



Tender enquiry no. Store/Tender/Drugs & Consumable for OT/1/2016 dated 19-05-2016

Corrigendum

Sr. No.	Page No/ Clause no/ point no	Existing sentence	To be read as				
1	As mentioned tender document	Last date of submission: 25-05-2016 at 3:00 PM, technical bid will be opened on the same date at 3:30 PM	Last date of submission: 07-06-2016 at 3:00 PM, technical bid will be opened on the same date at 3:30 PM				
2	Page no. 2 Para 1	I/We, the under signed, here by submit my/our tender for the Registration of firm / company for supply of <u>Drugs & consumables items for OT</u> on two years rate contract basis.	I/We, the under signed, here by submit my/our tender for the Registration of firm / company for supply of <u>Drugs & consumables items for OT</u> on three years rate contract basis The contract may be extended for further period of one year on mutually agreed terms.				
3	Page no. 7 Point no. 21	The successful tenderer will be required to furnish a Performance Security Deposit of 10% of contract amount in the form of Demand Draft, Fixed Deposit Receipt or Bank Guarantee from any Scheduled Bank...	The successful tenderer will be required to furnish a Performance Security Deposit of 05% of contract amount in the form of Demand Draft, Fixed Deposit Receipt or Bank Guarantee from any Scheduled Bank...				
4	Page no. 29 schedule-1 point no. 128	<table border="1" style="width: 100%;"> <tr> <td style="width: 50%;">Diclofenac Acq IV</td> <td style="width: 50%;">75 µg /ml</td> </tr> </table>	Diclofenac Acq IV	75 µg /ml	<table border="1" style="width: 100%;"> <tr> <td style="width: 50%;">Inj. Diclofenac sodium Inj IP</td> <td style="width: 50%;">75 mg/ml 1ml IV Bolus</td> </tr> </table>	Inj. Diclofenac sodium Inj IP	75 mg/ml 1ml IV Bolus
Diclofenac Acq IV	75 µg /ml						
Inj. Diclofenac sodium Inj IP	75 mg/ml 1ml IV Bolus						
5	Addendum for schedule-4 point no. 52 – 54 on Page no. 34	Double Lumen endotracheal tubes	Left sided Double Lumen endotracheal tubes: <ol style="list-style-type: none"> 1. Should contain 4 suction catheters, carlyn's type Y connector, Inbuilt Stylet, Separate clamp for tracheal and bronchial side. 2. Should be CE approved. 				
6	Page no. 35 schedule-4 point no. 62 – 63	Gum-elastic bougie (Sunmed/Rusch)	Gum-elastic bougie to assist endotracheal intubation				
7	Page no. 36 - 37 schedule- 4 point no. 99 – 104	Taper Cuff Sub-Glottic ETT Tube	Sub-Glottic ETT Tube				

