

TECHNICAL EVALUATION REPORT FOR PROCUREMENT OF HEPATITIS-B PCR KITS/DEVICES, PUNJAB HEPATITIS CONTROL PROGRAM, FY 2020-21			
PURCHASE CELL, DGHS PUNJAB			
PC-7/HCP/2020-21/HBV-PCR		1	2
Sr #	NAME OF THE BIDDER	M/s ROCHE PAKISTAN	M/s PAKISTAN MICROBIOLOGICAL ASSOCIATES
	QUOTED PRODUCT	ROCHE COBAS 6800 HBV 96 T IVD (HBV PCR KITS) Germany	SYSTAQ AB Quantgene (Amp Unit), Extraction System (Extraction Unit) USA
a	Original Tender Purchase Receipt obtained by Depositing Rs. 2000/- (Non-Refundable) to Cashier, Accounts Branch, DGHS	Yes	Yes
b	The bidder will submit 2% bid security (Copy with Technical Bid and Original Financial Bid) in the form of Bank Guarantee/CDR/Pay order from any scheduled bank.	Yes	Yes
c	Valid Drugs Manufacturing License(for manufacturers)/Establishment Registration Certificate(for Sole Agents/Authorized agent). However, the registration holding firm will be responsible for quantity & quality of the product.	No	No
d	Valid Drug Sale license. (where applicable)	Yes	No
e	Valid Drug Registration Certificate / Medical Device enlistment Certificate, Whichever is applicable as per Medical Devices Rules 2017 of the quoted product issued by DRAP Pakistan	Yes	No
f	Valid GMP certificate issued by DRAP (for local/manufacture).	N / A	N / A
g	Valid quality certification of US FDA/WHO/CE/JMHLW prequalified/approval of the quoted product.	Yes (FDA Approved)	Yes (CE Approved, Origin is USA)
h	Valid Free Sale Certificate indicating that the quoted brand is freely available in the country of manufacturer. This certificate must be issued by relevant authority of the country of origin duly legalized/notarized by embassy of Pakistan in country of manufacturer.	Yes	Yes
i	The firm undertakes that currently it is not Blacklisted/Debarred by any Government, or its organization or project on valid Rs. 100 judicial stamp paper duly legalized/verified by notary public.	No (Copy of stamp paper attached)	Yes
j	National Tax Number(NTN) Sales Tax Registration Number Certificate with documentary proof of ATL shall have to be provided by bidder.	Yes	Yes
k	The bid must comply with the advertised technical specification of the quoted item. Incomplete offer will straightaway be rejected.	Yes	No (System is semi automated having two seprate Extraction & Amplification Units)
l	The bidder shall provide 04 sample of quoted package, as per requirement for evaluation / satisfaction of the Committee along with its bid/offer. Sample, will be evaluated by the End User by analyzing its Production quality, Design, Reliability, Conformance to the specification & quality standards and safe for the usage. End user evaluation will be knowck down criteria.	Yes	Yes
STATUS		NON-RESPONSIVE	NON-RESPONSIVE
Reasons of Rejection		1. Valid Drugs Manufacturing License/Establishment Registration Certificate is not provided. 2. Copy of stamp paper (undertaking) has been attached instead of original one.	1. Valid Drugs Manufacturing License/Establishment Registration Certificate is not provided. 2. Valid Drug Sale license is not provided. 3. Valid Drug Registration Certificate / Medical Device enlistment Certificate is not provided. 4. The quoted item is semi-automated having separate Extraction and Amplification Units.