

संख्या / F. No. 19(78)/2014/Div. II/NPPA  
भारत सरकार  
Government of India  
रसायन और उर्वरक मंत्रालय  
Ministry of Chemicals & Fertilizers  
औषध विभाग  
Department of Pharmaceuticals  
राष्ट्रीय औषध मूल्य निर्धारण प्राधिकरण  
National Pharmaceutical Pricing Authority

तीसरी/पांचवी मंजिल 5<sup>th</sup> / 3<sup>rd</sup> Floor,  
वाई.एम.सी.ए. सांस्कृतिक केन्द्र बिल्डिंग  
YMCA Cultural Center Building,  
1, जय सिंह रोड, नई दिल्ली-110001  
1, Jai Singh Road, N. Delhi – 110001.  
दिनांक : 16.02.2017

**OFFICE MEMORANDUM**

**Subject: Guidelines for review order implementation-IPDMS compliance**

The undersigned is directed to refer to the O.M. No. 8/34/2014/Div.II/NPPA dated 07.02.2017 and the pending review orders of Department of Pharmaceuticals for compliance. Most of the orders are for reconsideration of the PTR and other data by NPPA. This was discussed in the 41<sup>st</sup> meeting of the Authority held on 13.02.2017 and Authority decided that in future, the companies challenging the PTR taken by AIOCD/NPPA, must supplement the evidence with their IPDMS submissions, in order to be considered for price revision.

2. Accordingly, petitioner(s)/concerned company(ies) are required to submit copies of supporting IPDMS submissions. A certificate to the effect that all requisite forms for all formulations have been filed online through Integrated Pharmaceutical DataBase Management System (IPDMS) may also be submitted to NPPA, **positively by February 23, 2017** so as to enable NPPA to take appropriate action in the matter. Such submissions may also be made in all future cases without fail. These instructions shall apply to all future cases as well.

  
(ए. के. खुराना)  
निदेशक

Copy to (for information and necessary action please):

- (i) All Apex Pharma organizations/Associations i.e. OPPI, IDMA, AISSPMA, PICCI, CII, IPA and FOPE.
- (ii) Sh. M.K. Bhardwaj, Deputy Secretary, DOP, Shashtri Bhawan, New Delhi.