Enabel

Tender Specifications

Public contract for the supply, delivery, installation and testing of medical equipment for health facilities as well as conducting user training.

Open Procedure

Reference number: UGA22009-10015

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DEROGATIONS FROM THE GENERAL IMPLEMENTING RULES

Section 4, 'Specific contractual and administrative conditions' of these Tender Specifications (CSC/Cahier Spécial des Charges) holds the specific administrative and contractual provisions that apply to this public contract by way of derogation from the Royal Decree of 14 January 2013 or as a complement or an elaboration thereof.

These tender documents derogate from Art. 25-33 of the General Implementing Rules (see point 4.7 "Performance bond (Art. 25-33)"). This is motivated by the need to provide equal opportunity for local and international tenderers to participate with a view to increasing competition.

1 Technical Specifications

1.1 Requirements for the goods

Technical requirements

The detailed technical specification of the items and quantities are listed below;

Lot 1: Supply and delivery of Cemonc equipment

Device Name	Oxygen Concentrator 10LPM	Quantity
General description	An oxygen concentrator is a device used to produce an enriched oxygen gas from atmospheric air; for use by people requiring medical oxygen due to low oxygen levels in their blood.	8 pcs
Composition (Per set):	Main unit: 1No. Standard accessories: 1 set	
Features/Performance Specifications	**Main unit** Oxygen concentrator capable of producing medical grade oxygen from atmospheric air. The unit should be mobile on castors and capable of supplying oxygen to two patients at a time. It should incorporate oxygen monitor facility complete with patient tubings. **Detailed Specifications** • Oxygen production technology: Pressure Swing Adsorption (PSA) • Oxygen outlets: Dual flow complete with flow meter • Oxygen flow rate: 2 – 10 LPM • Oxygen concentration: 93%±3% at maximum flow range. • Oxygen purity: Medical grade, oil free oxygen, 93%±3% oxygen concentration at maximum flow rate. • Alarm system alert: Audible/Visible light alarm on low oxygen concentration and power failure.	

	• Safety: Shutdown with power failure, high or low pressure and high temperature. • Dimensions (mm): Approx : Min 600(H) x 300(W) x 340(D) • Power supply: 100 - 240VAC, 50Hz with an inbuilt suitable surge Protector or Stabiliser suitable for running oxygen concentrator. **Standard Accessories** • Power supply cable: 1No. x 3m length with 3-pin top plug (British Standard) • Nosal cannulas, paediatric	
	Humidifier bottles, 150ml x 2No.	
	Voltage Stabilizer with surge protection: 240VAC/50Hz output.	
Recommended Spare parts	Air intake/gross particle bacterial filter: 1No. x Pack of 5Pcs	
	Fine/final bacterial filter: 1No. x Pack of 5Pcs	
	Fuses: Set	
Recommended Consumables	Oxygen tubings – Adult and Neonatal sizes	
Standards and Certifications	CE Mark, ISO9001, ISO13485	
Device Name	**Vacuum Extractor**	**Quantity**
General description	The device used to assist in the delivery of babies during complicated labour (e.g. delayed second stage).	14 pcs
Composition (Per set):	• Main unit • Standard accessories	
Features/Performance Specifications	**Main unit** • Type: Hand-operated vacuum pump • Bird type, stainless steel • Material: All parts should be autoclavable at 121^0C	

	• Soft vacuum extractor cup size: Ø50mm, 60mm & 940mm • Extraction handle: 2No. stainless steel • Cover plug for collection bottle with 3 holes • Connector, angled, chrome plated • Connector, angled with screw valve, chrome plated screw, cap and gasket. • Vacuum gauge: 0 to -100 kPa/- 800mmHg) with straight connector, chrome plated • Vacuum bottle: 500ml **Standard Accessories** • Suction tube: 2No. x silicone, transparent, inner/outer diameter 6/11mm, length 50 and 150cm • Carrying case: 1No.	
Recommended Spare parts	• Vacuum bottle: 500ml x 2No. • Spare gaskets for the screw valve: 1 set	
Recommended Consumables	• Suction tube: 2No	
Standards and Certifications	CE Mark, MDD93/42/EEC	
Device Name	**Dilation and Curettage (D&C) Set**	**Quantity**
General description	A set of instruments is used to carry out the D&C procedure.	14 Sets
Composition (Per set):	Set of instruments	
Features/Performance Specifications	Material: Stainless steel, hospital grade 304, reusable. **Composition of set:** • Dilators, Uterine, Hegar, Double Ended, Set of 8 — 1No. • Catheter, Metal, Female, s/s, Ch.12 — 1No. • Towel Clip, Cross Action — 4No. • Curette, Uterine, Sims, Blunt, D.E. — 1No.	

