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

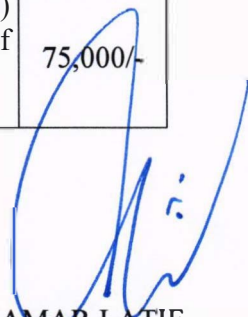
Islamabad, the 16th April, 2020

NOTIFICATION

No. F.6-18/2020-QA/DRAP.-- In exercise of the powers conferred by sub-section (I) 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 local manufacturing with validity for a period of two years.	50,000/-
2	Certificate of Good Manufacturing Practices (GMP) for export purposes with validity for a period of three years.	75,000/-


AAMAR LATIF,
Deputy Director (Legal Affairs).

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