F.No.5-8 /2025-MIS-P-001
Government of Pakistan
Ministry of National Health Services, Regulations & Coordination
Drug Regulatory Authority of Pakistan
*****

Islamabad, 22nd January, 2026.

ATTENTION: ALL MANUFACTURERS AND IMPORTERS OF MEDICAL DEVICES.

SUBJECT: TRAINING SESSION FOR MEDICAL DEVICE STAKEHOLDERS ON THE MDMC LICENSING & PRODUCT REGISTRATION PORTAL.

In line with its vision to promote transparency, digitalization, and ease of doing business, and as part of its continued commitment to streamline regulatory procedures through digital transformation, the Drug Regulatory Authority of Pakistan (DRAP) in collaboration with Healthcare Devices Association of Pakistan is pleased to conduct a refresher training for additional/updated features of Post Registration Variation(PRV) and Post Licensing Variation (PLV) on the MDMC Licensing & Product Registration Portal for medical device importers and manufacturers of Pakistan.

2. This training is specifically aimed at facilitating the smooth adoption and effective use of the **MDMC Licensing & Product Registration Portal**, developed to assist applicants in efficiently submitting, tracking, and managing their licensing and product registration, along with Post Registration Variation (PRV) and Post Licensing Variation (PLV) applications.

3. The training session will be conducted as per the following schedule:

Date	Time	Venue
Monday 26th January, 2026	02:00 PM to 05:00 PM	FPCCI Islamabad Office (FPCCI Capital House, G-8/1, Aiwan – e – Sanat – o – Tijarat Road, Mauve Area, Islamabad – Capital Territory)
Wednesday 28th January, 2026	11:00 AM to 02:00 PM	FPCCI Peshawar Office (Office No – 4, 2nd Floor, Al-Fatah Medicine Center, New Krishanpura, Near Bank of Punjab, Main G-T Road, Peshawar Khyber Pakhtunkhwa (KPK)
Friday 30th January, 2026	02:00 PM to 05:00 PM	FPCCI Lahore Office (22 – West Canal Bank Road,Near Tech Society,Lahore – Punjab.)
Tuesday 03rd February, 2026	02:00 PMs to 05:00 PM	FPCCI Karachi Office (Federation House, Tariq Sayeed Complex, Main Clifton, Block 5, Abdullah Shah Gazi Road, Karachi-Sindh.)

4. Session Highlights will include:

- Step-by-step guidance on portal usage.
- Document submission procedures.
- Application tracking system.
- Common errors and how to avoid them.
- Live Q&A session with DRAP experts.

5. All relevant stakeholders are encouraged to actively participate and nominate the employee who is directly responsible for submitting applications on the MDMC portal to attend the training session on the above mentioned date and time. Your cooperation is crucial in strengthening the medical device regulatory ecosystem in Pakistan and aligning it with international best practices.

6. This is issued with the approval of the CEO, DRAP.

(Dr. Ghayour Ahmed)
Deputy Director (MD&MC)

Distribution:

1. Healthcare Devices Association of Pakistan (HDAP).
2. Pakistan Electro-Medical Equipment Manufacture and Distribution Association.
3. Surgical Instruments Manufacturers Association of Pakistan (SIMAP).
4. Pakistan Pharmaceutical Manufacturers Association.
5. Pharma Bureau.
6. Pakistan Chemists and Druggists Association.

Copy to:-

1. PS to CEO, DRAP.
2. Director (MDMC), DRAP.
3. Director MIS, DRAP (with the request to post the letter on the official website of DRAP for the stakeholder's)
4. Additional Director Peshawar, DRAP (with the request to coordinate for the training).
5. Additional Director Lahore, DRAP (with the request to coordinate for the training).
6. Additional Director Karachi, DRAP (with the request to coordinate for the training).