	• Forceps, Dissecting, Straight, 1/2 Teeth, Serrated	1No.	
	• Forceps, Dissecting, Straight, Plain, Serrated	1No.	
	• Forceps, Uterine, Dressing, Bozeman	1No.	
	• Forceps, Ovum, McClintock	1No.	
	• Forceps, Sponge Holding, 240mm	3No	
	• Forceps, Uterine, Vulsellum, Teale	2No	
	• Sound, Uterine, Malleable	1No.	
	• Speculum, Vaginal, Auvard	1No.	
	• Speculum, Vaginal, Sim's	1No.	
	• Instrument Container, s/s, With Cover	1No.	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485		
Device Name	**Portable Ultrasound Scanner**		**Quantity**
General description	The device used to examine internal body organs (e.g. liver, gall, kidney, pancreas, thyroid, breast, uterus, bladder, ovary) using ultrasound technology.		1 Pc
Composition (Per set):	**Standard Configuration** • Main unit with digital scan converter: 1No. • 10" high-resolution monitor 1No. • 2.5MHz ~ 5.0MHz convex probe: 1No. • 5.0MHz ~ 10.0MHz linear probe: 1No. • 5.0MHz ~ 12.0MHz trans-vaginal probe: 1No. • Thermal Printer: 1No. • Communication and RS-232C interface • Trolley: 1No. • UPS unit: 1No. • Ultrasonic gel: 5 Lt.		
Features/Performance Specifications	Features: • High-quality image processor • Cineloop • Probe frequency conversation option		

	• Computer image communication • Various measuring function • Backlit keyboard • 10” high resolution monitor • Dual probe connector		
	Performance Specifications		
	1	Appearance	Mobile ultrasound system, 10-inch high-resolution monitor
	2	Scanning Mode	Linear, Convex and trans-vaginal scanning
	3	Display Mode	B, B/M, M, B+B
	4	Scanning Angel	Max. 90°
	5	Probe Frequency	2.5MHz ~ 5.0MHz (convex probe), 5.0MHz ~ 10.0MHz (linear probe), 5.0MHz ~ 12.0MHz (trans-vaginal probe), broad band, tri-frequency.
	6	Focusing Method	Multi-step focusing with variable aperture
	7	Zooming	Multi-zoom rate and depth shift
	8	Image Processing	Pre-processing, Correlation-processing, interpolation, Y correction, image reverse of right/left, up/down and positive/negative
	9	Grayscale	256
	10	Measurement & calculation functions	Distance, area, circumference, volume, angle, time, speed, heart rate, heart, heart functions, OB/GYN
	11	Image annotation	Display with many body marks and probe marks, image annotation
	12	Character display	ID, date, time, hospital name, probe type, frequency, gain, zoom,

		focus, and name of prescriber, as a minimum.	
	13 Body Mark	Body marks with probe mark display	
	14 Cine loop	64 frames for B mode, 256 seconds for M mode	
	15 Trolley	Mobile on castors, diameter 4" minimum	
	16 UPS	240VAC output with battery backup capacity of at least 30 minutes.	
	17 Communication	Image data can be transferred to the PC through interface RS-232C to save and process in JPG, BMP or TIF format	
	18 Power supply	AC 100 - 240V/ 50Hz	
	19 Printer	Thermal printer type supplied Sono paper roll	
	20 Gel	Ultrasound gel: 5Lt bottle	
Recommended Consumables	• Ultrasound gel: 5Lt bottle x 2No. • Sono paper roll: 5No.		
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO60601-1		
Device Name	**Universal Anaesthesia Machine (UAM)**		**Quantity**
General description	A device used to administer anaesthetic gases during a surgical procedure on a patient.		6 Units
Composition (Per set):	• Main unit • Standard accessories		
Features/Performance Specifications	**Main unit** Anaesthesia machine with the capacity to generate oxygen using an inbuilt oxygen generator or external oxygen source (e.g. oxygen cylinder).		

