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The Gazette of Pakistan

EXTRAORDINARY
PUBLISHED BY AUTHORITY

ISLAMABAD, TUESDAY, OCTOBER 15, 2019

PART II

Statutory Notifications (S.R.O.)

GOVERNMENT OF PAKISTAN
Ministry of National Health Services, Regulations and Coordination
Drug Regulatory Authority of Pakistan

NOTIFICATION

Islamabad, the 15th October, 2019

S.R.O. 1225(I)/2019.– 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 clause (n) of section 7 and clause (d) of section 11 thereof, the Drug Regulatory Authority of Pakistan, with the approval of the Policy Board, is pleased to direct that the fee specified in column (3) of the Table below shall be payable in respect of the services specified in column (2) thereof, namely:--

TABLE

Sr.	Description of services	Fee (Rs.)
(1)	(2)	(3)
1	Certificate of Good Manufacturing Practices (GMP) for export purposes for a period of three years.	15,000/-

[File No. 6-26/2011-QA(Pt.)]

AAMAR LATIF,
Deputy Director (Legal Affairs).