaceuticals and Biologics: Quality Assurance and Regulations' 2<sup>nd</sup> and 3<sup>rd</sup> January, 2026

1. Name of the event: "International Conference on Global Landscape of Pharmaceuticals and Biologics: Quality Assurance and Regulations"
2. Dates of the event: 02/01/2026 and 03/01/2026
3. Number of participants (also geographical locations indicative of the reach of the event):  
220

Venue: J. C. Bose Auditorium, Department of Zoology, Gujarat University, Navrangpura, Ahmedabad.

4. Name of collaborating agency/ Agency providing financial assistance for the conduct of the event.:

**Agency providing financial assistance:** Gujarat State Biotechnology Mission (GSBTM), Gujarat Council on Science & Technology (GUJCOST), Department of Science and Technology (DST).

**Knowledge Partners:** AIC-LMCP Foundation and Indian Pharmaceutical Association-Gujarat State Branch (IPA-GSB).

**Collaborating agencies:** Kesar Control systems, Ahmedabad; Intervein Research Labs, Ahmedabad; Auxilla Pharmaceuticals and Research LLP, Ahmedabad; SOTAX India (Pvt) Limited; SPINCO and Grey Matters.

5. Details of the Chief Guest, Guest(s) of Honour during the Inaugural and Valedictory functions.:

**Inaugural Function:**

Chief Guest: Dr. S. P. Adeshara, Former Commissioner, FDCA, Gujarat.

Guest of Honour: Mr. Mehul Patel, Director, Kesar Control Systems.

**Valedictory Function:**

Chief Guest: Dr. Hemant Koshia, Former Commissioner, FDCA, Gujarat.

Guest of Honour: Shri Gaurang Oza, Managing Partner, Vaibhav Analytical Services.

6. Details of the Speakers/ Resource Persons along with their affiliations and the title of their talks.

Sr. No.	Details of the Speakers/ Resource Persons	Affiliations	Title of the talk
<b>Track I: Pharmaceuticals</b>			
1	<b>Dr. Murlidhar Zope</b>	Head & Associate VP, Analytical R&D for Complex generics and Specialty Pharmaceutical Products, Sun Pharmaceutical Industries Ltd	Characterization and In-Vitro Bioequivalence for Complex Pharmaceuticals
2	<b>Dr. Viranchi Shah</b>	Saga Laboratories, Ex-President IDMA	Quality & Compliance – Navigating Some Recent Developments
3	<b>Mr. Kiran Kolhe</b>	VP-Regulatory Affairs, Amneal Pharmaceuticals	Regulatory Considerations for Complex Generic Drug Products – Peptides
4	<b>Ms. Sampada Walimbe</b>	IT Quality Lead, Abbott Healthcare	Emerging Trends in Software Quality Assurance for the Life Sciences Industry
5	<b>Mr. Vipul Doshi</b>	Chief Quality & Compliance Officer, Zydus Lifesciences Limited	Cost of Quality with Business Perspective
<b>Track II: Biologics and Biosimilars</b>			
6	<b>Dr. Samir Sangitrao</b>	Vice President and Head of Regulatory Affairs and R&D Quality Assurance – Biologics, Zydus Lifesciences Limited	QMS: Levelling Up Biologics & Biosimilars Excellence
7	<b>Dr. Ravishankara MN</b>	Senior General Manager, Sun Pharmaceutical Industries Limited	From Characterization to Comparability: Challenges in Biosimilar Analytical Assessment
8	<b>Dr. Sudheendra Kulkarni</b>	AVP – Regulatory Policy, Government Affairs & Audit Liaison, Biocon Biologics Limited.	Trends in Biosimilars: Development and Adoption in an Evolving Regulatory Environment
9	<b>Dr. Ketan Patel</b>	Professor of Pharm Sciences, St. John's University, NY, USA.	RRR of next generation biologics: Revolution, Research and Regulation

10	<b>Dr. Lalitkumar Vora</b>	Lecturer in Pharmaceutical Sciences, School of Pharmacy, Queen's University, Belfast.	When medicines meet medical devices: Regulatory framework and perspectives
<b>Panel Discussion</b>			
<b>Theme: Emerging Quality and Regulatory Trends Transforming Clinical Development</b>			
	<b>Dr. Shrinivas Savale</b>	Moderator	CEO, AIC – LMCP.
	<b>Dr. Kiran Marthak</b>	Panelist	Director -Regulatory Affairs & Phase 1 studies, Veeda Lifesciences.
	<b>Mr. Akshaya Nath</b>	Panelist	Executive Vice President, Intas Pharmaceuticals.
	<b>Dr. Dharmesh Domadia</b>	Panelist	VP Clinical Research, Cliantha.
	<b>Dr. Pawan Singh</b>	Panelist	Vice President and Head of Clinical Development, Kashiv Biosciences Pvt. Ltd.
<b>Track III: Herbal Medicines and Medical Devices</b>			
11	<b>Mr. Saurabh Rawat</b>	Head, Quality and Regulatory Affairs, Axio Biosolutions Pvt. Ltd.	Effective Quality Management Systems for Medical Devices in a Changing Regulatory Landscape
12	<b>Dr. Akshay Srivastava</b>	Head, Department of Medical Devices and In-charge R&D for Center of Excellence in Medical Devices, NIPER - Ahmedabad	Regulatory perspective in Medical Devices: Current challenges and future directions
13	<b>Dr. Pravin Nalawade</b>	PHS Lifesciences Pvt Ltd.	Global Opportunities and Quality Paradigms in the Nutraceutical Industry: An Indian Perspective
14	<b>Dr. Nidhi Sapkal</b>	Professor, Guru Nanak College of Pharmacy	Building Regulatory Strategy for Herbal Medicines and Nutraceuticals: Case Studies

