Amendment no. 03 Dated 24.12.2018

HSCC/PUR/CNCI/Kolkata/Medical Equipment/04/B dt. 05.11.2018

Procurement of Medical Equipment CNCI 2nd Campus All bidders are requested to note the following:

| Name of Item | Last date & time sale/downloading of Tender document | Last closing/submission date & time for receipt of tender | Date of opening of Techno – Commercial bids. | Earlier date of opening | Amendment Status for opening date |
|--|---|--|--|----------------------------|---|
| Syringe Infusion Pump Volumetric Infusion Pump Pulse Oximeter Patient Warming System ICU ventilator Multipara monitor Paediatric Flexible Video Bronchoscope | 10.01.2019, up to 13.00 hrs IST | 10.01.2019, 14.00 hrs IST | 10.01.2019,14.30 hrs IST | 03.01.2018 | Date Extended |

| | Item No. 01 SYRINGE INFUSION PUMP Qty - 50 No.s | |
|-----|---|--------------------|
| Sr. | Tender Technical Specification | May please read as |
| | Operational Requirements | |
| 1. | The syringe pump should be programmable, user friendly, safe to use and should have comprehensive battery backup and comprehensive alarm system. | No change |
| | Technical Specification | |
| 1. | Flow rate for 50 ml syringe should be programmable from .1 to 1000ml/hr or more in steps of 0.1ml/hr with user selectable flow set rate option. SAVE last infusion rate even when the AC power is switched off. | No change |
| 2. | Should accept all makes of 5ml,10ml,20ml,50ml &60ml syringes with accuracy should be minimum of +/-2% or better | No change |
| 3. | Manual Bolus rate should be programmable in the range of 40-1000ml/hr or more with infused volume display and one key press bolus. Reminder audio after every 1 ml delivered. | No change |
| 4. | Display of drug name and total infused volume with a provision of memorising 50 names or more by the operator | No change |
| 5. | Key board locking system for patient safety. | No change |
| 6. | Should have bright display of drug name, flow rate, battery indicator, infused volume all at a time. | No change |
| 7. | Keep Vein Open(KVO) must be available 1.0ml/hr or set rate if lower than 1.0ml. User should have a choice to disable KVO when required | No change |
| 8. | Selectable Occlusion pressure trigger levels selectable from 300/500/700 mmHg or at least three selectable levels. | No change |
| 9. | Manual pusher with plunger protection guard. | No change |
| 10. | Automatic detection of syringe size and proper fixing .Must provide alarm for wrong loading of | No change |

| | syringe such as flanges out of slot, disengaged plunger, unsecured barrel etc | |
|-----|---|---|
| 11. | Anti bolus system to reduce pressure on sudden release of obstruction. | No change |
| 12. | Should have comprehensive alarm package including: Occlusion limit exceed alarm, near end of infusion pre-alarm &alarm, volume limit pre-alarm & alarm, KVO rate flow, Low battery pre-alarm & alarm, Occlusion pressure pre alarm& alarm, AC power failure, Drive disengaged & preventive maintenance. | Should have comprehensive alarm package including: Occlusion limit exceed alarm, near end of infusion pre- alarm & alarm, volume limit pre-alarm & alarm, KVO rate flow, Low battery pre-alarm & alarm, Occlusion pressure pre alarm& alarm, AC power failure, Drive disengaged & maintenance alarm. |
| 13. | Should have digital & analog display of occlusion pressure indicator | Should have digital & visual display of occlusion pressure indicator |
| 14. | Should display remaining battery life in hrs & minutes on operating flow rate. | Should display remaining battery life in hrs & minutes (preferable) on operating flow rate. |
| 15. | Should have Universal mounting accessory on both vertical & horizontal stand. | No change |
| 16. | Should have facility of auto dose calculation. | No change |
| | System Configuration Accessories, Spares & Consumables for each pump | |
| 1. | Syringe infusion pump-01 | No change |
| 2. | Mounting device/Docking Station for at least four pumps as per requirement so as to enable to power up to up to 4 pumps with one power cord when mounted on IV pole(price to be quoted separately). | No change |
| | Environmental factors | |
| | The unit shall be capable of operating continuously in ambient temp of 10-40 deg C and relive humidity of 15-90% | No change |
| | Power Supply | |
| 1. | Power input to be 220-240VAC, 50Hz. Power cable should be fitted with Indian plug & adapter. | No change |
| 2. | Should have a rechargeable battery for 2 hr or more back up for about 5ml/hr flow rate with 50 ml syringes. Larger battery life and indication of residual life will be preferred. | No change |
| 3. | Resettable over current breaker shall be fitted for protection | Deleted. |
| | Standards, Safety and Training | |
| 1. | The product should be FDA approved/European CE certified with four digit number. | No change |
| 2. | Drop Test-Withstands 1 meter drop to any edge, corner or surface. | No change |
| 3. | Should conform to international test protocols on exposure to shock forces and to vibration forces. The standard should be documented. | No change |
| | Documentation | |
| | User & service manual in English. | |
| | List of important spare parts & accessories included in the warranty with their part number & costing | |
| | Certificate of calibration & inspection from factory. | |
| | Log book with instruction for daily, weekly, monthly &quarterly maintenance checklist. | |
| | List of equipments available for providing calibration & routine maintenance support as per manufacturer documentation in service/technical manual. | |
| | Must submit user list & performance report within last 5 yrs from major hospitals of West Bengal. | |