Sr. No.	Page No/ Clause no/ point no	Existing sentence		To be read as																					
8	Page no. 37-38 schedule-4 point no. 109 – 111	Fenestrated Double Cannula tracheostomy Tube Cuff: The tube should have reusable fenestrated inner cannula with 15mm twist lock connector and cap.		Fenestrated Double Cannula tracheostomy Tube Cuff: The tube should have reusable																					
9	Page no. 39 schedule-5 point no. 1	Medicated Triple lumen central venous line 7 Fr		Medicated Triple lumen central venous line 7 Fr, length: 13 – 15 cm																					
10	Page no. 39 schedule-5 point no. 1	Medicated Triple lumen central venous line 5.5 Fr		Medicated Triple lumen central venous line 5.5 Fr, length: 6 – 8 cm																					
11	Page no. 40 schedule-7 point no. 1 – 2	<p>Combined spinal epidural sets</p> <p>Specification: Epidural Minipack</p> <ul style="list-style-type: none"> • Should have tuohy needle of 80mm length and 10mm graduations with internal stylet and optional snap on wings • Should have a 10ml Epidural ‘Loss of Resistance’ syringe • Should have an closed ended epidural catheter with 3 lateral eyes, a smooth tip, distal tip mark and 1cm marking to facilitate accurate catheter positioning • Should have a catheter connector with a flat profile for increased patient comfort • Should have a flat filter with a 0.2µm hydrophilic supported membrane that allows two way filtration • Should have CE approval • Size: 16 & 18 for Adult use 		<p>Combined spinal epidural sets sets – 16G & 18G</p> <ol style="list-style-type: none"> 1. Should have tuohy needle of 80mm length and 10mm graduations with internal stylet and optional snap on wings 2. Should have a 10ml Epidural ‘Loss of Resistance’ syringe 3. Should have an epidural catheter with a smooth tip, distal tip mark and 1cm marking to facilitate accurate catheter positioning 4. Should have a catheter connector with a flat profile for increased patient comfort 5. Should have a flat filter with a 0.2µm hydrophilic supported membrane that allows two way filtration 6. Should have CE approval 7. Size: 16 & 18 for Adult use 8. Should contain a pencil point spinal needle of 115mm length with internal stylet. 9. Should have graduations on locking sleeve that helps in indicating the distance spinal needle has advanced out of the Tuohy needle. 																					
12	Page no. 43 schedule-11 point no. 5	Disinfectant for endoscopes surgical instruments, etc	2.4% Gluteraldehyde (EPA USFDA)	Disinfectant for endoscopes surgical instruments, etc	2.45% Gluteraldehyde (EPA USFDA)																				
13	Page no. 43 schedule-12 point no. 2 to 5	<table border="1"> <tr> <td data-bbox="451 1666 518 1727">2</td> <td data-bbox="518 1666 662 2007" rowspan="4">Percutaneous Tracheostomy sets</td> <td data-bbox="662 1666 762 1727">5 mm</td> <td data-bbox="762 1666 997 2007" rowspan="4">Tracheotomy tube cuffed-CE/FDA/ISO mark, PVC high volume low pressure cuff, Radio opaque line, rounded tip.</td> </tr> <tr> <td data-bbox="451 1727 518 1787">3</td> <td data-bbox="662 1727 762 1787">5.5 mm</td> </tr> <tr> <td data-bbox="451 1787 518 1848">4</td> <td data-bbox="662 1787 762 1848">6.5 mm</td> </tr> <tr> <td data-bbox="451 1848 518 1908">5</td> <td data-bbox="662 1848 762 1908">9 mm</td> </tr> </table>	2	Percutaneous Tracheostomy sets	5 mm	Tracheotomy tube cuffed-CE/FDA/ISO mark, PVC high volume low pressure cuff, Radio opaque line, rounded tip.	3	5.5 mm	4	6.5 mm	5	9 mm		<table border="1"> <tr> <td data-bbox="1005 1666 1072 1727">2</td> <td data-bbox="1072 1666 1216 2007" rowspan="4">Percutaneous Tracheostomy sets</td> <td data-bbox="1216 1666 1316 1727">5 mm</td> <td data-bbox="1316 1666 1540 2007" rowspan="4">Single dilator percutaneous Tracheotomy tube cuffed-CE/FDA/ISO mark, PVC high volume low pressure cuff, Radio opaque line, rounded tip.</td> </tr> <tr> <td data-bbox="1005 1727 1072 1787">3</td> <td data-bbox="1216 1727 1316 1787">5.5 mm</td> </tr> <tr> <td data-bbox="1005 1787 1072 1848">4</td> <td data-bbox="1216 1787 1316 1848">6.5 mm</td> </tr> <tr> <td data-bbox="1005 1848 1072 1908">5</td> <td data-bbox="1216 1848 1316 1908">9 mm</td> </tr> </table>	2	Percutaneous Tracheostomy sets	5 mm	Single dilator percutaneous Tracheotomy tube cuffed-CE/FDA/ISO mark, PVC high volume low pressure cuff, Radio opaque line, rounded tip.	3	5.5 mm	4	6.5 mm	5	9 mm	
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14	Addendum point		For schedule no. 4, 5, 6, 7, 8 ,9 10 & 12 manufacturers distributors as well as importers can bid
15	Page no. 5 point no. 5	<p>The tenders are to be submitted by the manufacturers only. Tenders quoted by suppliers on behalf of manufacturers will not be entertained even if they are authorized by the manufacturers. However, manufacturers can give authority letter to the supplier/ distributor/ stockiest for the purpose of making supplies, raising bills, collecting payment etc. only after selection in the tender. In such cases, the manufacturer has to accept responsibility for any lapse on the part of the distributor/supplier and an undertaking to this effect from the manufacturer will have to be submitted. Failure to submit such an undertaking will lead to rejection of authorization and manufacturer will have to supply Drugs & Consumables Items directly. This authorization should be valid for the entire duration of the contract. No change in the authorized supplier/distributor will be allowed during the rate contract period. Sub authorization further to any other agent for delivery of the goods or for raising bills/collecting payment etc. will not be accepted.</p>	<p>For schedule 1, 2 and 3 the tenders are to be submitted by the manufacturers only, Tenders quoted by suppliers on behalf of manufacturers will not be entertained even if they are authorized by the manufacturers. However, manufacturers can give authority letter to the supplier/ distributor/ stockiest for the purpose of making supplies, raising bills, collecting payment etc. only after selection in the tender. In such cases, the manufacturer has to accept responsibility for any lapse on the part of the distributor/supplier and an undertaking to this effect from the manufacturer will have to be submitted. Failure to submit such an undertaking will lead to rejection of authorization and manufacturer will have to supply Drugs & Consumables Items directly. This authorization should be valid for the entire duration of the contract. No change in the authorized supplier/distributor will be allowed during the rate contract period. Sub authorization further to any other agent for delivery of the goods or for raising bills/collecting payment etc. will not be accepted.</p> <p>For schedule 4 to 12 manufacturers as well as their authorised importers can submit the tender.</p> <p>In case of third party manufacturer, the legal owners of the product / brand shall be responsible for the product quality. A declaration should be submitted to that effect by the bidders.</p>

**Store Officer,
AIIMS, Raipur**