

## INDIAN PHARMACOPOEIA COMMISSION

National Coordination Centre- Pharmacovigilance Programme of India (PvPI) MINISTRY OF HEALTH & FAMILY WELFARE, GOVERNMENT OF INDIA SECTOR-23, RAJ NAGAR, GHAZIABAD- 201 002.

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## OFFICE ORDER

This is to inform all Adverse Drug Reactions Monitoring Centres under Pharmacovigilance Programme of India, that from 1st April 2017 onwards you will be required to send source documents with respect to Serious ICSRs and Unlabelled ADRs Cases i.e. Scan Copy of Suspected Adverse Drug Reaction form to NCC-PvPI on email i.d. <a href="mailto:pvpi@ipcindia.net">pvpi@ipcindia.net</a>. This step is aimed at improving the Quality of ICSRs, timely action on any suspected event due to use of Medicinal Products and also to facilitate Signal Review Processes.

(Dr. V.Kalaiselvan) Principal Scientific Officer

Copy to-

- 1. PS to Secretary-cum-Scientific Director, IPC, Ghaziabad.
- 2. PS to DCG (I), FDA Bhawan, New Delhi.
- 3. All Patient Safety Pharmacovigilance Associates/Staff of NCC-PvPI.