



# Rajasthan Medical Services Corporation Limited

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Ref. No.:- F.02(219)/RMSC/PROCUREMENT/DRUG/NIB-22/2016/109

Dated: 27/01/2017

## Corrigendum – 2

Subject :- Amendments in technical specifications, conditions and date extension

Ref. No. :- Bid Ref. No. F.02(219)/RMSC/PROCUREMENT/DRUG/NIB-22/2016/1597 Dated:29.12.2016

and

Corrigendum-1 Ref. No. F.02(219)/RMSC/PROCUREMENT/DRUG/NIB-22/2016/77

Dated:20.01.2017

S. No	Existing condition / technical specification/Packing Unit/Quantity (clause no.)	Amended condition / technical specification/ Packing Unit/Quantity/Shelf Life/Date Extension (clause no.).																
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*[Handwritten signatures and marks]*

5	<p><b>Clause 5(i) TECHNICAL BID</b>          WHO-GMP (WHO - Good manufacturing practices Certificate) Certificate issued by the Licensing Authority. The WHO-GMP certificate must not be older than one year from the due date of Bid submission in the case where validity is not mentioned in the certificate. The WHO-GMP certificate of all the manufacturing plants, of which products have been quoted, should be submitted. The Bidder shall also furnish an undertaking in the format given in Annexure-VII point no.8 declaring that the Bidder complies with the requirements of WHO-GMP. The Importer should produce WHO- GMP /COPP of the manufacturing firm or a certificate which is at par with WHO-GMP issued by exporting countries like US-FDA approval, etc. In the case of imported drugs, labels and product literature of all quoted products must be submitted.</p>	<p><b>Clause 5(i) TECHNICAL BID</b>  <b>In the existing condition of WHO-GMP in the clause, it is relaxed to GMP for below mentioned item codes :-</b></p> <table border="1" style="width: 100%;"> <thead> <tr> <th style="width: 15%;">Code No.</th> <th>Name of item with specification</th> </tr> </thead> <tbody> <tr> <td style="text-align: center;">440</td> <td>Dextromethorphan Hydrobromide Syrup IP 13.5mg / 5ml</td> </tr> </tbody> </table> <p>GMP (Good manufacturing practices Certificate) Certificate issued by the Licensing Authority. The GMP certificate must not be older than one year from the due date of Bid submission in the case where validity is not mentioned in the certificate. The GMP certificate of all the manufacturing plants, of which products have been quoted, should be submitted. The Bidder shall also furnish an undertaking in the format given in Annexure-VII point no.8 declaring that the Bidder complies with the requirements of GMP. The Importer should produce WHO- GMP /COPP of the manufacturing firm or a certificate which is at par with WHO-GMP issued by exporting countries like US-FDA approval, etc. In the case of imported drugs, labels and product literature of all quoted products must be submitted.</p>	Code No.	Name of item with specification	440	Dextromethorphan Hydrobromide Syrup IP 13.5mg / 5ml
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**Note:-**

- It may be noted that if any type of amendments required than further corrigendum will be published and informed.
- Rest terms and conditions will remain the same.

*Handwritten initials/signature*

  
**Executive Director (Proc.)**  
**RMSC**