| Service station should be in Kolkata. | |
|---|---|
| Compliance report to be submitted in a tabulated & point wise manner clearly mentioning the para/page no. of original catalogue/datasheet. Any point, if not substantiated with authenticated catalogue/manual, will not be considered. | |
| To be added | Technical evaluation will be finalised only after functional demonstration of the offer equipment. The functional demonstration will not be a substitute for technical evaluation of the document submitted along with the bid. |

| | Item No. 03 Pulse Oximeter with Heart Rate | |
|-----|--|--|
| Sr. | Tender Technical Specification | May please read as |
| 1. | Functional Requirement | |
| | 1. Continuously capable of displaying oxygen saturation in real time using an external probe on skin. | No change |
| | 2. Should have reusable, sterilisable (using medical grade disinfectant) robust probe which can be | No change |
| | easily connected & disconnected. | |
| | 3. Should operate from main voltage or from internal rechargeable battery. | No change |
| 2. | Technical Characteristics | |
| | a)SPO2 measurement range at least 20-70 and 70-99%, minimum gradation 1%. | No change |
| | b)Accuracy of SPO2 better than +_1% for range 20-70 and better than +_3% for range 70-99. | b)Accuracy of SPO2 better than +/-2% for range 40-70 and |
| | | better than +/-3% for range 70-99. |
| | c)Pulse rate range at least 30 to 240 bpm, minimum gradation 1 bpm. | No change |
| | d) Accuracy of pulse rate better than +_5bpm. | No change |
| | e) Signal strength or quality to be visually displayed. | No change |
| | f)Audio-visual alarms required high & low SPO2 and pulse rate (operator variable settings), sensor | No change |
| | disconnected, sensor failure, low battery. | |
| | g) TFT screen size 5" or more. | No change |
| | h) Plethysmograph display is mandatory, | No change |
| | i) Should have minimum 24 hrs trend memory for SPO2 & PR. | No change |
| | j) Should have easily accessible touch button & dial/touch screen to operate the machine. | No change |
| | k) Weight should be less than 5 kg. Case is to be hard & splash proof. Display must allow easy viewing | No change |
| | in all ambient light levels. | |
| | I) Noise should be <50Dba. | No change |
| | m)Voltage(value, ac or Dc, monophase or triphase) 220 to 240 V,50Hz | No change |
| | n) Internal replaceable rechargeable battery allows operation for at least two hrs in event of power | No change |
| | failure. Battery charger to be internal to the system and integral to mains power supply & to charge | |
| | battery during mains power operation unit. | |
| | o) Power consumption to be 50-100W. Mains supply cable to be at least 3m in length. | No change |
| 3. | Accessories, Spare parts & Consumables | |
| | Accessories (mandatory, standard, optional): | No change |

