



Dy.No. 63/2026-DIR(MIS)
Government of Pakistan
Ministry of National Health Services Regulations & Coordination
Drug Regulatory Authority of Pakistan
Islamabad

Dated: 21-09-2026

SUBJECT: TRAINING OF ONLINE PHARMACEUTICAL TRACK AND TRACE SYSTEM FOR IMPLEMENTATION OF 2-D BARCODE AND SERIALIZATION. (FOR ISLAMABAD/RAWALPINDI/AJK INDUSTRY)

Please refer to DRAP's previous communications including the amendments to the Drugs (Labelling and Packing) Rules, 1978 vide S.R.O. 963(I)/2026 dated 9th June 2026, and the subsequent DRAP's Guidelines on Data Matrix Barcode and Serialization issued on 28th July 2026, DRAP is transitioning to operational enforcement of its Pharmaceutical Track & Trace framework. To meet the mandatory deadline of **9th October 2026** for 2-D barcode printing and unique product serialization across all manufactured and imported pharmaceutical and biological products. DRAP has deployed the unified digital platform for pharmaceutical Track and Trace System. This software serves as the central operational backbone to enable end-to-end supply chain visibility, product verification, and the eradication of counterfeit drugs from the market.

2. An online Pharmaceutical Track and Trace system for implementation of **2-D barcode and serialization requirements** has been developed and deployed, enabling effective traceability and verification of regulated products manufactured after 9th October 2026. The implementation is being undertaken in accordance with the applicable regulatory requirements of **S.R.O. 963(I)/2026** dated June 9th June 2026.

3. In order to facilitate the pharmaceutical industry and other relevant stakeholders, hands-on training session on online system have been scheduled as per details given as follows.

DRAP Focal Point	Date	Venue
Mr Ijaz Mustafa Deputy Director MIS, DRAP Head Office 0300-0914044	23rd Sep 2026 (Wednesday)	Rawalpindi Chamber of Commerce & Industries (RCCI)

4. Members of all concerned pharmaceutical associations, manufacturers, importers, and other relevant stakeholders are hereby invited to participate in the aforementioned training session. The session will provide detailed guidance and practical demonstrations regarding serialization, 2-D barcode requirements, data submission, system workflows, and other related operational procedures to facilitate effective and uniform implementation of the applicable regulatory requirements.

5. This issues with the approval of CEO, Drug Regulatory Authority of Pakistan

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(Atta ur Rahman Ch)
Director, MIS Division
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Distribution:

1. Chief Executive Officer, DRAP.
2. Director QA< DRAP Head Office
3. Additional Director (MIS) DRAP Head Office
4. All Additional Directors (Field Offices) with the request to arrange the logistic support.
5. All concerned Focal Points, Field Offices DRAP
6. All concerned Pharmaceutical Associations (Human/Veterinary)
7. All manufacturers/importers.
8. Office File.



Director, MIS Division