Date : 3rd September, 2024

Time : 10:00 AM - 1:30 PM

Venue : Lecture Theatre-1, Nehru Hospital,
Postgraduate Institute of Medical Education
& Research (PGIMER)
Sector-12, Chandigarh-160012

Registration Details:

Registration Fee	:	Rs 2000 (Including GST)
Bank Details	:	Bank of Baroda
Account No.	:	21860100013540
Branch	:	Sanjay Nagar, Ghaziabad
IFSC Code	:	BARB0SANGHA
SWIFT Code	:	BARBINBBGHA
Type of Bank Account	:	Super Saving, MICR Code of Bank : 110012204

Scan QR & Register

Contact information for registration and inquiries
shatrunjay.ipc@gov.in , mvpi-ipc@gov.in

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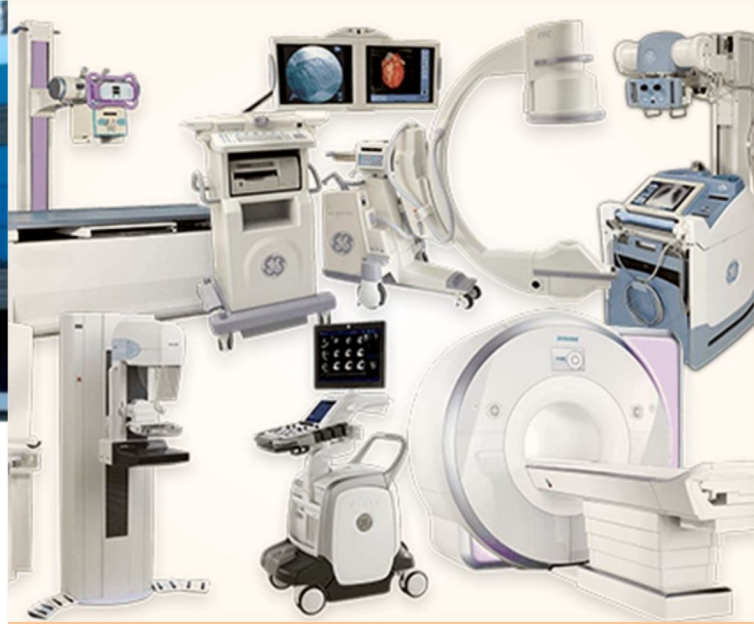
Sector-23, Raj Nagar
Ghaziabad, Uttar Pradesh - 201002

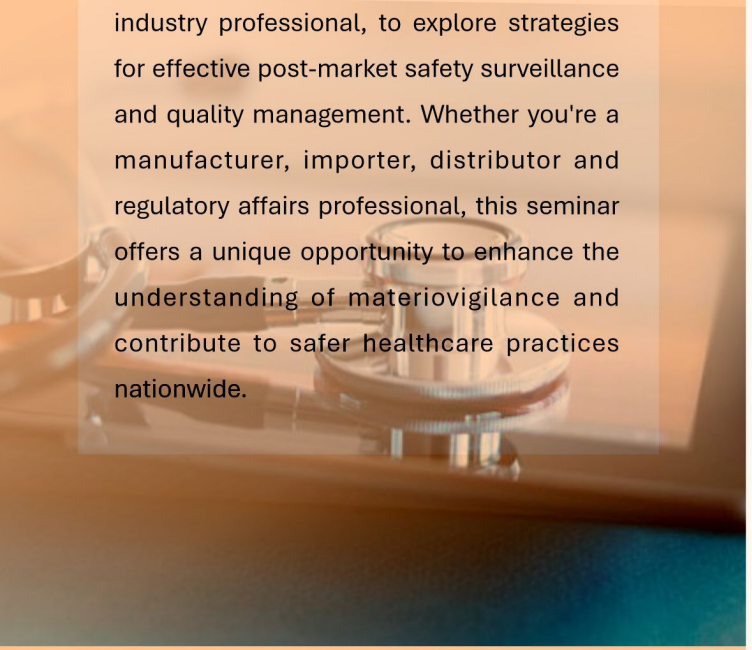


सत्यमेव जयते
IPC

Seminar on Strengthening Medical Device Licence Holders on Post Market Safety Surveillance Practices in India

Introduction

Join us for an insightful “Seminar on Strengthening Medical Device Licence Holders on Post Market Safety Surveillance Practices in India" on 3rd September, 2024 organized by Indian Pharmacopoeia Commission - Materiovigilance Programme of India (IPC-MvPI). This seminar aims to equip medical device manufacturers with essential knowledge to navigate the regulatory landscape of adverse event reporting in India. Participants will engage with experts from the Central Drugs Standard Control Organization (CDSCO), IPC-MvPI, healthcare professionals, and reputed industry professional, to explore strategies for effective post-market safety surveillance and quality management. Whether you're a manufacturer, importer, distributor and regulatory affairs professional, this seminar offers a unique opportunity to enhance the understanding of materiovigilance and contribute to safer healthcare practices nationwide.



Seminar Agenda

10:00 AM - 10:10 AM	Welcome Address and Objective
10:10 AM - 10:40 AM	Medical Devices Rules 2017 - Post Market Surveillance Requirements and Challenges
10:40 AM - 11:10 AM	Medical Device Safety Surveillance Systems in India - Materiovigilance Programme of India (MvPI)
11:10 AM - 11:40 AM	Quality Management System Requirement for Domestic Medical Device Manufacturers
11:40 AM - 12:10 PM	Mitigating Potential Risks: Healthcare Professional's Perspective
12:10 AM - 12:40 PM	MvPI Reporting Tools
12:40 PM - 01:10 PM	Best Practices for Materiovigilance: Insights from Medical Device Manufacturers
01:10 PM - 01:20 PM	Closing Remarks
01:20 PM onwards	Lunch

Background

Indian Pharmacopoeia Commission (IPC), an Autonomous body of Ministry of Health and Family Welfare, Government of India, has entrusted with National Coordination Center (NCC) responsibilities related to Materiovigilance Programme of India (MvPI) since 2018 with an objective to improve Indian patient safety by monitoring, recording and analyzing the root cause of adverse events or risks associated with the use of medical devices and suggesting regulatory bodies for appropriate action with the sole intention of improving patient safety. MvPI aims to promote and facilitate adverse event reporting of medical devices and subsequently evaluating these events.

Objectives

- To equip medical device manufacturers with essential tools and techniques for effective adverse event reporting
- To ensure compliance with MvPI guidelines
- To promote patient safety and device efficacy

Expected Outcome

- It will help in decision making for medical device stakeholders on how to report, when to report and what to report and what not to report to IPC, NCC-MvPI.
- Strengthening of effective materiovigilance system across the country.
- Development of the better quality management systems.

Who Should Attend?

- Medical device manufacturers, importers & distributors
- Quality assurance and regulatory affairs professionals