INDIAN PHARMACOPOEIA COMMISSION

National Coordination Centre, Pharmacovigilance Programme of India

MINISTRY OF HEALTH & FAMILY WELFARE, GOVERNMENT OF INDIA
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OFFICE ORDER

The Pharmaceutical Industries are hereby informed to submit Adverse Drug Reactions (ADRs) due to their respective pharmaceutical product to National Coordination Centre (NCC), Pharmacovigilance Programme of India (PvPI) in xml-E2B format to hasten the process of uploading Individual Case Safety Reports (ICSRs) in VigiFlow, a WHO-UMC, Web database. Also this will enhance the process of assessment and signal detection.

This shall be effective from 1st June 2015.

(Dr. G. N. Singh)
Secretary-cum-Scientific Director

Copy To,

1) Drugs Controller General (India), FDA, Bhawan, Kotla Road, New Delhi. - To made available on CDSCO Website.
2) Dr. S. K. Gupta, Chairman, Quality Review Panel, Emeritus Professor & Head, Department of Clinical Research, DIPSAR, New Delhi. – For information.
3) Dr. Urmila Thatte, Chairman, Signal Review Panel, Professor & Head, Department of Clinical Pharmacology, Seth GS Medical College & KEM Hospital, Mumbai. – For information.

To be upload in website