7. A brief description of the outcomes achieved through the event or the message from the event.

L. M. College of Pharmacy (LMCP), Ahmedabad hosted the International Conference on “Global Landscape of Pharmaceuticals and Biologics: Quality Assurance and Regulations” on 2–3 January 2026. This conference was organised to mark about 50 years of PG education and 30 years of PG education in Pharmaceutical Quality Assurance in particular. The conference witnessed participation from over 250 delegates representing students, faculty members and industry professionals from India and abroad. The conference was structured into thematic tracks covering Pharmaceuticals, Biologics and Biosimilars, Herbal Medicines and Medical Devices, with a dedicated panel discussion on clinical development and its regulations.

The conference provided a comprehensive platform for academicians, researchers, regulators, and industry professionals to deliberate on emerging trends, challenges, and innovations shaping quality systems and regulatory frameworks in the life sciences sector. The programme emphasized key contemporary themes such as global regulatory harmonization, current Good Manufacturing Practices (cGMP), quality risk management, and advances in analytical and manufacturing technologies, reflecting the growing need for alignment between scientific innovation and compliance-driven development.

The conference was structured into thematic tracks covering Pharmaceuticals, Biologics and Biosimilars, Herbal Medicines and Nutraceuticals, Medical Devices, and Intelligent Manufacturing with AI/ML applications. A major highlight of the conference was the extensive poster presentation segment. In total, 110 poster presentations were presented out of 124 accepted for presentation, demonstrating strong research engagement and practical relevance. These included 56 posters in Track I (Emerging Techniques for Quality Control of Pharmaceuticals and Biologics), 31 posters in Track II (Changing Paradigms in Quality Assurance of Herbals, Cosmetics and Nutraceuticals), 23 posters in Track III (Evolving Landscape in Quality Aspects of Medical Devices), and 14 posters in Track IV (Intelligent Manufacturing and Quality Control: AI/ML for Next-Gen Pharma and Biologics). The poster sessions showcased applied and compliance-oriented research across areas such as analytical method development, stability studies, regulatory science, herbal standardization, medical device quality, and Pharma 4.0 initiatives, reinforcing the integration of innovation with regulatory expectations and real-world quality challenges.

The expert sessions collectively offered an integrated perspective on how science, quality systems, and regulation interact across modern product development. Dr. Viranchi Shah highlighted the evolving regulatory landscape and the need for proactive compliance supported by continuous improvement, emphasizing that quality compliance today must remain current, resilient, and responsive. Mr. Vipul Doshi provided a strong business perspective through “Cost of Quality,” demonstrating how quality-related costs influence operational efficiency and profitability, and how prevention-focused systems strengthen sustainable performance without compromising compliance. A science-driven focus was

evident in sessions addressing complex products and analytical evidence. Dr. Murlidhar Zope emphasized scientific rigor and regulatory relevance in establishing bioequivalence for complex pharmaceuticals, stressing advanced characterization and in-vitro methodologies aligned with global regulatory expectations. Complementing this, Mr. Kiran Kohle addressed regulatory considerations for peptide-based complex generics, focusing on challenges in characterization and quality attributes, and outlining guideline-based strategies for successful development and submission.