	• Construction: Aluminium frame, Wide worktop and drawers • Dimensions (LXWXH): 650-800 x 50-60 x 120-135 cm • Monitor: In-built patient monitor with touch screen control and information display • In-built anaesthesia ventilator with touch screen control and information display. Automatic and manual, drive type should be a turbine, battery backup time 4-6 hours, display be 10-11 inches touch screen, mode [Volume, pressure and spont]. Tidal volume of 20- 2000Ml, minute volume 0.3-15L/min, Pressure control 0-60CmH_20, PEEP 0-25 CmH_20; I: E RATION 1.9-5.2, Rare1-100bpm. • Worktop shall be chemical resistant and rustproof • Vaporizers: Isoflurane and sevoflurane, should work with and without compressed gas, Have low resistance; Capacity of 110-125mL • Drawers: 2 lockable drawers • Castor Wheels: 4 castor wheels, Ø4" minimum; antistatic non-conductive material, with 2 lockable. • Oxygen sensor: Paramagnetic oxygen sensor • Oxygen flow rate: 0.1-10 LPM • Should be able to work with an auxiliary oxygen supply • Additional gas: Air • Cylinder pin index yokes: O2, Air • O2 flush: (at 50 psi): max. 35 - 50 L/min • Inbuilt Anaesthetist lamp. • Total fresh gas flow meter: 0 to 10 L/min, • Power Supply: 100 - 240 VAC, 50/60 Hz, with integrated AVS protection	

	• Alarm; High and low setting values, breathing circuit disconnect. **Standard Accessories** • Compatible Oxygen cylinder with cylinder head with wheel-type knob valve. • Power supply cables. • Cylinder trolley with a restraining chain. • Patient breathing circuits and tubings • Soda lime: 5Kg pack	
Recommended Spare parts	• Filter: 10Pc pack	
Recommended Consumables	• Patient breathing circuits [Adult and Peadrartic] and tubings – 1 set	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Autoclave, Electric, 75 Lt (sterilizer)**	**Quantity**
General description	The device is used to sterilize equipment and medical supplies to kill microorganisms using steam under pressure at 121/132/134°C.	5 Pc
Composition (Per set):	Main unit Standard Accessories	
Features/Performance Specifications	**Main unit** • Control: Microprocessor controlled • External finish: Powder-coated steel body • Type: Vertical with casters for excellent mobility • Sterilizing temperature: Max. 136 deg. Celsius • Material (Internal chamber): Stainless steel grade 304 • Loading capacity: 75 litres, Sterilization Chamber. • Heat generation: Electric heater elements • Gasket type: Airtight Silicon gasket • Thermal printer, with record paper roll	

	• Internal dimensions (mm): Approx. Diam. 490 x 540(H) • Display: Digital backlight LCD • Timer: 0-60min • Working Pressure. 0.22 to 0.23 MPa • Programs: Pre-programmed sterilization cycles • Power supply: 100 - 220 VAC/50Hz **Accessories** • Supplied with two wire gaskets, • Sterilizing drums: Small (6in x 6in), Medium (11in x 9in), Large (12in x 15in); all in stainless steel grade 304.	
Recommended Spare parts	• Heating element: 1 set • Gasket: 1 Pack • Water sensor: 2No. • Safety Valve: 1 set	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Operation Table, Hydraulic /theatre bed**	**Quantity**
General item description	A device for performing surgical operations on patients.	5 Pc
Composition (Per set):	• Main unit • Standard Accessories	
Features/ Performance Specifications	**Main unit** • Universal operation table, mobile with simple mechanical and hydraulic system. • Material: Stainless steel, hospital grade 304. • Table top: X-ray translucent and with head, back, pelvis and leg plate. • Guide rails: Equipped. • Height adjustment (mm): 650 to 1,040	

	• Table height adjustment: Foot pedal
	• Lateral tilt: By hand drive.
	• Table Dimension Minimum : 2000(L) x 500(w) x 1,040(H)
	• Bed section Movements
	  • Leg Sectors 22, -90
	  • Back Section +75, -45
	  • Head Section +50, -90
	  • Tilting to both 20
	  • Trendelenburg 45
	  • Anti-Trendelenburg 20
	Accessories
	• Arm support
	• Leg support
	• X-ray cassette tray
	• Anaesthesia brackets
	• Body Belt	

Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Operating Light, Mobile (LED)**	**Quantity**
General item description	Used to focus light to illuminate a patient's specific body area or cavity during a surgical procedure.	8 Pc
Composition (Per set):	Main unit	
Features/ Performance Specifications	• Single light head with LED lights in cluster. • Adjustable field of view including at least 0.17 - 0.30m range (depending on the model supplied). • Focal length of at least 0.7 m. • Maximum illumination of at least 120,000 lux at 1 m. • Colour temperature between 3000°K and 5000°K. • Colour rendering index (CRI) of 95.	

