



O.M. No.28/2026-Dir(MIS)
Government of Pakistan
Ministry of National Health Services Regulations & Coordination
Drug Regulatory Authority of Pakistan
Islamabad

Dated: 06-08-2026

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.

MEMORANDUM

SUBJECT: DIRECTIVE FOR MANDATORY DATA VALIDATION ON eAPP SOFTWARE FOR INTEGRATION WITH TRACK AND TRACE SYSTEM.

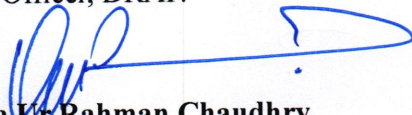
The undersigned is directed to refer to this Authority's earlier Memorandum of even number dated 10th June 2026 regarding the subject cited above, wherein the implementation of 2-D barcode and serialization requirements was notified following the Drugs (Labelling and Packing) Rules, 1986 vide S.R.O. 963(I)/2026 dated 09.06.2026.

2. In continuation of the ongoing system deployment, it is hereby communicated that the eAPP Software (pertaining to Licensing and Product Registration) is being integrated with the national Drug Track and Trace System. Under the integrated mechanism, the Track and Trace System shall automatically fetch manufacturing license and product registration details (mandatory required for 2-D barcode generation and serialization). Consequently, no manual entry of product or licensing data shall be permitted within the Track and Trace System.

3. In view of above, all pharmaceutical manufacturers, importers, and relevant industry stakeholders are directed to perform complete verification and validation of their respective licensing and product data on the eAPP portal on or before 25th August 2026 positively. The eAPP system shall be automatically locked after 25th August 2026. Thereafter, no request for data validation, correction, or modification shall be entertained.

4. In case of any technical, operational difficulty or guidance required during the validation process, the concerned authorised persons may contact to **PE&R Division (Mr. Sayyad Hussain Khan, Director, 0334-5352661)** and **MIS Division (Mr. Saqeb Arshad, Additional Director, 0300-9553737)** of DRAP immediately.

5. This issues with the approval of the Chief Executive Officer, DRAP.


Atta Ur Rahman Chaudhry

Director (MIS)
Drug Regulatory Authority of Pakistan

Copy to:

1. CEO, Drug Regulatory Authority of Pakistan.
2. Director, PE&R, DRAP, Islamabad.
3. Director, Licensing, DRAP, Islamabad.
4. All Additional Directors Filed Offices, DRAP.
5. All Chief Drug Controllers, Islamabad, Punjab, Sindh, KPK, Balochistan, Gilgit Baltistan and AJK.
6. Deputy Director (V), MIS Division, DRAP with the request to upload on official website of DRAP.