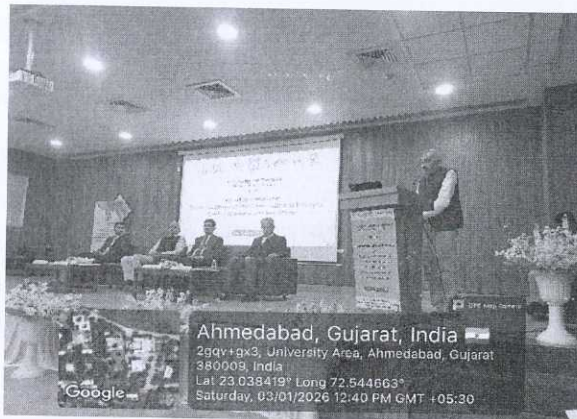
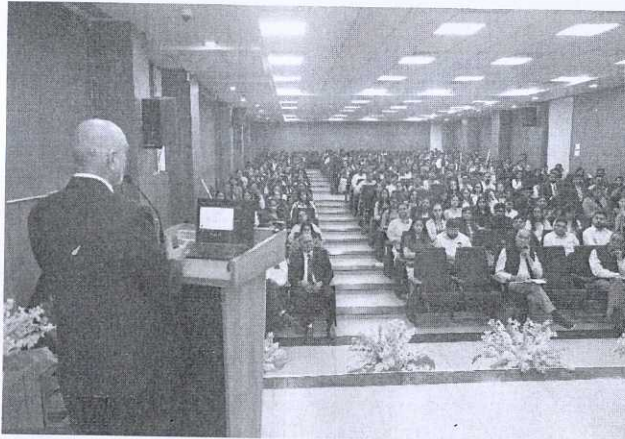
In the biologics and biosimilars domain, Dr. Sameer Sangitrao discussed strengthening QMS maturity using best practices and risk-based approaches to ensure consistency and compliance. Dr. Ravishankar M.N highlighted challenges in biosimilar analytical comparability and the need for robust analytical strategies amid evolving regulations. Dr. Sudheendra Kulkarni presented global trends in biosimilar development and adoption, while Dr. Ketan Patel discussed innovation and regulatory readiness in next-generation biologics. Sessions also covered regulatory coordination for combination products and medical devices, with Dr. Lalitkumar Vora, Mr. Saurabh Rawat, and Dr. Akshay Srivastava addressing pathways, QMS strengthening, and future regulatory directions. Ms. Sampada Walimbe highlighted emerging trends in software quality assurance, and Dr. Pravin Nalawade and Dr. Nidhi Sapkal addressed global opportunities and regulatory strategy for nutraceuticals and herbal medicines.

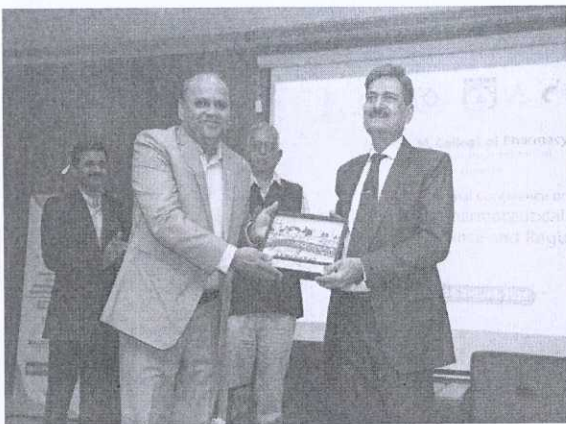
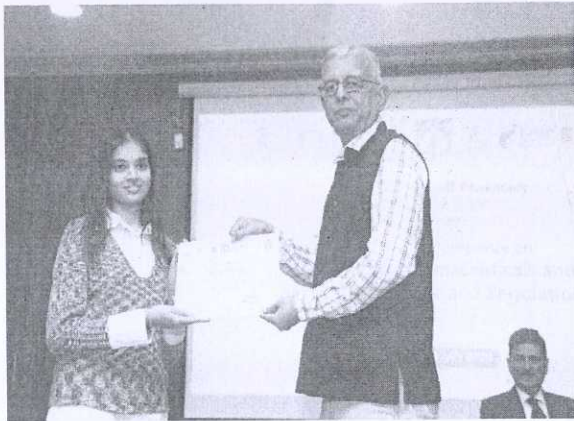
In addition, the conference included a felicitation ceremony to honour the founding faculty members of the institution, namely Prof. C. G. Shishoo, Prof. S. A. Shah, Dr. B. N. Suhagiya, Shri I. S. Rathod, Dr. R. K. Parikh, Dr. G. K. Jani, Dr. Dushyant Shah, and Shri S. K. Vyas, in recognition of their foundational contributions.

Overall, the conference successfully strengthened participants' understanding of evolving quality and regulatory paradigms, while promoting research dissemination and practical, compliance-aligned innovation through expert deliberations and a robust poster platform.

#### 8. A few representative photographs







**List of Annexures:**

- Annexure I: List of Participants
- Annexure II: Brochure
- Annexure III: Detailed schedule of the event
- Annexure IV: News Clippings

**10. Signatures:**

**Prepared By**

**Dr. Mansi N. Athalye**

**Name of the Organizing Secretary**

**Dr. Shrinivas S. Savale**

**Name of the Convener**

**Dr. M. T. Chhabria**

**International Conference on "Global Landscape of Pharmaceuticals and Biologics: Quality Assurance and Regulations**  
**2-3 January 2026**

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 GOPALBHAI,  
 DHOKIA HARSH  
 JAGDISH,

