



Dy. No.01/2026-Dir(MIS)
Government of Pakistan
Ministry of National Health Services Regulations & Coordination
Drug Regulatory Authority of Pakistan
Islamabad

Dated: 24-07-2026

Pakistan Pharmaceutical Manufacturers Association (PPMA) House No.474, Street No.34, I-8/2, Islamabad.	Pharma Bureau, Chamber of Commerce Building, Talpur Road, Karachi.
Pakistan Chemist and Druggist Association PCDA), Suit No.504, 5 th Floor, Mashriq Centre, Near Civic Centre, Gulshan Iqbal, Block 14, Karachi.	KPK Pakistan Pharmaceutical Manufacturers Association (KPPMA) KPK.

Subject: **Intimation Regarding Capacity Building & Technical Awareness Sessions on 2-D Barcode and Serialization Implementation.**

Dear Sir,

Drug Regulatory Authority of Pakistan (DRAP) is pleased to inform that, pursuant to promulgation of the revised **2-D Barcode and Serialization Rules**. DRAP intends to arrange **technical session (QA Session)** for the pharmaceutical industry.

2. The objective of these sessions is to provide comprehensive guidance on regulatory requirements, technical standards and practical implementation of **2-D Barcodes and Serialization** in order to ensure effective compliance and elimination of counterfeit medicines from the market and access of safe and authentic medicines.

3. The training sessions are schedules as per the details given below.

Dates	City	Venue	Trainers
27th July 2026	Peshawar	To be decided by Industry Associations	Mr. Saqeb Arshad (Addnl Director)/Mr. Ijaz Mustafa (DD-MIS)
29th July 2026	Islamabad/Rawalpindi	To be decided by Industry Associations	Mr. Saqeb Arshad (Addnl Director)/Mr. Ijaz Mustafa (DD-MIS)
3rd August 2026	Lahore	To be decided by Industry Associations	Mr. Saqeb Arshad (Addnl Director)/Mr. Abdur Rehman (DD-MIS)
6th August 2026	Karachi	To be decided by Industry Associations	Mr. Saqeb Arshad (Addnl Director)/Mr. Yasir Mehmood (DD-MIS)

4. PPMA, KPPMA, PCDA and Pharma Bureau are requested to **nominate focal persons**, disseminate this information among member companies and encourage active participation of relevant technical and regulatory staff. The cooperation in this important regulatory initiative is highly appreciated.

5. This issues with the approval of CEO DRAP.


Atta Ur Rahman Chaudhry
Director (MIS)

Drug Regulatory Authority of Pakistan

Copy to:

1. CEO, Drug Regulatory Authority of Pakistan.
2. All Additional Directors Filed Offices (For the logistic support and coordination with associations for the finalization of venue).