

Rajasthan Medical Services Corporation Limited

Gandhi Block, SwasthyaBhawan, Tilak Marg, C-Scheme, Jaipur-302005 (Raj.)

Phone No: 0141-2228066 .2228064 Fax No. 0141-2228065 Website: http://rmsc.health.rajasthan.gov.in

CIN:U24232RJ2011SGC035067

E_mail:edprocurement@gmail.com, andrmsc@nic.in

Ref. No.:- F.02(275)/RMSCL/PROCUREMENT/DRUG/NIB-17/2019/07

Dated: - 06.01.2020

Corrigendum - I

Subject:-Amendments in BOQ/ technical specification/Eligibility Criteria.

Ref.:- Prebid meeting dated 26.09.2019 and NIB No. F.02(275) / RMSCL / PROCUREMENT / DRUG / NIB-17 / 2019 / 1559 Dated:-20.12.2019 (Technical bid opening due on dated -23.01,2020)

	23.01.2020)		
S.	Existing condition / technical specification/Packing	Amended condition / technical specification/ Packing	
No	Unit/Quantity (clause no.)	Unit/Quantity/Shelf Life/Date Extension (clause no.).	
1,	Clause 2(e) ELIGIBILITY CRITERIA	Clause 2(c) ELIGIBILITY CRITERIA	
	Clause 5 (i) (j) TECHNICAL BID	Clause 5 (i) (j) TECHNICAL BID	
	Annexure-XV Ref. Clause No. 2 (c)	Annexure-XV Ref. Clause No. 2 (c)	
	Code Name of item with specification	Code Relaxation in Condition of Market Standing	
	No.	No. Certificate and Performance Statement	
	NE16 Dual Rapid Test kit for HIV & Syphilis	*NE16 Firm has to submit Market Standing Certificate	
	NEIO Dual Rapid Test Rit for Til v & Syphins	of Dual Rapid Test kit for HIV & Syphilis for 6	
		months and Performance Statement of at least	
		one batch in financial year 2018-19 or 2019-20.	
		OR	
		Firm has to submit Market Standing Certificate	
		of both test kits i.e HIV and Syphilis for 6	
		months and Performance Statement of at least	
		one batch in financial year 2018-19 or 2019-20.	
		*However, for all above items, the firm has to submit	
		with bid, the product permission (from the Licensing	
		Authority) same as per bid specifications.	
2.	Clause 5(I) TECHNICAL BID	Clause 5(I) TECHNICAL BID	
	WHO-GMP (WHO – Good manufacturing practices	In the existing condition of WHO-GMP for	
	Certificate) Certificate issued by the Licensing	manufacturer in the clause, it is relaxed to QMS	
	Authority. The WHO-GMP certificate must not be	Certificate / ISO 13485 for below mentioned item	
	older than one year from the due date of Bid	codes:-	
	submission in the case where validity is not mentioned	Code Name of item with specification	
	in the certificate. The WHO-GMP certificate of all the	No.	
	manufacturing plants, of which products have been	NE16 Dual Rapid Test kit for HIV & Syphilis	
	quoted, should be submitted. The Bidder shall also		
	furnish an undertaking in the format given in	As item included under Medical Device Rules 2017 hence	
	rathon an undertaking in the format given in	for manufacturer QMS Certificate issued by the competent	

authority OR ISO 13485 issued by CDSCO registered notified body acceptable. The QMS Certificate / ISO 13485 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 QMS Certificate / ISO 13485 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 QMS Certificate / ISO 13485. 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.

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Annexure-VII point no.8 declaring that the Bidder

complies with the requirements of WHO-GMP. The

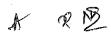
Importer should produce WHO- GMP /COPP of the

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.

	S. Existing condition / technical specification/Packing No Unit/Quantity (clause no.) Amended condition / technical specification/ Packing Unit/Quantity/Shelf Life/Date Extension (clause no	ng
3		dicines he last 7-18 or ss than annual 2016- should ling in Eye 72,588 of last Crore.
4	Annexure - VIII (List of Drugs with Specifications) and BoQ Annexure - VIII (List of Drugs with Specifications) BoQ	s) and
	Code No. No. Code Name of item with specification No. Code Name of item with specification No. Code No. Code Name of item with specification No. 682 Teneligliptin Tablet IP 20 mg	
	Notes	

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.

Executive Director (Proc.)
RMSCL

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