	• Sterilizable and removable light handle for positioning. • Minimum LED life of 50,000 hours. • Minimum of 100° horizontal turning. • Articulating arm between light and base stand. • Base with at least four (4) anti-static swivel castors in a star formation. • A minimum of two (2) castors are equipped with brakes. • Heavy base with low centre of gravity, indicate weight of base relative to total device weight. • Floor to light height adjustable to include at least range of 1.1 m to 2.12 m. • On/off switch, battery status indicator (depending on model) and controls to adjust light intensity. • Dimming function with a minimum of 5 settings or fully adjustable. • Built in backup rechargeable battery with minimum 3 hours of battery life, battery type lead-acid. • Automatic switch from mains to batteries in case of power failure. • Power requirements: 100 - 240 Volts - 50/60 Hz.	
Recommended Spare parts	LED Lamp (set).	
Standards and Certifications	CE Mark, ISI, ISO 9001:2008	
Device Name	**Delivery Bed**	**Quantity**
General item description	A medical device for conducting deliveries. General item description Fixed height bed for conducting deliveries.	8 Pc
Composition (Per set):	NA	

Features/ Performance Specifications	Purpose of use Features/Performance Specifications • Construction: 2/3 Sections with adjustable backrest and waste receptacle can. • Constructed from round/rectangular mild steel sections, 2.0mm thick/gauge steel plate section • Under the body section, the leg section is partially extendable to serve as a shelf for receiving infant or basin/ • Leg section can be completely recessed 25mm. • Equipped with snaps and straps to hold pads to table. • Dimensions: 1900Lx650Wx800H (mm) • Well finished and epoxy paint coated • Body section stands mounted on PVC stumps/shoes. • Leg section stands mounted on 2" castor wheels, 2 with brakes. • Supplied with removable mattress 75mm thick, with vinyl leather cover with heavy-duty zipper. • Leg holder: Equipped, mounted on bed frame.	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Wheel Chair, Adult**	**Quantity**
General item description	For moving patients who are unable to walk on their own or persons with disability. General item description Foldable wheelchair with footrest.	1 Pc
Composition (Per set):	Composition (Per set): • Main unit • IV Pole	

Features/ Performance Specifications	Features and specifications: Main Unit • Type: Foldable wheelchair for adults • Front wheels: Free rolling, 360 degrees' swivel. • Rear Wheels: Equipped with hand-operated brakes. • Two push handles at the rear. • Frame: chrome-plated or epoxy-coated tubular steel. • IV Pole: Equipped. • Frame, diameter: 2.2cm. • Frame: 2.2cm (outside, across), 1.2mm thickness • Side-to-side leg support: Yes • Swing-away foot and arm supports: Equipped • Upholstery for Armrests, seat and back: Washable plastic, flexible, tear resistant, anti-static, flame retardant, disinfectant and liquid proof. • tyres: heavy-duty solid rubber • Carrying capacity: 150 Kg. **Dimensions** • Overall Dimension (D x W x H).: 45 x 60 x 89 cm • Seat Depth: 42-43cm • Back Rest: 43.5-49 x4 0.5-42 cm (W X H). • Side-to-side legs support: 47-50 x 8.5-23 cm. • Front Wheel diameter: 18-22cm, • Rear Wheel diameter: 58-60cm.	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Quantity	1 Pc	
Device Name	**Patient Stretcher/Trolley**	**Quantity**
General item description	Purpose of use • Designed specifically for patient transport, 2 sections	24 Pc

