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Government of Pakistan
Ministry of National Health Services, Regulations and Coordination
(Drug Regulatory Authority of Pakistan)
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NOTIFICATION

Islamabad, the 15th August, 2023.

S.R.O. 1067(I)/2023.-- In exercise of the powers conferred by sub-section (1) of section 20 of the Drug Regulatory Authority of Pakistan Act, 2012 (XXI of 2012) read with sub-rule (3) of rule 4 of the Drug Regulatory Authority of Pakistan (Fee and Levy) Rules, 2022, and in partial modification of its previous Notification No. S.R.O.496(I)/2023 dated the 17th day of April, 2023, the Drug Regulatory Authority of Pakistan, with the approval of the Policy Board, is pleased to direct that the fee specified in column (4) of the Table below shall be levied in respect of functions specified in column (2) and (3) thereof, subject to conditions prescribed in this Notification, namely:-

TABLE

Sr.	Regulatory function	Description	Fee (Rs.)
(1)	(2)	(3)	(4)
1	Issuance of certificate of Good Manufacturing Practices (GMP) for all therapeutic goods	For GMP certificate requiring panel inspection	25,000/- per annum
		For a subsequent GMP certificate for any other country on the basis of already conducted inspection for GMP certificate.	12,500/-

(i) One GMP certificate for a country or group of countries shall be issued with single fee;
(ii) One free sale certificate for a country or group of countries shall be issued with single fee; and
(iii) GMP certificate shall be valid for three years.

[No. F.6-20/2023-QA&LT]

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Deputy Director (Legal Affairs).

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