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17th International Conference and Exhibition on

Pharmacovigilance & Drug Safety

June 13-14, 2024 | Rome, Italy



SCIENTIFIC PROGRAM

DAY 1
Thursday, 13th June

08:30-09:00 Registrations

09:00-09:30 Introduction

09:30-09:50 COFFEE BREAK

09:50-11:50 Meeting Hall 01

KEYNOTE LECTURES

11:50-13:10

MEETING HALL 01	MEETING HALL 02
Talks On: Pre-Clinical and Clinical Trials	Talks On: Adverse Drug Reactions
Clinical trials on drugs used in respiratory disorders	Acute and chronic adverse drug events
Regulatory authorities and ethics committee	Classification of various adverse drug reactions
Multi center trials and monitoring	Mechanism of adverse drug reactions
Alternative trials design and models	Assessing and reporting adverse drug reactions
Data collection and quality control	Post marketing reports on adverse effects

13:10-13:15 GROUP PHOTO

13:15-14:00 LUNCH BREAK

14:00-16:00

MEETING HALL 01	MEETING HALL 02
Talks On: Pharmacovigilance and Risk Management	**Talks On:** Good Pharmacovigilance Practice
Information technology in pharmacovigilance and promoting companies	Role of pharmacovigilance and pharmacoepidemiology in risk management
Review of softwares used in pharmacovigilance and clinical trials	Good reporting practices
Implementation of Risk Minimization Plans	Interpreting safety signals
Risk communication: Interface between pharmacovigilance, sales and marketing	Randomized clinical trials

16:00-16:20 COFFEE BREAK

MEETING HALL 01 (16:20-17:00)
Young Researchers in Pharmacovigilance and Drug Safety

DAY 2
Friday, 14th June

09:00-10:30 Meeting Hall 01 — **KEYNOTE LECTURES**

10:30-10:50 COFFEE BREAK

10:50-12:50	MEETING HALL 01	MEETING HALL 02
	Talks On: Pharmacy Practices and its Challenges	**Talks On:** Clinical Trials on Various Disorders
	Pharmacy practice and its guidelines	Emerging technology in clinical trials
	Challenges in compounding and dispensing practice	Recent clinical trials on AIDS
	Dosage regimen, drug toxicity and drug safety measures	Clinical trials on benign and malignant tumors
	Pharmacoepidemiology of drug shortages Drug Labelling	Clinical trials on cardiovascular diseases

12:50-13:35 LUNCH BREAK

13:35-15:55	MEETING HALL 01	MEETING HALL 02
	Talks On: Data Quality Management and Analysis	**Talks On:** PV Consultings and Business Opportunity
	Data Quality Control and Assurance	Pharmacovigilance software Devlopment Companies
	Quality Planning and Improvement	Bussiness Opportunities for PV Software Devloping Companies
	Data Safety Monitoring	Opportunities for PV Software Services companies

15:55-16:15 COFFEE BREAK

MEETING HALL 01 (16:15-17:00)	MEETING HALL 01 (17:00-18:00)
Poster Presentations	Workshop

Awards & Closing Ceremony

PAST AFFILIATES

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Nathan Cummins
Mayo Clinic, USA

Silvia Leone
University of Genoa, Italy

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Professor of Microbiology University of Kansas, USA

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Medical Safety Program Team Leader at Saudi Patient Safety Centre, Saudi Arabia

Dr. Jacob Joseph
Hardvard Medical School, USA

Dr. M .Wasif-Khan
Head Global Pharmacovigilance Evolet Healthcare Pvt. Ltd, India

Dr. Zhou Tom Hui
Senior Principle Scientist Medical Safety at Takeda, USA

Dr.Ashok Srivastava
Chief Medical Officer Clinfo Matrix, USA

Dr.Ujwala Salvi
Chief Executive Officer, Nucleon Therapeutics LLP, India

Dr. Meenal Patwardhan
Therapeutic Area Head Pharmacovigilance & Patient Safety, Abbive, USA

Eleni Konstantinou
Hardvard Medical School, USA

Essam Ghanem
Head of Pharmacovigilance Celyad Biopharmaceutical (Immuno-Oncology), Belgium

PAST AFFILIATES

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Qimat Tayba Pharmaceutical Biotechnology Factory , Saudi Arabia



Glimpses of Pharmacovigilance Conferences








