



**INDIAN PHARMACOPOEIA COMMISSION**  
MINISTRY OF HEALTH & FAMILY WELFARE, GOVERNMENT OF INDIA  
SECTOR-23, RAJ NAGAR, GHAZIABAD- 201 002.  
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No. T.11012/01/2017-AR&D

Date: 30.09.2019

To,

1. The Drugs Controller General (India)
2. All Zonal Offices/Port Offices of CDSCO
3. All State Drug Controllers
4. Directors of Central Drug Laboratories
5. Directors of State Drug Laboratories
6. Members of Scientific Body of the IPC
7. Government Analysts
8. IDMA/OPPI/BDMA/FSSAI/Small Scale Industry Associations

**NOTICE**

The competent authority has approved the extension of implementation date of Indian Pharmacopoeia (IP) Addendum 2019 to IP 2018 for a period of 03 months i.e. up to 31<sup>st</sup> December, 2019.

This is for notice and compliance.

Yours faithfully,

  
(Dr. Jai Prakash) 30/09/2019  
Sr. Principal Scientific Officer

**Copy to:**

1. PPS to Secretary (H&FW), Ministry of Health & Family Welfare, Govt. of India, Nirman Bhawan, New Delhi.
2. PS to Additional Secretary & DG (CGHS), Ministry of Health & Family Welfare, Govt. of India, Nirman Bhawan, New Delhi.
3. PS to Joint Secretary (R), Ministry of Health & Family Welfare, Govt. of India, Nirman Bhawan, New Delhi.

*Indian Pharmacopoeia (I.P.)*

– *The book of standards for drugs.*

*National Formulary of India (N.F.I.)*

– *The reference book that promotes rational use of generic medicines.*

***On Path of Evolving a Modern Scientific Institution***