	General item description • Patient trolley with side rails and stretcher	
Composition (Per set):	One Main unit	
Features/ Performance Specifications	**Features/Performance Specifications** • Heavy duty carriage mounted on 4 swivel castors, two with brake and four with anti-static wheel. • Backrest angle adjustable via secured pawl and gear ratchet, safe for patient and operator • Transfer bars connect all lower distal portions of the 4 castors, providing maximal structural strength • With fold-away side rails • Head-end side has a removable height adjustable IV-pole; height is set with a robust clamp with a heavy knob • Fixings of the fold-away side rails and IV-pole are solid steel and welded to the frame of the stretcher **Material:** • Frame: epoxy-coated tubular steel • Cover: plastic, flexible highly tear resistant, anti-static, flame retardant, disinfectant- and liquid proof, washable • Caster frame/bracket: steel or nylon **Dimensions:** • Stretcher, two sections extended, including upholstery: 160-200x50-60x72-88cm (l x w x h) • Fold away side rails: 50-100 x 30-45cm (l x h) • Frame: 2.7-3.3cm (outside, across), 1.8-2.2mm (thickness) • Swivel castor wheel: 2.3-3 x11-15cm (w x diameter) • Upholstery: 4.5-55cm (h) • Carrying capacity: minimum 160 kg	

Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Patient Monitor 5 parameters with a stand**	**Quantity**
General item description	Portable 5-parameter patient monitor. Used to measure and monitor patients' basic physiologic parameters (“vitals”) and display them on screen.	8 Pc
Composition (Per set):	Composition (Per set): • Main unit • Standard accessories	
Features/ Performance Specifications	Purpose of use Features/Performance Specifications **Main unit** • The monitor should be equipped with appropriate software. • Display: At least 15 inches’ touch screen TFT display with a minimum of 6 waveforms • Measured parameters: SpO2, ECG, NIBP, Temperature, Respiratory rate, Heart rate. • Resolution of screen: 1200x700, minimum. • NIBP: (Manual/Auto/STAT); • Measurement type: (adult, pediatric, neonatal); • Measurement range: Systolic, Diastolic, Mean; Mean Values - (2.6 - 35.0 kPa); Resolution - (0.1 kPa); Accuracy - (± 0.4 kPa or 5 %); • SPO2 : Measurement range: 0 - 100 % • Pulse measurement range: 30- 250 bpm; • BPM Accuracy: ± 2 %; • Temperature: surface and rectal; Measurement range -Minus 1 to 45C; Accuracy: ± 0.1 • Alarms: Alarm sound (Crisis/warning/Advisory);	

	• Alarm silence; Alarm suspend; Alarm Volume adjustable; Vital sign alarm; Arrhythmia; Technical alarms. • Up to five hours battery backup. • Trend recorder storage of at-least 48 hours. • Power Supply: 100-240VAC/50Hz.	
Recommended Spare parts	Standard Accessories • Trolley on castor wheels with brakes. • ECG/Resp: 5 lead ECG cable with Clip – 2 set per monitor • NIBP; Adult cuff – 2No • NIBP Pediatric Cuffs – I set • SpO2: Adult SpO2 sensors – 2 No. • Temperature Probe: Central temperature Probe- 2 No. • Skin temperature probe – 2 No. • Recommended Consumables •NIBP; Adult cuff – 1 set • NIBP Pediatric Cuffs – 1 set • SpO2: Adult SpO2 sensors – 2 No. • Temperature Probe: Central temperature Probe- 2 No. • Skin temperature probe – 2 No. • ECG electrodes – 50Pc pack	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Foetal Doppler**	**Quantity**
General item description	Used to detect the foetal heartbeat of a foetus or unborn baby in a mother's womb. **General item description**	20 Pc

	A handheld Doppler foetal monitor used to provide an audible simulation of the heartbeat of a foetus or unborn baby in a mother's womb	
Composition (Per set):	Composition (Per set) • Main unit • Standard Accessories	
Features/ Performance Specifications	Features/Performance Specifications • Type: Handheld • Probe Frequency: 2 - 2.5 MHz • Output Power: 28 mW/cm2 or less. • Fatal heart rate (FHR) range: 50 ～ 220 bpm or wider • Display: LCD, energy indicator, FHR, Battery status. • Speaker: Inbuilt speaker; 0.6W or more • Power source: Rechargeable Battery. • Battery charging: Automatic charging of battery while on AC line power. • Audio mute switch to prevent noise • Energy conservation: Automatic power off when not in use. • Power cord (Set)	
Recommended Spare parts	Battery set	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Patient Bed**	**Quantity**
General item description	**Purpose of use** To provide safety, comfort and mobility for hospitalised patients in need of health care.	8 Pc
Composition (Per set):	Main unit: 1	