Utsav  
 6-1-11  
 received

Prakash

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




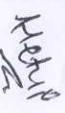
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LMCP/79	Mr., MEETRAJSINH, DODIYA, JITENDRASINH	Student	₹1,000	Institute of Pharmacy Nirma University	919574309696	meetraisindhodiya73 @gmail.com	Shah Vanshika Niraj	Received
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LMCP/82	Ms., Nikita, Diwakar, Rajkishor	Student	₹1,000	A. R. College of Pharmacy and G. H. Patel Institute of Pharmacy, Anand-388120	916358821264	nikitradiwakar57@gm ail.com		Received
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LMCP/86	Mr., MILIN, PATEL, GIRISHBHAI	Student	₹1,000	Dharmsinh Desai University	919974922808	patelmilin24@gmail.com		
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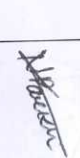



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LMCP/133	Ms., Mitwa, Patel, Rakeshkumar	Student	₹1,200		919601618106	mitwapatel1810@gm ail.com		<i>(Signature)</i>
LMCP/134	Ms., Trisha, Sarkar, Hemant	Student	₹1,200	SVKM's NMIMS Shobhaben Prataphbai College oh Pharmacy and Technology	918356080347	trishasarkar257@gm ail.com		<i>(Signature)</i> 02/12/25
LMCP/135	Ms., Suhani, Pandya, Paragkumar	Student	₹1,200	Shobhaben Prataphbai Patel School of Pharmacy & Technology Management	9199408761829	suhani26pandya@gm ail.com		<i>(Signature)</i> 21/12/25
LMCP/136	Ms., Patel, Jahanvi, Jigar	Student	₹1,200	Institute of Pharmacy, Nirma University	919913377051	20ptphdp127@nirma uni.ac.in		<i>(Signature)</i> 21/12/25
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LMCP/138	Mr., Jannay, Patel, Ashish	Student	₹6,000	KBIPER (KSV UNIVERSITY)	918980010106	jannaypatel4@gmail. com	Jannay Patel, Krupa Aghav, Khushi Babariya, Nancy Panchal, Vidhi Patel, Yashvi Dodiya	<i>(Signature)</i>

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LMCP/141	Ms., Miss, Nikita, Agarwal	Student	₹1,200	School of Pharmacy and Technology Management, SVKM's	919602657901	na788936@gmail.co m	
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			y	Kesar Control System				
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	Dr. Shyam Sunder			DMISH	7023459497			<i>[Signature]</i>
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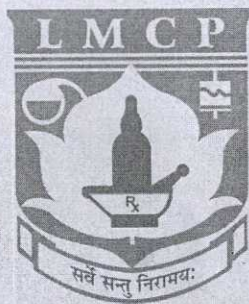
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L. M. College of Pharmacy  
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**International Conference on  
Global Landscape of  
Pharmaceuticals and Biologics:  
Quality Assurance and Regulations**



2-3 January

2026



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## About L. M. College of Pharmacy

L. M. College of Pharmacy (LMCP) is a premier Pharmacy institute of India that was established in 1947 with a vision to impart quality Pharmacy education. The institute is managed by Ahmedabad Education Society, a philanthropic trust established in 1935. The institute offers D. Pharm., B. Pharm., Pharm. D., M. Pharm. and Ph.D. programs; all approved by the Pharmacy Council of India, New Delhi, Govt. of India. The institute is accredited by NAAC with "A" grade and has been conferred Autonomous status for a period of ten years from 2024-25 academic year. LMCP is the only Pharmacy college in the country to get financial assistance from NITI Aayog's Atal Innovation Mission (AIM) to set up the 'Atal Incubation Centre'-AIC-LMCP Foundation (ALF) that focuses on supporting innovators in Pharmaceutical and Healthcare sector. With the support of highly qualified and experienced faculty members, a great number of industrialists, researchers and academicians have emerged from the institute, contributing significantly to Pharmaceutical and Healthcare sector in India as well as on the global horizon.

## About the Conference

The **International Conference on 'Global Landscape of Pharmaceuticals and Biologics: Quality Assurance and Regulations'** aims to bring together leading experts, academicians, researchers, regulators, and industry professionals to share knowledge about the evolving global landscape of pharmaceuticals, biologics, and medical technologies in the context of Quality and Regulations. This event will provide an opportunity for exchanging knowledge, presenting policies, and bringing professionals together between industry and academia, with a focus on quality assurance, regulatory alignment, and innovation. Participants will gain knowledge about international standards, new rules and regulations, approaches to ensuring quality, and the latest developments in the pharmaceutical, herbal, biologic, and medical device industries. The conference is organized into thematic tracks to facilitate cross-disciplinary discussions among participants from different fields.



