F.No. 9-2/2022-PEC
Government of Pakistan
Ministry of National Health Services, Regulations and Coordination
(Drug Regulatory Authority of Pakistan)
TF Complex Sector G-9/4
****

Islamabad, 26th January, 2024

CIRCULER

ACCEPTANCE OF IAEA TECHNICAL REPORTS ON RADIOPHARMACEUTICALS AS REGULATORY REFERENCE

It is hereby circulated for information of all concerned that the DRAP Authority in its 176th meeting held on 6th December, 2023 has observed that the International Atomic Energy Agency (IAEA) has published various documents of technical guidance for member states on radiopharmaceuticals which are considered as a reference in addition to those specified in pharmacopoeia. Therefore, the Authority approved the IAEA Technical Report 466 & other technical reports on Radiopharmaceuticals Production & Quality Control issued by IAEA as one of the regulatory references of DRAP for Radiopharmaceutical products.

2. All applicants are, therefore, advised to comply the aforesaid decision of the Authority.

26/01/2024.

(ASIF JALIL)
Deputy Director (PE&R) /
Incharge PEC

Distribution:

1. Chairman, Pakistan Pharmaceutical Manufacturer's Association, Islamabad.
2. Executive Director, Pharma Bureau, Karachi.
3. All Members of the Registration Board.
4. Director General (PINSTECH), Islamabad

Through E-Office:

1. All Directors of DRAP.
2. Additional Director (MIS), DRAP for information and uploading on DRAP's website.
3. Additional Director / Officer In-charge DRAP Karachi, Lahore, Islamabad, Peshawar, Quetta for circulation to pharmaceutical units and importers located in their respective area of jurisdiction.

Deputy Director (PE&R) /
Incharge PEC