

Madhya Pradesh Public Health Services Corporation Ltd.

Corrigendum No 03 for PreBid Query Resolution Tender No.02/Pharma

S. No.	Item Name	Drug Code	Unit (SKU)	Quantity of Supply in SKU	Recommendation from bidder with Justification	Proposed Resolution/Amendment
1	Hydroxy Propyl Methyl Cellulose Injection 2%	120855	5ml Prefilled Syringe	30000	3 ml Prefilled Syringe instead of 5 ml Prefilled Syringe, 3 ml Prefilled Syringe, It is mentioned as 5 ML Prefilled Syringe in the Tender. The normal industry practice is either 2 ML or 3 ML only.	Specification amended as Hydroxy Propyl Methyl Cellulose Injection 2% 3ml Prefilled Syringe
2					In given tender you have said that if any product is of NSQ for 3 times the company would be blacklisted, regarding this we want to put a point that if same product is of NSQ and it fails for 3 times then only you should blacklist the company	No Change
3					Please refer to tender no. 002 for drugs. We would like to please attention on submitting bank guarantee, you have given time for only 7 days which is practically not possible because the amount value is high bank takes its own time in issuing the BG, hence it's a request to kindly look into the matter and change the maximum day criteria to 30 days.	No Change
4	Acyclovir tab. IP - 200mg	110167	10x10	9980	Item specification allow to quote:- ACICLOVIR 200 mg DT Tab.	DT Tablet also acceptable
5	Acyclovir (800mg), Tablet	110168	10x10	7988	Item specification allow to quote:- ACICLOVIR 800 mg DT Tab.	DT Tablet also acceptable
6	Micronised Progesterone(100 Mg), Tablet	120702	10x10	1272	Item specification allow to quote:- MICRONISED PROGESTERONE(100MG), CAPSULES.	No Change
7	Risperidone Tab 2 mg	110359	10x10	2000	Item specification allow to quote:- RISPERIDONE 2MG MD TAB.	DT Tablet also acceptable
8	Salbutamol Nebulizing solution(5 mg/ 2.5 ml), Ampule	110442	2.5 ml Amp	1610800	Item specification allow to quote:- SALBUTAMOL NEBULIZING SOLUTION (5 MG/15 ML), BOTTLE.	No Change

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9	FACTOR VIII with Vwf	120862	250 IU/Vial	4500	Request for review and revision: In case wherein bidder is - the Indian subsidiary of a global manufacturer, turnover of the parent global company should be considered. In our case, the turnover statement of Grifols S.A. may be considered, as Grifols India is newly setup Indian subsidiary. - OR please allow authorized distributor (of Indian subsidiary), to quote/bid with relevant documents to meet above requirements.	No Change
10					Request for review and revision: in case wherein bidder is the Indian subsidiary of a global manufacturer, turnover of the parent global company should be considered. In our case, the turnover statement of Grifols S.A. may be considered, as Grifols India is newly setup Indian subsidiary. - Exemption of the “Eligibility Criteria – Clause h” , since the product has not been commercially sold / available in India. -Or Required certificate may be submitted for current year.	No Change
11					Request for review and revision: In case wherein bidder is the Indian subsidiary of a global manufacturer, turnover of the parent global company should be considered. In our case, the turnover statement of Grifols S.A. may be considered, as Grifols India is newly setup Indian Subsidiary. OR please allow authorized distributor (of Indian subsidiary), to quote/bid with relevant documents to meet above requirements.	No Change

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12					<p>Request for review and revision: In the case wherein the product is manufactured by an internationally renowned organization, with safety & efficacy data, but not commercially available in India for preceding 3 years, - Exemption of the clause “TECHNICAL BID 4.1.n Cover A” , since the product has not been commercially sold / available in India. - Or Global Turnover certificate may be submitted by the parent organization.</p> <p>- Documented evidence of use of the item to be quoted, on India population / patients through humanitarian aid programs sponsored by World Federation of Hemophilia may be considered.</p>	No Change
13					<p>Request for review and revision: In the case wherein the product is manufactured by an internationally renowned organization, with safety & efficacy data, but not commercially available in India for preceding 3 years, the following may be considered: - Exemption of the “Eligibility Criteria – Clause e (i)/(ii)” , since the product has not been commercially sold / available in India for preceding 3 yrs. - Global Marketing / Registration Certificate for the item to be quoted.</p> <p>- Documented evidence of use of the item to be quoted, on Indian population / patients through Humanitarian Aid programs sponsored by WFH (World Federation of Hemophilia).</p> <p>- Published Clinical efficacy & safety data of the item to be quoted, on Indian population.</p>	No Change

