



F. No. 4-9/2017 RRR
Government of Pakistan
Ministry of National Health Services, Regulations and Coordination
Drug Regulatory Authority of Pakistan
(Division of Pharmaceutical Evaluation and Registration)
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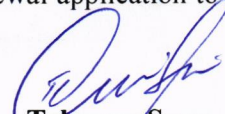
SAY NO TO CORRUPTION

Islamabad the 23rd August, 2019

1. Chairman Pakistan Pharmaceutical, Manufacturing Association (PPMA), PPMA House, House No. 474, Street No. 34, Sector I-8/2, **Islamabad**
2. Executive Director Pharma Bureau, Chamber of Commerce & Industries Building Talpur Road, **Karachi**
3. Executive Director Pakistan Chemist & Druggist Association (PCDA), 504, 4th Floor Mashriq Centre, Near Civic Centre, Gulshan-e-Iqbal, Block-14, **Karachi**

Subject: **SUBMISSION OF RENEWAL APPLICATION OF BIOLOGICAL DRUGS IN BIOLOGICAL DIVISION**

I am directed to refer to the subject captioned above and to inform that since promulgation of DRAP Act, 2012, Division of Biological Evaluation & Research has been established to carry out functions specifically related to Biological Drugs (both registration and renewal thereof). However, it has been observed that majority of manufacturers and importers of Biological products are still submitting their renewal applications in the RRR Section of Pharmaceutical Evaluation & Registration Division. Accordingly all the manufacturers and importers of registered biological products are hereby advised to address their renewal application to Biological Evaluation & Research Division for further processing.


Tehreem Sara
(Deputy Director RRR)
Pharmaceutical Evaluation &
Registration Division

Copy to: -

1. Assistant Director to CEO, DRAP, Islamabad.
2. Director PE&R, DRAP, Islamabad
3. Director Biological Evaluation & Research, DRAP, Islamabad
4. Director QA & LT DRAP, Islamabad for onward communication to DRAP sub offices to inform all importers and manufacturers of biological products in their area of jurisdiction.
- ✓ 5. Director (MIS) DRAP, Islamabad for placing on the DRAP website
6. Master file.



*upload on website.
DD MS- 2/1/2019*