

NIT 20 Prebid Queries and Resolution

Sr. No.	Tender Technical Specification	Queries Raised by Bidder	Amendment/ Resolution
1	If a bidder is MSME/Udhyog Aadhar/SSI registered manufacturer of Madhya Pradesh then they will be exempted from submitting EMD and Bid document fee. However tender processing fee is not exempted. If MSME/Udhyog Aadhar/SSI registered bidder wishes to avail above facility then they should follow necessary processes with E Procurement Portal www.mptenders.gov.in and if necessary, take help of help line on E Procurement Portal.	Request for Relaxation of Norms as per Policy of Government of India there is a provision to relax the norms Related to Prior Experiences - Prior Turnover and Exemption for Bid Security / EMD ,As per attached memorandums of Govt. Of India , Department of Expenditure Procurement Policy Division.	Not Accepted
2	2. ELIGIBILITY CRITERIA 2. (d). The Analytical Laboratory should have achieved an average annual turnover of at least Rs.25 Lakhs during last three years i.e. 2015-16, 2016-17 and 2017-18. The Government Laboratories, Research & Development Laboratories, Laboratories run by Co-operative body & Educational Institutions are exempted from the turnover criteria. 2.(c). The Analytical Testing Laboratory should have at least 3 (Three) years experience in the analysis of drugs (biological / Chemical). The bidder should provide details of past experience in the proforma given in Annexure - I.	We are a indore based testing lab. We have started operation of this lab in 16-17 hence we have market standing of 3 years i.e 2016-2017 , 2017-18 & 2018-19. As 2018-2019 year is over this may also been included for the Annual statement. We can give the CA certificate of all the 3 years. Kindly amend the tender format . We request the committee to look into the matter.	Not Accepted
3	2. ELIGIBILITY CRITERIA a) The Analytical Testing Laboratory should be accredited by "National Accreditation Board of Laboratories" (NABL) and such accreditation should be valid on the date of submission of Tender. Tenderer should submit copy certificate of Accreditation issued by NABL (duly certified by a Chartered Accountant) for Chemical/ Biological testing including testing of drugs and medical supplies as the case may be.	In Addition to that "Analytical laboratory must be approved/registered by USFDA or WHO: Pre qualified.	Not Accepted
4	12. Description of Services and Performance Requirements: d) The laboratory should report on packaging and general description of the product. All the tests mentioned in IP/BP/USP/Drugs & Cosmetics Act/BIS/ISO. Etc., (as the case may be) should be carried out for each and every sample. The results obtained in the test should be mentioned in figure (wherever possible). In case of a non-pharmacopoeial drug, the laboratory should follow standards and validated protocols as per ICH guidelines including stability indicating assays and provide the details with the report. In all those cases where sustained release or delayed release profiles are claimed, dissolution should invariably be performed.	All the List of Drug Items should be complying to respective Pharmacopoeia (IP/BP/USP/Drugs & Cosmetics Act/BIS/ISO. Etc.) or In House Method. If it is In-House Method then MOA (Method of Analysis) / Specification to be Provided of respective Drug of Item.	Not Accepted

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CGM (T)
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