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S. No.	Item Name	Drug Code	Unit (SKU)	Quantity of Supply in SKU	Recommendation from bidder with Justification	Proposed Resolution/Amendment
14	Rituximab -100mg Inj	120268	Vial	4500	In Online price bid only Basic rate and GST Column is there but our company plant is in SEZ area so Basic custom duty and Social Welfare Tax is also levied other than GST. Kindly change in online price bid accordingly.	No Change
15	Rituximab -500mg Inj	120269	Vial	4500	In Online price bid only Basic rate and GST Column is there but our company plant is in SEZ area so Basic custom duty and Social Welfare Tax is also levied other than GST. Kindly change in online price bid accordingly.	No Change
16	Rituximab -100mg Inj	120268	Vial	4500	1. QUANTITY SHOULD BE DECREASED AS PER LAST YEAR CONSUMPTION 2. TENDER QUOTATION SHOULD BE BY MANUFACTURER AND BILLING ALLOWED FROM SISTER CONCERN COMPANY WHERE MANUFACTURING COMPANY HAS GIVEN MARKETING RIGHT TO SISTER CONCERN COMPANY	Quantity Ameded as 3000, No other changes
17	Rituximab -500mg Inj	120269	Vial	4500	1. QUANTITY SHOULD BE DECREASED AS PER LAST YEAR CONSUMPTION 2. TENDER QUOTATION SHOULD BE BY MANUFACTURER AND BILLING ALLOWED FROM SISTER CONCERN COMPANY WHERE MANUFACTURING COMPANY HAS GIVEN MARKETING RIGHT TO SISTER CONCERN COMPANY	Quantity Ameded as 3000, No other changes

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S. No.	Item Name	Drug Code	Unit (SKU)	Quantity of Supply In SKU	Recommendation from bidder with Justifiacion	Proposed Resolution/Ame ndment
18	Surgical Spirit IP	120599	50ml Bottle	230107	<p>We request you to modify the eligibility criteria for the product Surgical Spirit IP (Drug Code- 120599) with the following pionts:</p> <ol style="list-style-type: none"> 1. Relax from WHO-GMP to GMP. 2. Turnover criteria- From Rs 10Cr to Rs 2Cr. The Surgical spirit comes under disinfectant category various state governments across the country have relaxed turnover criteria. 3. Delivery schedule 60 days the state govt like Telangana Govt, Odisha govt, have issued time to supply 60 to 70 days from the date of purchase order has the product we are dealing with surgical spirit IP having 95% alcohol content it requires lot of procedure to follow up its take time to procure. Hence we request extend the time to 70 days and oblige. 	Pack amended as 500ml Bottle, No other changes.
19	Aluminium Hydroxide+Magnesium aluminium silicate+Magnesium Hydroxide+Simethecon(Tab 300 mg+50mg + 25 mg+25 mg),Tablet	110439	10x10	2000		Item code 110439 is deleted since Antacid EDL item is available in Live RC
20					<p>As per your NIT Tender Ref Number MPPHSCL 002 for drugs due dt 11. 02. 2019 . pre bid queries .</p> <p>In your NIT Point number 7. In case any one batch is found NSQ than particular product of the firm will be black listed/debarred for not less than two year. Upon blacklisting / debarrement of such 3 products or found such 3 NSQ batches(of one or more products) under a tender then firm will be blacklisted not less than 3 years.</p> <p>Please clearly the metter.</p>	No Change