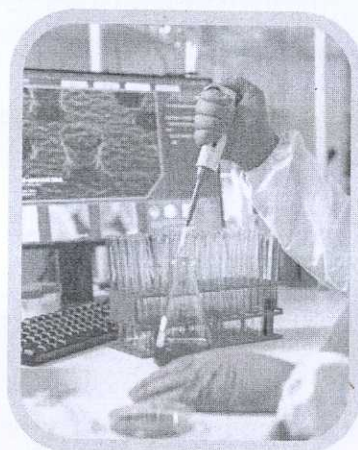
*Shiv*

# Conference Tracks

## Track I: Pharmaceuticals

This track integrates industrial pharmacy, and regulatory science to address:

- Harmonization of global regulations concerning pharmaceutical formulations
- Compliance with Current Good Manufacturing Practices (CGMP), and relevant pharmacopoeial standards for Active Pharmaceutical Ingredients (APIs), and excipients
- Quality risk management in pharmaceutical products with focus on assessing safety, efficacy, and validation processes in the manufacturing of pharmaceuticals



## Track II: Biologics and Biosimilars

Biologics and biosimilars are going to shape the future of therapeutics. The primary areas of focus are as follows:



- Regulatory pathways for biologics and biosimilars including the USFDA, EMA, WHO, CDSCO, and ICH
- Characterization and comparability exercises utilizing advanced analytical tools
- Assessment of immunogenicity and monitoring of safety
- Scale-up and process development for manufacturing of biosimilars
- Management of cold chain logistics and advancements in formulation techniques
- Analysis of patent landscape and challenges

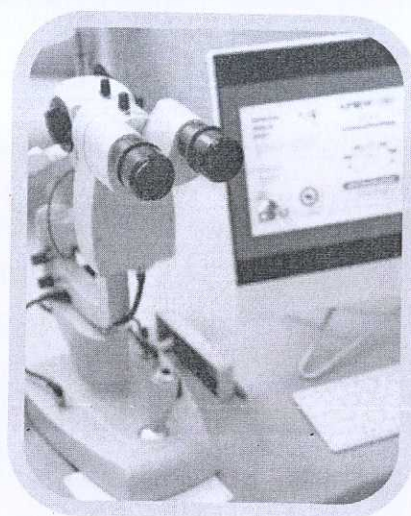


## Track III: Herbal Medicines and Medical Devices

This track addresses the expanding field of herbal medicines, medical devices, diagnostics, implants, and combination products and also promotes collaboration among device manufacturers, pharmaceutical innovators, quality professionals, and regulatory authorities to achieve patient-centric and technology-driven solutions.

Key aspects of the track are:

- Standardization and authentication guidelines for herbal formulations
- Quality risk management in herbal products with focus on assessing safety, efficacy, and validation processes in the manufacturing of nutraceuticals and phytopharmaceuticals
- Following aspects of Medical Devices/ Diagnostics/ Drug Device Combination
  - (i) Design controls and lifecycle quality management
  - (ii) Processes for risk classification and certification
  - (iii) Clinical evaluation, validation, and design engineering
  - (iv) Standards for biocompatibility testing and sterilization
  - (v) Combination products and drug delivery devices
  - (vi) Post-market surveillance and digital integration



## Registration Details

Category	Registration fees* up to 15/12/2025	Registration fees* After 15/12/2025
UG and PG students	1000/-	1200/-
Academicians	1500/-	1800/-
Working Professionals from Industry	2000/-	2500/-

\*Inclusive of GST; includes access to all conference sessions and hospitality

Link for registration form:

<https://zfrmz.com/LliMK4ISTv5ugVxzNFDo>

OR

Scan the QR code



*Signature*

## Poster Presentation

The Organising Committee invites delegates to take part in focused, productive discussions with Subject Matter Experts during the dedicated poster sessions. Delegates are encouraged to submit abstracts for poster presentation and engage in meaningful conversations with experts, get constructive feedback, and start collaborations.

The goal of the Poster Session is to bring forth innovative research, advanced technologies, and modern practices that enhance quality assurance and regulatory standards in the fields of pharmaceuticals, biologics, herbals and medical devices.

## Key Thrust Areas

### 1. Emerging Techniques for Quality Control of Pharmaceuticals and Biologics

This track is focused on advanced analytical tools and quality systems that improve safety, accuracy, and compliance in drug development and manufacturing processes. Posters may centre around the following topics:

- Advanced chromatographic and spectroscopic techniques
- Process Analytical Technology (PAT) and real-time release testing
- Rapid microbiological methods and endotoxin testing
- Bioassays, potency determination, and stability evaluation
- Automation and high-throughput analytical platforms
- Regulatory expectations for QC of biologics, vaccines, and complex generics

### 2. Changing Paradigms in Quality Assurance of Herbals, Cosmetics and Nutraceuticals

This track encourages translational approaches combining traditional knowledge, innovation, and regulatory discipline.

Posters could be focused on:

- Standardization and phytochemical profiling of herbal formulations
- Safety and efficacy validation of botanicals and nutraceuticals
- GxP and regulatory compliance for herbal and cosmetic industries
- Quality risks, adulteration control, and contaminant testing
- Integrating AYUSH practices with scientific evidence and quality assurance

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### 3. Evolving Landscape in Quality Aspects of Medical Devices

This track emphasizes quality systems, safety standards, and regulatory developments that are influencing the medical device sector.

