**F. No : 10-2/2018-MIS**
**Government of Pakistan**
**Ministry of National Health Services, Regulations & Coordination**
**Drug Regulatory Authority of Pakistan**
**TF-Complex G-9/4 Islamabad**
**********

Islamabad the 10th July, 2019

**"SAY NO TO CORRUPTION"**

1) The Executive Director
   Pharma Bureau - Karachi
2) The Chairman
   Pakistan Pharmaceutical Manufacturer Association – Islamabad
3) The Chairman
   PCDA Karachi
4) Chairman KPPMA
   Peshawar

**SUBJECT: TRAINING OF DRUG- INTEGRATED REGULATORY INFORMATION MANAGEMENT SYSTEM (D-IRIMS)**

The undersigned is pleased to inform that the Management Information Services Division(MIS) of Drug Regulatory Authority of Pakistan (DRAP) is in the process to establish and strengthen the IT system for all regulatory functions which will lead to ease of doing official business. As a first step the MIS Division has developed a software i.e. Drug-Integrated Regulatory Information Management System (D-IRIMS) in consultation with PE&R, Drug licensing and QMS Division of DRAP. The software includes the following features.

1) Online Fee Challan Generation
2) Online application for drug registration
3) Online application for renewal of drug registration
4) Online application for post registration changes
5) Online Data submission of 2-D Barcode system
6) Mobile App for the verification of 2-D Barcode

2. In view of above a hands-on training will be started for the relevant professionals regarding the use of D-IRIMS at Karachi, Lahore, Peshawar and Islamabad. Our team will be available in the provincial headquarters on the following dates.

**PTO**

| Sr. No. | Date & Time | City | Participants |
|---|---|---|---|
| 01 | 22nd and 23rd July, 2019 (Monday & Tuesday) | Karachi | Pharma Bureau, PCDA, PPMA and Pharma Industry of Sindh and Balochistan |
| 03 | 24th July, 2019 (Wednesday) | Lahore | Pharma Industry of Punjab |
| 04 | 29th July, 2019 (Monday) | Islamabad | Pharma Industry of Islamabad and AJK |
| 05 | 31st July, 2019 (Wednesday) | Peshawar | Pharma Industry KPK |

3. It has been decided that venue and other arrangements will be made by the pharma associations and training on the software will be provided by DRAP. You are therefore requested to make arrangements on the dates of the availability of DRAP team at your city to deliver hands on training. Arrangements shall be finalized in consultation with the Mr. Atta-Ur-Rahman, Additional Director(MIS) DRAP Islamabad through your focal person at the following contact details.

Email : addl-directormis@dra.gov.pk
Phone #:051-9107414

4. After the training of software, the applications regarding the registration, renewal and post registration changes will be received online through the D-IRIMS software.

5. Moreover all the pharmaceutical companies and importers are also requested to upload all the data of their registered products on Drug Regulatory Information System (DRIS) which is integrated with the D-IRIMS. Usernames and passwords have already been issued to the registered firms. At present 430 companies are registered on software and the remaining

companies / importers are requested to get registered on system and upload the registered product data up to 30th July 2019.

6. This issues with the approval of competent authority.

**(Atta-Ur-Rehman Ch)**
**Additional Director(MIS)**
**Phone # 051-9107414**

Copy to: -

1) The CEO DRAP
2) All Directors DRAP.
3) All Additional Director (DRAP Field offices) [with request to attend the training and provide the logistic support to DRAP Officials]
4) DD(MIS) to upload on official website of DRAP.
5) Office file