



AIIMS, Guwahati

# Capacity Building & Strengthening of Materiovigilance Programme of India in North Eastern States



April 12, 2025



10:00 AM - 4:30 PM



**Deadline for Registration**  
April 7, 2025



**Who should attend ?**

- Healthcare Professionals (HCPs), e.g., Doctors, Nurses, Pharmacists
- Bio-Medical Engineers
- Regulatory Affairs Executive in health industries
- All other professionals working in the hospital settings



Scan and Register  
at no cost



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All India Institute of Medical Sciences  
(AIIMS)  
Guwahati, Assam

**NOTE:** e-certificates will be provided to all the participants

## Indian Pharmacopoeia Commission

National Coordination Centre - Materiovigilance Programme of India

Ministry of Health and Family Welfare, Government of India



## Introduction

Join us for the “Capacity Building & Strengthening of Materiovigilance Programme of India in North Eastern States” focused on Medical Device Safety Surveillance Systems in India, tools and techniques for adverse event reporting under the Materiovigilance Programme of India (MvPI) and Causality Assessment. This training is designed to equip healthcare professionals with the knowledge and skills needed to effectively report and manage adverse events, ensuring the highest standards of safety and compliance in the medical device industry.

## Objectives

- To equip HCP (Healthcare Professionals) with essential tools and techniques for effective adverse event reporting.
- To ensure compliance with regulatory guidelines.
- To safeguard public health by ensuring device quality and safety.

## About us

**Indian Pharmacopoeia Commission (IPC)**, an Autonomous body under 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 tracking, documenting, 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.

**AIIMS Guwahati** is a premier medical institute established with the vision of providing world-class healthcare, advanced medical education, and research in North East India. As part of the All India Institutes of Medical Sciences (AIIMS) network, it is committed to developing the healthcare infrastructure in the region. The institute offers state-of-the-art facilities and a comprehensive curriculum that ensures holistic development for medical professionals. AIIMS Guwahati is dedicated to fostering excellence in medical training, empowering future healthcare leaders to meet the evolving needs of society.



# AGENDA

## Capacity Building & Strengthening of Materiovigilance Programme of India in North Eastern States

**April 12, 2025 (Saturday)**

10:00 am - 10:30 am	<ul style="list-style-type: none"><li>• <b>Programme Inauguration</b></li><li>• <b>Welcome Address</b> (AllMS, Guwahati Representative)</li><li>• <b>Opening Remarks</b> (Dr. Bikash Medhi, Professor, Dept. of Pharmacology, PGIMER, Chandigarh)</li><li>• <b>Special Address</b> (State Licensing Authority)</li><li>• <b>Special Remarks</b> (Prof. Ashok Puranik, Executive Director, AllMS-Guwahati)</li></ul>
10:30 am - 11:00 am	<b>Overview of the Regulatory Framework for Medical Devices in India</b> (Representative from CDSCO)
11:00 am - 11:15 am	<b>TEA BREAK</b>
11:15 am - 11:45 am	<b>Medical Device Safety Surveillance Systems in India—Materiovigilance Programme of India (MvPI)</b> (Dr. V. Kalaiselvan, Senior Principal Scientific Officer, Indian Pharmacopoeia Commission (IPC), Ghaziabad)
11:45 am - 12:45 pm	<b>Medical Device Adverse Events Reporting Tools with Decision Tree</b> (Dr. Shatrunjay Shukla, Scientific Assistant, IPC, Ghaziabad)
12:45 pm - 01:15 pm	<b>Causality Assessment</b> (Mr. Naveen V, Scientific Assistant, IPC, Ghaziabad)
01:15 pm - 02:15 pm	<b>LUNCH</b>
02:15 pm - 02:45 pm	<b>Role of Healthcare Professionals in ensuring quality and safety of medical devices</b> (Dr. Bikash Medhi, Professor, Dept. of Pharmacology, PGIMER, Chandigarh)
02:45 pm - 03:15 pm	<b>Mitigating Potential Risks: A Healthcare Professional's Perspective</b> (Dr. Phulen Sharma/Dr. Abhilash Goyal, AllMS Guwahati)
03:15 pm - 04:00 pm	<b>Case Studies with Reportable and Non- Reportable Adverse Events</b> (Representative, IPC, Ghaziabad)
04:00 pm - 04:15 pm	<b>Q &amp; A Session</b>
04:15 pm - 04:20 pm	<b>Vote of Thanks</b> ( Dr. Phulen Sharma, Coordinator, MDMC, AllMS Guwahati)