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21					<p>In case any one batch is found NSQ than particular product of the firm will be blacklisted/debarred for not less than two year. Upon blacklisting / debarment of such 3 products or found such 3 NSQ batches(of one or more products) under a tender then firm will be blacklisted not less than 3 years.</p> <p>7. In case any three batch is found NSQ than particular product of the firm will be black listed/debarred for not less than two year. Upon blacklisting / debarment of such 3 products or found such 3 NSQ batches(of one or more products) under a tender then firm will be blacklisted Generally accepted condition of other tender authorities</p>	No Change
22	Trihexyphenidyl 2mg tab	110191	10x10	3000	<p>Please consider per tablet cost while evaluating the price as our commercial pack size available as 30 Tablets in a Strip (80 X 30`s= 2400 Tablets in box)</p>	Refer clause number 10.15
23	Acetazolamide Tab(250mg),Tablet	110263	10x10	12186	<p>Please consider per tablet cost while evaluating the price as our commercial pack size is available as 15 Tablets in a Strip (36 x 15`s= 540 Tablets in box)</p>	Refer clause number 10.15
24	Phenytoin Sodium Oral Suspension - 25 mg/ml(100 ml Bottle),Suspension	110065	100ml Bottle	16552	<p>Requesting you to consider loan license for this product. As this preparation is available only few manufacturers fulfilling the technical requirement .</p>	No Change
25	Recombinant FVIII	120266	500 IU/Vial	6000	<p>Ø Presence in the Global Market more than 10 years</p>	As per tender terms only
26	Recombinant FVIII	120267	1000 IU/Vial	7500	<p>Ø USFDA Approval As our drug falls under the category of “New Drug as defied by CDSCO” – Kindly consider to waive the following</p> <p>Ø USFDA Approval As our drug falls under the category of “New Drug as defied by CDSCO” – Kindly consider to waive the following</p> <p>Ø Market Standing cum Performance Certificate</p> <p>Ø Bill of lading/Bill of Entry for last 3 years</p> <p>Ø Certificate of Analysis</p> <p>Ø Annexure III</p>	As per tender terms only

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27	Recombinant FVIII	120265	250 IU/Vial	6000	Relaxation on the Market standing clause of 3 years being new drug category. Form 45 , enclosed –issued from DCGI	As per tender terms only
28	FACTOR VIII with Vwf	120862	250 IU/Vial	4500	Change specification to Plasma derived Factor VIII 250 IU or Anti Hemophillic Factor VIII because as per Indian Pharmacopeia (I.P.) product description is “Dried Human Anti hemophilic Fraction. In the product description I.P. states that ‘ Dried Human Anti-hemophilic Fraction is a preparation of a plasma proteins fraction that contains the glycoprotein coagulation factor VIII together with varying amount of Von Willebrand Factor (Vwf) , depending upon method of preparation.” In the I.P. labeling guidelines Vwf content is not included separately . All the tender floated earlier by MPPHSCL has Anti Hemophillic Factor VIII 250 IU and 500 IU. And also add Plasma derived Factor VIII 500 IU / Anti Hemophillic Factor VIII 500 IU which is already in all the pervious tender flotted by MPPHSCL.	Specification amended as Plasma derived FACTOR VIII for Item Code 120862. Plasma derived Factor VIII 500 IU / Anti Hemophillic Factor VIII 500 IU will be included in next tender.
29	Trastuzumab Inj. 440 mg	120837	Vial	7500	Tender quantity is very high. It should be 1200	No Change
30	Rituximab -100mg Inj	120268	Vial	4500	Tender quantity is very high. It should be 2500	Quantity Ameded as 3000 for Item code 120268
31	Rituximab -500mg Inj	120269	Vial	4500	Tender quantity is very high. It should be 2500	Quantity Ameded as 3000 for item code 120269
32	Calamine Lotion IP (contains per 1000 ml:- calamine 150 gm,Zinc oxide 50 gm, Bentonite 30 gm, Sodium Citrate 5gm, liquified phenol 5ml, glycerin 50 ml purified waterfreshly boiled and cooled to produced 1000ml) 50 ml Bottle	110258	50ml Bottle	220320	Also exempt WHO GMP critieria to GMP certification in view of external application item.	No Change

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33	Surgical Spirit IP	120599	50ml Bottle	230107	In earliest tender it was procured in ml but now it was stated as 50ml request to amend the same to 500ml Also exempt WHO GMP criteria to GMP certification in view of external application item.	Specification amended as Surgical spirit IP 500ml Bottle for Item code 120599, No other change.
34					Sir, above application are ment for external usage and WHO GMP is esstential for latest molecules/internal usage/Indigenous consumption but not particular application The bidding prices of WHO GMP unit are expected to be abnormally higher when compared to the prices of GMP certified units, especially for these particular items.	No Change