Corrigendum

AIIMS Rishikesh tender Enquiry No. T.No 24/RC/Medicines/Pharmacy-604/2021-RISH(Admn)
"Rate Contract document for procurement of **Neoplastic Disorders Medicines / Immunomodulators, Immunosuppressants/ Antiemetics** for Central Pharmacy AIIMS, Rishikesh"

S.No.	Original point	Revised (To be read as)
1.	**Instruction to Bidders; Point1. c.** No third-party manufacturers are allowed to quote for any drug.	Bidders with products manufactured on P2P basis from third party but having quality control of such products by in-house Quality Assurance Department are eligible to participate in tender.
2.	**Technical evaluation; Point f.** Bidder(s) shall submit a Manufacturing & Market Standing Certificate / Experience Certificate issued by the Centre / State Drug Licensing Authority of the respective state that quoted product is manufactured and marketed by them since last 05 (Five) years.	Bidder(s) shall submit a Manufacturing & Market Standing Certificate / Experience Certificate issued by the Centre / State Drug Licensing Authority of the respective state that quoted product is manufactured and marketed by them since last 03 (three) years. Any firm who took over another manufacturer portfolio in past 3 years from date of publishing of this tender shall submit Manufacturing & Market Standing Certificate / Experience Certificate issued by the Centre / State Drug Licensing Authority stating time duration for which quoted product(s) was manufactured by either manufacturers totalling for 03 (three) years. In such scenario, current manufacturer must fulfil other eligibility and technical criteria of tender document.
3.	**Technical Evaluation Point E:** Bidder must have recent experience, within last two years, for supplying salt, for which bid submitted, to developed countries	Bidder must have recent experience, within last two years, for supplying salt, for which bid submitted, to developed country(ies). For imported drug manufacturer must be Supplying/ selling the product to/ in developed country(ies).
4.	**Technical Evaluation; Point j.:** Projection Capacity Assessment Certificate: Manufacturing firm must enclose certificate issued by Chartered Accountant/ concerned State Drug Controller indicating actual production detail of a particular molecule, batch-wise, for items quoted.	Capacity Assessment Certificate: Manufacturing firm must enclose certificate as per **Annexure B (Attached below)** ***If any bidder already got prepared the format as per earlier clause with all required information for last three financial years** then it will also be accepted.
5.	**Technical Evaluation; Point i.:** **Quality Related Action taken by drug regulatory authority of India** Bidder has to provide certificate issued from regulatory authority, India that no warning has been issued or action has been taken in last 5 years, in terms of quality of production and produced batches.	**Technical Evaluation; Point i.:** **Quality Related Action taken by drug regulatory authority of India** Bidder has to provide certificate issued from regulatory authority, India that no warning has been issued or action has been taken in last 5 years or from date of approval to market, whichever is latest, in terms of quality of production and produced batches.
6.	**Financial Evaluation; Point c:** Supplier has to submit a notarized affidavit on Indian Non Judicial Stamp Paper of Rs.10/- that bidder has not quoted price higher than previously supplied to any government Institute/ Organization/reputed Private Organization or	Supplier has to submit a notarized affidavit on Indian Non Judicial Stamp Paper of Rs.10/- that bidder has not quoted price higher than previously supplied to any government Institute/ Organization/ reputed Private Organization or DGS&D rate in *last three years or since*

