



Rajasthan Medical Services Corporation Limited

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Ref. No.:- F.02(227)/RMSC/PROCUREMENT/DRUG/NIB-07/2017/ 916

Dated: 08.08.2017

Corrigendum - 1

Subject:- Amendments in technical specifications and conditions etc.

Ref. -Pre bid meeting dated 01.08.2017 for the Bid reference no F.02(227)/RMSC/ PROCUREMENT/DRUG/NIB-07/2017/879 Dated:21.07.2017 (Technical bid opening due on dated - 22.08.2017)

S. No	Existing condition / technical specification/Packing Unit/Quantity (clause no.)	Amended condition / technical specification/ Packing Unit/Quantity/Shelf Life/Date Extension (clause no.).								
1.	<p>Clause 5(I) TECHNICAL BID 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>Clause 5(I) TECHNICAL BID In the existing condition of WHO-GMP in the clause, it is relaxed to GMP for below mentioned item codes :-</p> <table border="1"> <thead> <tr> <th>Code No.</th> <th>Name of item with specification</th> </tr> </thead> <tbody> <tr> <td>242</td> <td>VDRL Antigen (with +ve and -ve control) / RPR slide Kit</td> </tr> <tr> <td>399</td> <td>Concentrated Solution for Haemodialysis B.P Acetate concentrate in 10 Litre Cans.</td> </tr> <tr> <td>687</td> <td>Concentrated Solution for Haemodialysis B.P Sodium Hydrogen carbonate Concentrate (Part A and Part B)</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	242	VDRL Antigen (with +ve and -ve control) / RPR slide Kit	399	Concentrated Solution for Haemodialysis B.P Acetate concentrate in 10 Litre Cans.	687	Concentrated Solution for Haemodialysis B.P Sodium Hydrogen carbonate Concentrate (Part A and Part B)
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(Handwritten signatures and initials)

2.	BoQ:- Column number - 18 Column number - 19 Column number - 11 and TOTAL AMOUNT With Taxes (Total of 6+8+10)	BoQ:- Read as Column number - 11 Read as Column number - 12 Read as Column number - 13 and TOTAL AMOUNT With Taxes (Total of 6+8+10 or 6+12)
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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.

H → J R


Executive Director (Proc.)
RMSC