CENTRAL MEDICAL SERVICES SOCIETY

Ministry of Health and Family Welfare (Government of India)
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Dated 06.02.2023

AMENDMENT NO. 03

Tender No- CMSS/PROC/2022-23/NVBDCP/030 for Procurement of Cap. Miltefosine 50mg & Cap. Miltefosine 10mg for NVBDCP.

The following amendment in the subject bid document is made.

S. No.	Query Received from Prospective Bidders	Standard Tender Clause	Response
A	**M/s Centurion Laboratories Pvt. Ltd.**		
1.	With ref. to prebid report clause-1 of miltefosine cap's procurement enclosed , we came to know that you have given in eligibility criteria clause 4, to give order only to bidder having form -10, to import API - miltefosine issued by DCGI , but we have developed (arranged to get) this API mfged by local API drug manufacturer M/s Biopin-Aambal Life Sciences Private Limited, K 39, BIDC Industrial estate, Gorwa, Vadodara 390016 under 'ATMANIBHAR BHARAT' slogan of our PM , hence pl. amend eligibility creteria clause no 4 , to allow locally manufactured API,-	**Eligibility Criteria:** **Clause 4 (b)** Tenderer shall be a manufacturer of quoted product and having valid own manufacturing license for the offered product and it should comply as per technical specifications. The manufacturing license should be valid on the date of tender opening packet - 1. The raw material/API for manufacturing of Cap. Miltefosine shall be imported by way of Import License (Form-10 from DCGI). NCVBDC shall if required, facilitate the vendor in procuring import licenses (Form 10) from DCGI after award of LoA.	Amended as tender clause 4 (b); Tenderer shall be a manufacturer of quoted product and having valid own manufacturing license for the offered product and it should comply as per technical specifications. The manufacturing license should be valid on the date of tender opening packet - 1. The raw material/API for manufacturing of Cap. Miltefosine (i) Shall be imported by way of Import License (Form-10 from DCGI). NCVBDC shall if

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			available in any of the official pharmacopeia, 'In House' standards are applicable as per Drugs and Cosmetics Act 1940 and Rules there under. **Bidder is requested to submit an undertaking that the drug is not available in IP or any other approved pharmacopeia.** The manufacturing License should be valid on the date of tender opening packet 1. And Submit an undertaking for the requirement of facilitation for obtaining import License (Form 10) from CDSCO and requesting to CMSS/ Programme Division to facilitate the same. OR In case the API is domestically sourced, the Manufacturing License of the domestic manufacturer of the API issued by the relevant competent authority has to be submitted with the technical bid and same should be valid on date of tender opening.

Rest all Terms and Conditions remains same.

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General Manager (Procurement)