| | i) Reusable probes each for adult, pediatric & infant use, one number each | No change |
|----|---|-----------|
| | ii) Y probes with clips for infant use & forehead spo2 sensor for detection of low saturation level(<70%)/ lex probe with the provision of fixation, one number each for reusable type | No change |
| 4. | Spare parts : | |
| | Two sets of spare fuses (if non re-settable fuses used) | No change |
| 5. | Environmental consideration: | |
| | 5.1. Capable of operating continuously in ambient temperature of 0-50° C & relative humidity of 15- 90 % in ideal circumstances | No change |
| | 5.2. User's care, cleaning, disinfection & sterility: Cleanable with alcohol or chlorine wipes | No change |
| 6. | Standards & safety | |
| | 6.1 Should be USFDA Approved/European CE Certified with 4 digit certification | No change |
| | 6.2 Advanced maintenance task required shall be documented | No change |
| | 6.3 Warranty of 5 years with free servicing (min 5 during warranty) | No change |
| | 6.4 The spare price list of all spares & accessories (including minor) required for maintenance & repair in future after warranty period should be attached | No change |
| 7. | The Operating manual, Service manual, other manuals | |
| | 7.1 The user & maintenance manuals to be supplied in English | No change |
| | 7.2 Certificate of calibration & inspection to be provided Service Centre should be in Kolkata | No change |
| | 7.3 Any recommendation for use & supplementary warning for safety should be declared. | No change |
| 8. | Technical evaluation will be finalised only after functional demonstration of the offer equipment. The functional demonstration will not be a substitute for technical evaluation of the document submitted | No change |
| 5. | along with the bid. | |

| | Item No. 06 – Multi-para Monitor, Qty - 50 Nos | |
|-----|---|---|
| SN | Tender Specifications | May please read as |
| 1. | Advanced high end modular patient monitor of screen size 12 inches or more, having integrated non-invasive and invasive measurements and features suitable for paediatric and adult patients. | Advanced high end modular patient monitor of Touch screen and Trim Knot facility (preferable) with screen size 15" or more, having integrated non- invasive and invasive measurements and features suitable for paediatric and adult patients. |
| 2. | Monitor must have the facility to display minimum 6 waveforms or more, along with related numerical parameters on single screen. | Monitor must have the facility to display minimum 8 waveforms or more along with related numerical parameters on single screen. |
| 3. | Parameters: Monitor must have the facility to monitor ECG, low perfusion motion tolerant SpO2 (with waveform), HR (source: ECG/SpO2 pleth/Auto), NIBP, RR, dual temperature, IBP, CVP. | No Change |
| 4. | Monitor must have facility to display real-time 12 lead ECG with ST-T changes. | No Change |
| 5. | Monitor must have advanced arrhythmia detection and ST analysis as standard feature. | No Change |
| 6. | Monitor must have the time linked review function for both graphic & tabular trends of all parameters for 48 hours. Monitor must have arrhythmia recall facility. | No Change |
| 7. | Monitor must have inbuilt rechargeable battery for minimum one hour operation for both machine & display board. | No Change |
| 8. | Monitors must have printing facility through central stations. | No Change |
| 9. | Monitor must be able to connect to central monitoring station. | No Change |
| 10. | Should be USFDA Approved/European CE Certified with 4 digit certification. | No Change as per office memorandum No. X.11035/379/2015-DFQC(Pt) of Ministry of Health & Family Welfare Dated 20 th February 2018 |
| 11. | Each monitor to be supplied with. | Each monitor to be supplied with. |
| | a) Five lead ECG electrode cable – Two in number with adult & paediatric ECG electrodes. | a) Five lead ECG electrode cable – Two in number with adult & paediatric ECG electrodes. |
| | b) SpO2 soft probe: | b) SpO2 soft probe: |
| | · Adult Probe: 5 No.s. | · Adult Probe: 5 No.s. |
| | · Pediatric Probe: 3 No.s. | · Pediatric Probe: 3 No.s. |
| | · Neonatal Probe: 3 No.s (Wrap type). | · Neonatal Probe: 3 No.s (Wrap type). |
| | c) NIBP cuffs for adult & paediatric, large adult, thigh: Five each size. | c) NIBP cuffs for adult & paediatric, large adult, thigh: Five each size. |
| | d) Temperature probe: One skin and two core probes for each monitor. | d) Temperature probe: One skin and two core probes for each monitor. |
| | | e) IBP connection cable: Four in number |
| | | f) IBP Disposable pressure transducers: 20(Twenty) in number each monitor |
| 12. | Each monitor must be supplied with. | All 50 monitors must be supplied with |
| | | 25 interchangeable Side-stream CO2 Module with 50 etCO2 sampling line each module 20 water trap each module Or |
| | | 25 interchangeable Main-stream CO2 Module with 2 Mainstream sensors each module 5 adult reusable airway adapter |