Submissions for posters are encouraged on:

- Regulatory frameworks for medical devices including ISO, EU MDR, USFDA, and CDSCO
- Risk management and lifecycle documentation
- Requirements for biocompatibility testing and sterilization
- Quality systems for diagnostics, implants, and combination products
- Digital health, wearable devices, and cybersecurity considerations

### 4. Intelligent Manufacturing and Quality Control: AI/ML for Next-Gen Pharma and Biologics

This track encourages forward-thinking strategies that will improve efficiency, ensure compliance, and maintain quality by using advanced technologies in manufacturing processes, quality control measures, and regulatory compliance frameworks.

Participants are welcome to address following topics in the poster:

- Optimization of processes and predictive maintenance driven by AI and ML
- Smart sensors, and automation in the context of Pharma 4.0
- Real-time data analytics and quality management system digitization
- Utilization of robotics in bioprocessing and fill-finish operations
- AI applications for deviation analysis, batch release processes, and regulatory intelligence management
- Industry 4.0 and continuous manufacturing methodologies

#### Abstract submission:

#### Link for Abstract submission:

<https://forms.gle/Rt9SUR4t4APBSDHh9>

**Last date for Abstract submission: 15/12/2025**

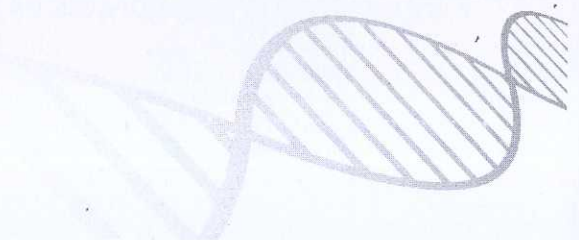
**Intimation of Acceptance: 22/12/2025**

For any query write to:



[conference@lmcp.ac.in](mailto:conference@lmcp.ac.in)

*Sh*



## Speakers

Dr. Sudheendra Kulkarni (Biocon Biologics)  
Mr. Kiran Kolhe (Amneal Pharmaceuticals)  
Dr. Ravishankara M. N. (Sun Pharmaceuticals)  
Dr. Akshay Srivastava (NIPER, Ahmedabad)  
Mr. Vipul Doshi (Zydus Lifesciences)

Mr. Akshaya Nath (Intas Pharmaceuticals)  
Dr. Lal Hingorani (Pharmanza Herbal)  
Ms. Sampada Walimbe (Abbott Healthcare)  
Dr. Ketan Patel (St. John's University, NY)  
**and MORE**

## Organising Committee

### Patron:

Dr. Shriram Nerlekar

### Chairman:

Dr. Mahesh T. Chhabria

### Organising Secretary:

Dr. Shrinivas S. Savale

### Members:

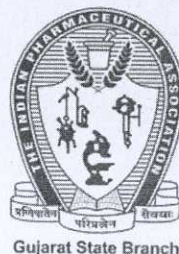
Dr. Gaurang B. Shah  
Dr. Anuradha K. Gajjar  
Dr. Manju Misra  
Dr. J. B. Dave

Mr. Gopal K. Joshi  
Dr. Ketan M. Ranch  
Dr. Devang B. Sheth  
Dr. Krupa V. Gadhvi

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*Shr*



Department of Science & Technology  
Government of Gujarat



Gujarat State Branch



We cordially invite you to the  
***Inaugural Function***  
of  
***International Conference on  
Global Landscape of  
Pharmaceuticals & Biologics:  
Quality Assurance and Regulations***  
on  
***2<sup>nd</sup> January 2026, Friday, 9:30 a.m.***

**Chief Guest**  
**Dr. S. P. Adeshara**  
Former Commissioner, FDCA, Gujarat

**Guest of Honour**  
**Mr. Mehul Patel**  
Director, Kesar Control Systems  
have kindly consented to grace the occasion.

**Venue: Dr. J. C. Bose Auditorium,  
Gujarat University, Navrangpura, Ahmedabad - 380009**

**Patron: Dr. Shriram Nerlekar, Director,  
Ahmedabad Education Society**  
**Convener: Dr. Mahesh T. Chhabria, Principal,  
L. M. College of Pharmacy**  
**Organizing Secretary: Dr. Shrinivas S. Savale, CEO,  
AIC - LMCP Foundation**



We cordially invite you to the

***Valedictory Function***

of

***International Conference on***

***Global Landscape of***

***Pharmaceuticals & Biologics:***

***Quality Assurance and Regulations***

on

***3<sup>rd</sup> January 2026, Saturday, 11:45 a.m.***

**Chief Guest\***

**Dr. Hemant Koshia**

**Former Commissioner, FDCA, Gujarat**

**Guest of Honour**

**Shri Gaurang Oza**

**Managing Partner, Vaibhav Analytical Services**

**have kindly consented to grace the occasion.**

***Venue: Dr. J. C. Bose Auditorium,***

***Gujarat University, Navrangpura, Ahmedabad - 380009***



**Patron: Dr. Shriram Nerlekar, Director,  
Ahmedabad Education Society**  
**Convener: Dr. Mahesh T. Chhabria, Principal,  
L. M. College of Pharmacy**