	DGS&D rate in recent past. Therefore, if at any stage it is found that supplier has quoted lower rates than those quoted in this tender; Institute (purchaser) would be given benefit of lower rates by Supplier and any excess payment if any, will become immediately payable to AIIMS, Rishikesh. If such affidavit is not submitted, tender will be summarily rejected. (Part of technical bid)	*the last change in M.R.P. whichever is latest*. Therefore, if at any stage it is found that supplier has quoted lower rates than those quoted in this tender; Institute (purchaser) would be given benefit of lower rates by Supplier and any excess payment if any, will become immediately payable to AIIMS, Rishikesh. If such affidavit is not submitted, tender will be summarily rejected. (Part of technical bid)
7.	Special Conditions of Contract (SCC) / Contract terms; Labelling; Point 5a. Before making supply, approved rate contract holder should ensure that all primary packaging labels (i.e. Ampoules, vials, bottles, jars, tubes etc.) should be embossed, imprinted, stamped with letters, or requirements like "AIIMS-RISHIKESH SUPPLY NOT FOR SALE" stamp with permanent ink on each item/strip up to primary level.	Before making supply, approved rate contract holder should ensure that all primary, secondary and tertiary packaging labels (i.e. Ampoules, vials, bottles, jars, tubes etc.) should be embossed/ imprinted with barcode carrying information about batch number, date of manufacture, expiry date etc. Packing should have labels mentioning "AIIMS-RISHIKESH SUPPLY NOT FOR SALE" in permanent ink on each item/strip up to primary level.
8.	**Eligibility criteria cum Administrative cum Technical Evaluation Checklist (Annexure I) :**	
	12. Please provide a notarized affidavit on Non Judicial stamp paper of Rs. 10/- in format given in Annexure-V that you have not quoted price higher than previously supplied to any government Institute/I/reputed Private I or DGS&D/GeM rate in last three years. If you don't fulfil this criterion, your tender will be out rightly rejected.	12. Please provide a notarized affidavit on Non Judicial stamp paper of Rs. 10/- in format given in Annexure-V that you have not quoted price higher than previously supplied to any government Institute/reputed Private Institute or DGS&D/GeM rate in *last three years or since the last change in M.R.P. whichever is latest*. If you don't fulfil this criterion, your tender will be out rightly rejected. * If the quoted product M.R.P. was changed within preceding three years of date of tender publish, then bidder has to submit documents as evidence of approval of such change from drug regulatory authority.
	21. Please submit duly filled Bid Security Declaration Form on letter head.	21. Invoice against claim of supply/ sell of quoted molecules to developed country (ies)
	22. Details of DD No of Tender fee: Date: Payable at:	22. In-house quality assurance lab with accreditation certificate in name of bidder in reference to Instruction to Bidders; revised Point1. c.
	-	23. Please submit duly filled Bid Security Declaration Form on letter head.
	-	24. Details of DD No of Tender fee: Date: Payable at:
9.	**Annexure III:** I agree to maintain the following information in the Company's record for a period of 8 years and shall make this available for verification to any statutory authorities	I agree to maintain the following information in the Company's record for a period till 01 (one) year after all contractual obligations are over and shall make this available for verification to any statutory authorities
10.	Last date of Bid Submission 29/04/2021	Last date of Bid Submission 10/05/2021
11.	Date of Bid Opening 30/04/2021	Date of Bid Opening 11 /05 /2021

Annexure B

CAPACITY ASSESSMENT CERTIFICATION

No.:

1. Name of the firm:
Address_____
Telephone_____
Email Id: _____

2. _____ (Name of Manufacturer) is properly registered to supply pharmaceuticals or vaccines or medical devices in India, is in good legal and statutory standing with the responsible health authorities and is licensed as a primary manufacturer of the range of pharmaceuticals or vaccines or medical devices to be offered.

3.The production capacities for _____ (name of Manufacturer) is as follows:

S.No. (As per tender document)	Medicine Name	Formulation	Strength	Production Capacity in Financial Year		
				2017-18	2018-19	2019-20

Certified by Charted Accountant
(Signature & Seal with date)

4. _____ (Name of Manufacturer) retains full records of production batches and quality control test results, has demonstrated compliance with the M GMP quality standards during the past three years, and will exhibit these on request.

5. We hereby certify that the above information is true and accurate to the best of our knowledge.
Signature & Seal of the Authority of company _____
Date:
Full name (Printed) _____
Position _____