F.No. 8-5/2022-AD (QMS)
Government of Pakistan
(Drug Regulatory Authority of Pakistan)
T.F Complex, 7th Mauve Area G-9/4, Islamabad.
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Islamabad, the 15th August 2022

1. **Chairman,** Pakistan Pharmaceutical Manufacturer Association, Islamabad.	2. **President,** Pharma Bureau, Karachi.

Subject: **TRAINING SESSION ON REGULATORY DATA STANDARDS**

Regulatory Data standard help Regulatory Authorities and Pharmaceutical Manufacturers for efficient exchange of regulatory data which ultimately reduces the processing time by avoiding clarifications due to variation in terminologies. These standards also facilitate data sharing between regulators and manufacturers in better understandable way, through both manual and electronic means, in a harmonized and consistent manner that is interoperable across the globe.

2. In this context, DRAP in collaboration with United States Pharmacopeia, Promoting the Quality of Medicines Plus Program has organized following training sessions on the regulatory data standards for the manufacturers. The sessions are: -

i. Training on Regulatory Data Standards for Pharmaceutical Manufacturers on 19th August 2022 at Karachi.
ii. Training on Regulatory Data Standards for Pharmaceutical Manufacturers on 22nd August 2022 at Lahore.

3. You are requested to share nominations of technical and regulatory workforce for each training by 16th August 2022 positively. Nominated person can register themselves free of cost at www.dra.gov.pk/events on a first-come basis.

15/08/22.

(Sayyad Hussain Khan)
Deputy Director to CEO

Copy to:

1. Director, Admin, HR& Logistics, DRAP, Islamabad
2. Mr. Waqas Ahmed, Chief of Party, USP PQM Plus program
3. PS to CEO, DRAP, Islamabad
4. Concerned Field Offices of DRAP