**Organizing Secretary: Dr. Shrinivas S. Savale, CEO,  
AIC - LMCP Foundation**





**International Conference on Global Landscape of  
Pharmaceuticals and Biologics: Quality Assurance and Regulations  
(2<sup>nd</sup> and 3<sup>rd</sup> January 2026)**

ORGANIZED BY

**L. M. College of Pharmacy  
(An Autonomous Institution)**

<b>Venue: J. C. Bose Auditorium, Department of Zoology, Gujarat University, Navrangpura, Ahmedabad</b>		
<b>Day-1, Friday 2<sup>nd</sup> January 2026</b>		
8:30 am – 9:30 am	<b>Registration and Breakfast</b>	
9:30 am – 10:15 am	<b>Inauguration ceremony</b>	
<b>Track I: Pharmaceuticals</b>		
<b>Time</b>	<b>Speaker &amp; Affiliation</b>	<b>Title of talk</b>
<b>Plenary Session</b>		
10.30 am to 11:10 am	<b>Dr. Murlidhar Zope</b> Head & Associate VP, Analytical R&D for Complex generics and Specialty Pharmaceutical Products, Sun Pharmaceutical Industries Ltd	Characterization and In-Vitro Bioequivalence for Complex Pharmaceuticals
<b>Scientific session</b>		
11:10 am to 11:35 pm	<b>Dr. Viranchi Shah</b> Saga Laboratories, Ex-President IDMA	Quality & Compliance – Navigating Some Recent Developments
11:35 am to 12:00 noon	<b>Mr. Kiran Kolhe</b> VP-Regulatory Affairs, Amneal Pharmaceuticals	Regulatory Considerations for Complex Generic Drug Products – Peptides
12: 00 noon to 12:25 pm	<b>Ms. Sampada Walimbe</b> IT Quality Lead, Abbott Healthcare	Emerging Trends in Software Quality Assurance for the Life Sciences Industry
12:25 pm to 12:50 pm	<b>Mr. Vipul Doshi</b> Chief Quality & Compliance Officer, Zydus Lifesciences Limited	Cost of Quality with Business Perspective
1:00 pm – 2:00 pm	<b>Lunch</b>	
12.45 pm-2.15 pm	<b>Poster Session (Track II and III)</b>	<b>Venue: J. C. Bose Auditorium Lawn</b>

<b>Track II: Biologics and Biosimilars</b>		
Time	Speaker & Affiliation	Title of talk
<b>Plenary Session</b>		
2:00 pm – 2:40 pm	<b>Dr. Samir Sangitrao</b> Vice President and Head of Regulatory Affairs and R&D Quality Assurance – Biologics, Zydus Lifesciences Limited	QMS: Levelling Up Biologics & Biosimilars Excellence
<b>Scientific session</b>		
2:40 pm to 3:05 pm	<b>Dr. Ravishankara MN</b> Senior General Manager, Sun Pharmaceutical Industries Limited	From Characterization to Comparability: Challenges in Biosimilar Analytical Assessment
3:05 pm to 3:30 pm	<b>Dr. Sudheendra Kulkarni</b> AVP – Regulatory Policy, Government Affairs & Audit Liaison, Biocon Biologics Limited	Trends in Biosimilars: Development and Adoption in an Evolving Regulatory Environment
3:30 pm to 3:55 pm	<b>Dr. Ketan Patel</b> Professor of Pharm Sciences, St. John's University, NY, USA.	RRR of next generation biologics: Revolution, Research and Regulation
3:55 pm to 4: 20 pm	<b>Dr. Lalitkumar Vora</b> Lecturer in Pharmaceutical Sciences, School of Pharmacy, Queen's University, Belfast	When medicines meet medical devices: Regulatory framework and perspectives
4:30 pm – 5:00 pm	<b>Tea Break</b>	
<b>Panel Discussion</b>		
<b>Theme: Emerging Quality and Regulatory Trends Transforming Clinical Development</b>		
5:00 pm to 6:00 pm	<b>Dr. Shrinivas Savale</b> (Moderator) CEO, AIC - LMCP	
	<b>Dr. Kiran Marthak</b> Director -Regulatory Affairs & Phase 1 studies, Veeda Lifesciences.	
	<b>Mr. Akshaya Nath</b> Executive Vice President, Intas Pharmaceuticals.	
	<b>Dr. Dharmesh Domadia</b> VP Clinical Research, Cliantha.	
	<b>Dr. Pawan Singh</b> Vice President and Head of Clinical Development, Kashiv Biosciences Pvt. Ltd.	
6:00 pm – 7:30 pm	Poster Session (Track I and IV)	Venue: J. C. Bose Auditorium Lawn
7:30 pm – 8:30 pm	<b>Dinner</b>	

