



# Rajasthan Medical Services Corporation Limited

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Ref. No.:- F.02(278)/RMSCL/PROCUREMENT/DRUG/NIB-03/2020/311

Dated:-19.02.2020

## Corrigendum – II

Subject:- Amendments in BOQ/ technical specification/Packing Unit/Quantity.

Ref.:- NIB No. F.02(278)/RMSCL/PROCUREMENT/DRUG/NIB-03/2020/112 Dated:-17.01.2020

(Technical bid opening due on dated –25.02.2020)

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	<p>Clause 2(c) ELIGIBILITY CRITERIA            Clause 5 (i) (j) TECHNICAL BID            Annexure-XV Ref. Clause No. 2 (c)</p> <table border="1" data-bbox="194 454 713 636"> <thead> <tr> <th>Code No.</th> <th>Name of item with specification</th> <th>Packing Unit</th> </tr> </thead> <tbody> <tr> <td>617</td> <td>Budesonide Powder for Inhalation IP 200 mcg</td> <td>30 Capsule (rotacap)</td> </tr> </tbody> </table> <p>Clause 5(i) TECHNICAL BID</p> <p>(l) 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.</p> <p>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>The Firm will continue to hold WHO-GMP Certificate for the product during entire rate contract period of the product. If WHO-GMP certificate expires, it is firm's responsibility to inform RMSCL about the same and not to accept any further purchase order till re-issue /renewal of WHO-GMP certificate. During the period of non validity of WHO-GMP certificate of the firm the rate contract will deemed to be suspended. If the firm fails to inform RMSCL about the expiry of WHO-GMP certificate and accept purchase order of RMSCL and later on it comes to the knowledge of RMSCL, in this situation firm shall be liable for a panel action.</p>	Code No.	Name of item with specification	Packing Unit	617	Budesonide Powder for Inhalation IP 200 mcg	30 Capsule (rotacap)	<p>Clause 2(c) ELIGIBILITY CRITERIA            Clause 5 (i) (j) TECHNICAL BID            Annexure-XV Ref. Clause No. 2 (c)</p> <table border="1" data-bbox="825 443 1367 591"> <thead> <tr> <th>Code No.</th> <th>Name of item with specification</th> <th>Packing Unit</th> </tr> </thead> <tbody> <tr> <td>617</td> <td>Budesonide Powder for Inhalation IP 200 mcg</td> <td>30 Capsule</td> </tr> </tbody> </table> <p>Market Standing Certificate and Performance Statement of Two year acceptable.</p> <p>Clause 5(i) TECHNICAL BID</p> <p>In the existing condition of WHO-GMP in the clause, it is relaxed to GMP for below mentioned item codes :-</p> <table border="1" data-bbox="848 842 1433 1267"> <thead> <tr> <th>Code No.</th> <th>Name of item with specification</th> <th>Packing unit</th> </tr> </thead> <tbody> <tr> <td>399</td> <td>Concentrated Solution for Haemodialysis B.P Acetate concentrate in 10 Litre Cans.</td> <td>10 Ltrs Plastic Can</td> </tr> <tr> <td>687</td> <td>Concentrated Solution for Haemodialysis B.P Sodium Hydrogen carbonate Concentrate (Part A and Part B)</td> <td>Part A (10 Ltrs Plastic Can) Part B (Two Packet of 820 to 900 gm each )</td> </tr> </tbody> </table>	Code No.	Name of item with specification	Packing Unit	617	Budesonide Powder for Inhalation IP 200 mcg	30 Capsule	Code No.	Name of item with specification	Packing unit	399	Concentrated Solution for Haemodialysis B.P Acetate concentrate in 10 Litre Cans.	10 Ltrs Plastic Can	687	Concentrated Solution for Haemodialysis B.P Sodium Hydrogen carbonate Concentrate (Part A and Part B)	Part A (10 Ltrs Plastic Can) Part B (Two Packet of 820 to 900 gm each )
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**Note:-**

- It may be noted that if any further amendments are issued then a corrigendum will be published and informed.
- Rest of the terms and conditions will remain the same.

*(Handwritten initials)*

*(Signature)*  
 Executive Director (Proc.)  
 RMSCL