| | | 2 infant reusable airway adapter |
|-----|---|--|
| | a) IBP connection cable: Four in number. | Deleted |
| | b) IBP Disposable pressure transducers: 20(Twenty) in number each monitor. | Deleted |
| | c) 50 etCO2 sampling line with 20 water trap each for 25 modules in side-stream type/ 2 mainstream sensors with 5 adult airway adapter & 3 infant airway adapter each for 18 modules | Amended as above |
| | | All 50 monitors must be supplied with |
| | | 5 interchangeable modules for third IBP |
| 13. | 6(Six) Central Nursing Station for every 8-10 beds must be provided for complete supply of total 50 monitors. | No Change |
| | a) Cabling, switches, hubs for central monitoring network has to be done by bidder in the ITU. | No Change |
| | b) Each Central Station should be supplied with 21" color LED medical grade display and one 21" slave display and one laserjet printer. | Each Central Station should be supplied with 21" or more color LED medical grade display and one 21" slave display and one laserjet printer. |
| | c) One Online UPS of 1 hour backup should be supplied with each Central Nursing Station. | No Change |
| 14 | 25 Interchangeable modules of etCO2 (Mainstream/sidestream) must be provided for complete supply of total 50 monitors & 4 modules for third IBP must be provided for complete supply of total 50 monitors. | Deleted |
| 15 | User and service manual must be in English. | No Change |
| 16 | Service centre should be in Kolkata. | No Change |
| 17. | Wall mounting bracket with flexible arm to be provided by company. | No Change |
| 18. | Monitor should be 12 inches display with touch screen and Trim knob. | Monitor should be 15 inches display with touch screen and Trim knob (preferable). |
| 19. | Monitors should have capable of future up-gradation through interchangeable modules to monitor and display. 4 channel EEG Cardiac output (both thermos-dilution, as well as minimally invasive) Spirometry | Monitors should have capable of future up-gradation through interchangeable modules to monitor and display. 4 channel EEG Cardiac output (both thermos-dilution, as well as minimally invasive) Spirometry SvO2 ADD ON: Monitor should preferably have a facility for ICU electronic charting and integration with other ICU equipments like ventilator & syringe pumps etc. Integration to be done considering one ventilator, one patient monitor, one rack of syringe pump at bedside. |
| 20. | Monitor should have non-volatile memory; the trend data should remain saved during power-off in the monitor/central station. | No Change |
| 21. | The monitors or central station should have HL7 output for future digital upgradation. | No Change |
| 22. | The monitor should be able to measure and display ECG, Respiration, NIBP, SpO2, Dual IBP, Dual Temperature & EtCO2 simultaneously. | No Change |
| 23. | The monitor and central station should be upgradable in future to HIS, PACS/ DICOM Viewer & Web browsing facility through mobiles/laptop. | No Change |
| 24. | The monitor should have the capability to measure & display QT/QTc. | No Change |
| 25. | Technical evaluation will be finalized only after functional demonstration of the offer equipment. The functional demonstration will not be a substitute for technical evaluation of the document submitted along with the bid. | No Change |

| To be added | The manufacturer should preferably have the capability with proven track |
|-------------|--|
| | record to upgrade the monitoring system to integrated paperless e-charting |
| | to connect all ICU devices at one point. |
| | |

All other terms and conditions of the tender enquiry document shall remain unchanged. Prospective bidders are advised to regularly visit HSCC website/ CPP as corrigendum /amendments etc. if any, will be notified on this portal only, no separate advertisement will published in the news papers.

Sr. CGM-I, HSCC (I) Ltd For & on behalf of Director CNCI, Kolkata