<b>Day 2, Saturday 3<sup>rd</sup> January 2026</b>		
<b>Track III: Herbal Medicines and Medical Devices</b>		
8:30 am – 9:30 am	<b>Breakfast</b>	
<b>Time</b>	<b>Speaker &amp; Affiliation</b>	<b>Title of Talk</b>
<b>Plenary Session</b>		
9:30 am – 10:10 am	<b>Mr. Saurabh Rawat</b> Head, Quality and Regulatory Affairs, Axio Biosolutions Pvt. Ltd	Effective Quality Management Systems for Medical Devices in a Changing Regulatory Landscape
<b>Scientific session</b>		
10:10 am to 10:35 am	<b>Dr. Akshay Srivastava</b> Head, Department of Medical Devices and In-charge R&D for Center of Excellence in Medical Devices, NIPER -Ahmedabad	Regulatory perspective in Medical Devices: Current challenges and future directions
10:35 am to 11:00 am	<b>Dr. Pravin Nalawade</b> PHS Lifesciences Pvt Ltd.	Global Opportunities and Quality Paradigms in the Nutraceutical Industry: An Indian Perspective
11:00 am to 12:00 noon	<b>Dr. Nidhi Sapkal</b> Professor, Guru Nanak College of Pharmacy	Building Regulatory Strategy for Herbal Medicines and Nutraceuticals: Case Studies
12:15 pm – 1:00 pm	<b>Valedictory function</b>	
1:00 pm – 2:00 pm	<b>Lunch</b>	

# LM ફાર્મસીમાં આજથી ઈન્ટરનેશનલ કોન્ફરન્સ

એલ.એમ. કોલેજ ઓફ ફાર્મસી દ્વારા 'ગ્લોબલ લેન્ડસ્કેપ ઓફ ફાર્માસ્યુટિકલ્સ એન્ડ બાયોલોજિક્સ: ક્વૉલિટી અશ્યોરન્સ એન્ડ રેગ્યુલેશન્સ' વિષય પર આજથી બે દિવસીય ઈન્ટરનેશનલ કોન્ફરન્સનું આયોજન કરાયું છે. ઈન્ટરનેશનલ કોન્ફરન્સમાં ફાર્માસ્યુટિકલ્સ, બાયોસિમિલર્સ, હર્બલ મેડિસિન્સ અને ન્યુટ્રાસ્યુટિકલ્સ, મેડિકલ ડિવાઈસીસ તેમજ એઆઈ-એમએલ આધારિત ઈન્ટેલિજન્સ મેન્યુફેક્ચરિંગ વિશે તજજ્ઞો દ્વારા માહિતી આપવામાં આવશે.



## એલ.એમ.ફાર્મસીમાં આંતરરાષ્ટ્રીય પરિસંવાદ



એલ.એમ. કોલેજ ઓફ ફાર્મસીમાં 'ફાર્માસ્યુટિકલ્સ અને બાયોલોજિક્સનું વૈશ્વિક પરિદ્રશ્ય: ગુણવત્તા ખાતરી અને નિયમન' વિષય પર આંતરરાષ્ટ્રીય પરિસંવાદ યોજાયો હતો, જેમાં પ્રોફેસર મહેશ છાબરિયા, ગુજરાત ફૂડ એન્ડ ડ્રગ કન્ટ્રોલ એડમિનિસ્ટ્રેશનના નિવૃત્ત કમિશનરો ડૉ. સુબોધ આદેશરા અને ડૉ. હેમંત કોશિયા, બાયોકોન બાયોલોજિક્સના ડૉ. સુધીન્દ્ર કુલકર્ણી, ન્યૂયોર્કની સેન્ટ જોન્સ યુનિ.ના ડૉ. કેતન પટેલ ઉપસ્થિત રહ્યા હતા. આ પરિસંવાદમાં ભારત તેમજ વિદેશના વિદ્યાર્થીઓ, પ્રાધ્યાપકો અને ઉદ્યોગ ક્ષેત્રના વ્યાવસાયિકો મળીને કુલ ૨૫૦થી વધુ પ્રતિનિધિઓએ ભાગ લીધો હતો. પરિસંવાદમાં ફાર્માસ્યુટિકલ્સ, બાયોલોજિક્સ અને બાયોસિમિલર્સ, હર્બલ ઔષધિઓ અને મેડિકલ ડિવાઈસીસ જેવા વિષય આધારિત સત્રો ગોઠવવામાં આવ્યા હતા. ઉપરાંત ક્લિનિકલ ડેવલપમેન્ટ અને તેના નિયમન પર વિશેષ પેનલ ચર્ચાનું આયોજન કરવામાં આવ્યું હતું. પરિસંવાદનું એક મુખ્ય આકર્ષણ પોસ્ટર પ્રેઝન્ટેશન હતું, જેમાં ૧૧૦થી વધુ પોસ્ટરો રજૂ કરવામાં આવ્યા હતા.