Five Days International Conference on Advancements in Drug Regulatory Affairs: Pharmaceutical Industry-Academia Perspective (ADRA-2022)

26-30 September, 2022

Convenor: Thakur Gurjeet Singh

Total No. of Participants: 105

Resource Persons :

Sr. no.	Name of the resource person	Full affiliation	Place
1	Prof (Dr) Bhupinder Singh Bhoop	Em. Professor and Ex-Chairperson & Dean, UIPS, Panjab University, Chandigarh	Chandigarh
2	Dr Paolo Rocco	Professor, University of Milan	Italy
3	Dr Pirthi Pal Singh	Vice President, Tirupati Group, Paonta Sahib	HP
4	Dr Umberto Mussazi	Professor, University of Milan,	Italy
5	Dr Harish Dureja	Professor, Department of Pharmaceutical Sciences, MDU	Rohtak, Haryana
6	Dr Praveen Topale	Head DRA, Panacea Biotec Ltd	Punjab
7	Ms Sonia Sharma	RA Head, Morepen Laboratories	HP
8	Dr Sanju Nanda	Professor, Department of Pharmaceutical Sciences, MDU	Rohtak, Haryana
9	Dr Rashmi Panda	GM R&D, Mankind Research Centre,	Gurugram
10	Dr MinakshiGarg	Associate Professor, DPSRU	New Delhi
11	Mr RohanRathi, DRC	Sandoz Development Centre	Hyderabad
12	Dr N Jawahar	Assistant Professor, JSS College ofPharmacy	Tamil Nadu
13	Dr Sandeep Kumar	Assistant Professor, NIPER	Hyderabad

The conference aimed at highlighting the need for drug regulation to ensure the safety and efficacy of drugs for the general public. The pharmaceutical industry is considered as the most highly regulated industries worldwide. The regulatory bodies like, FDA, EMA, CDSCO ensure compliances in various legal and regulatory aspects of a drug. Countries possess their own regulatory authority, which is responsible for enforcing the rules and regulations and issue the guidelines to regulate drug development process, licensing, registration, manufacturing, marketing, labeling and the product life cycle of pharmaceutical products. In an ever-changing regulatory environment, the role of regulatory affairs becomes more critical.

Pharmaceutical regulations, thus, across the world play an important role in ensuring the safety and efficacy of the approved drugs. These not only regulate the pricing of drugs but the quality as well. The regulations are required both for new innovations and already existing products, in order to improve health status.


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