

TECHNICAL EVALUATION REPORT OF INFUSION DEXTRAN-40 FOR EP&C PROGRAM FOR FY 2018-19, DGHS

T.E No. PC-15/EP&C/2018-19

Sr. No.	Name of Item	Name of Firm	Compulsary Parameters								Status	Reason(s) of Rejection
			The bidder must possess valid Drug Manufacturing License issued by DRAP (manufacturers) and valid Drug Sale License (in case of importers).	The bidder will provide valid GMP Certificate from Country of Manufacturer/Country of Origin.	In case of Sole agent bidder will provide Valid Sole Agency Certificate/Authorization Certificate for the quoted item.	Specifications quoted in the technical offer will be verified from samples provided with the bid. Product that comply 100% with the advertised specifications and fulfill the requirements as per rules shall be considered.	Undertaking Regarding "Declaration of No Spurious/Adulterated Batch" by any notified Drug Testing Laboratory of quoted item within last three years on valid Rs.100 judicial stamp paper duly verified by notary public	The firm undertakes that currently it is not Blacklisted/Debarred by any procuring agency on valid Rs.100 judicial stamp paper duly verified by notary public.	Sample for evaluation by the technical committee.	The item other than registered products under Drug Act 1976/DRAP Act 2012 can also be quoted subject to valid registration under relevant law of the country of origin/manufacturing shall be pre-requisites.		
1	Infusion Dextran-40 Saline	M/s The Medicine Company	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Responsive	