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Date: November 11, 2024

NOTICE

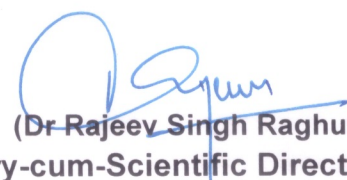
To,

1. The Drugs Controller General (India)
2. The States/UTs Drug Regulators
3. IDMA, OPPI, etc.

Subject: Implementation of Pharmacovigilance Guidance Document for Marketing Authorization Holders (MAHs) of Pharmaceutical Products, Version 2.0 with effect from 1st February 2025 - Reg.

Sir/Madam,

It is brought to the notice of all concerned that the National Coordination Centre-Pharmacovigilance Programme of India (NCC-PvPI), Indian Pharmacopoeia Commission (IPC) in collaboration with Central Drugs Standard Control Organization (CDSCO) released the Pharmacovigilance Guidance Document for Marketing Authorization Holders (MAHs) of Pharmaceutical Products, Version 2.0 on 17th September 2024. This PV Guidance Document for MAHs will be effective from 1st February 2025 to strengthen the Pharmacovigilance system at MAHs organization in India.



(Dr. Rajeev Singh Raghuvanshi)
Secretary-cum-Scientific Director, IPC

Copy to:

1. PS to Secretary-cum-Scientific Director, Indian Pharmacopoeia Commission, Ghaziabad.
2. Library and Information Officer, Indian Pharmacopoeia Commission to upload it on the website of IP Commission, Ghaziabad