Features/ Performance Specifications	**Main Feature** • Backrest, Footrest and Trendelenburg movements by 3 cranks/3 movements manual operation • Epoxy-coated mild steel bed frame • 3 section bed board • Detachable head/footboard with easy lock • ABS side rails with angle indicator • ABS head and footboard with holes for moving the bed easily • Diameter of 125mm noiseless castors, all with individual brakes • Bumper castors on 4 corners for protection • Should have Collapsible side rails **Accessories** • IV pole with adjustable height • Drainage hooks at the side of the bed **Specification** • Overall size（L*W*H）2160*960*500mm • Back section adjustable 0-75° • Foot section adjustable 0-40° • Trendelenburg 0-12° • Caster Dia with break 125mm (5") • Safe working load 140-240 kg	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Blood Refrigerator**	**Quantity**
General item description	Purpose of use Used to store and preserve blood and blood components at temperatures ranging from 2°C to 6°C. Cabinet-type Blood Bank refrigerator for safe storage of blood in a hospital.	1 pc
Composition (Per set):	Composition (Per set): • Main unit	

	• Standard accessories (Set)	
Features/ Performance Specifications	**Features/Performance Specifications** Type: Cabinet type, vertical. Cooling Method: Forced Air Cooling. Defrosting: Automatic. Refrigerant: CFC-Free. Number of Shelves: 2 or more. Operating Temperature: 4ºC to 6ºC. Capacity: 450ml Blood bags x 60No. • Internal Light: Yes • Temperature Display: Digital. • Alarms: High and low-temperature alarms, Sensor error alarms, Power failure alarms, Door ajar alarms, low battery alarms. Power Consumption: Max: 240W. • Power Supply : 100-240VAC/50Hz Gross Weight: 55Kg Internal Size (W*D*H)mm: 505*560*610 External Size (W*D*H)mm: 625*820*1150 • Accessories • Testing hole • Outdoor latch light. • Thermal printer/Temperature Recorder Supply with Supplied with Automatic voltage regulator suitable for the size above	
Recommended Spare parts	Digital Thermometer LCD Display, Light tubes, Fuse (set).	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Hemocue HB**	**Quantity**

General description	Hand-held, glucose photometer HemoCue Glucose starter set. automated point-of-care device for measuring glucose in capillary whole blood.	7 Pc
Composition (Per set):	Main unit Standard accessories	
Features/Performance Specifications	**Main unit** • Automated point-of-care device for measuring haemoglobin in capillary whole blood • Technology: Dual wavelength optical absorption photometry (506 nm and 880 nm for Hb measurement and turbidity compensation). • Both wavelengths are at isobestic points for oxyhaemoglobin and deoxyhaemoglobin • Provides direct reading from inserted micro-cuvette. • Removable micro-cuvette holder to allow for easy cleaning • Accuracy: Correlation with the ICSH (International Committee for Standardisation in Haematology) reference method: 0.99 • Factory calibrated and built-in auto self-test each time the device is switched on. Needs no further calibration. • Sample size: 5 to 10uL capillary, venous or arterial whole blood. • Sample collected directly from skin surface into single-use micro-cuvette by capillary action. • Uses dedicated micro-cuvette (closed system). • Measuring range: 0 to 26 g/dL. • Read-out in: g/dL or mmol/L • Reading time: <20 sec.	

	• Display informs: haemoglobin reading, reading errors, systems errors, battery status. • Symbols in display facilitate multi-lingual understanding • Interfaces: RS 232 to printer or computer. • Power supply: 220V /50 or 4 x type AA / R6 1.5 V batteries (allowing for 100h to 150h continued usage) **Standard Accessories** • 1x Point-of-care haemoglobin meter • 1 x Set of 200 micro-cuvettes (4x50) • 1 x Box of 200 safety lancets (sterile single-use, auto-disable, incision 2.25 mm) • 1 x Set of 5 cleaners • 1 x Set of 4 AA / R6 1.5 V batteries (separately packed) • 1 x power adaptor • 1 x Storage and transportation box (hard-case) • 1 x Instructions for use in English, including a pictogram showing step-by-step the measurement procedure • 1 x Operator's manual in English	
Standards and Certifications	• CE Mark, MDD93/42/EEC	

Lot 2: Supply and delivery of NICU equipment

Device Name	Oxygen Concentrator 10LPM Dual Flow	Quantity
General description	An oxygen concentrator is a device used to produce an enriched oxygen gas from atmospheric air; for use by people requiring medical oxygen due to low oxygen levels in their blood.	13 Pc
	• Main unit: 1No.	

Composition (Per set):	• Standard accessories: 1 set
Features/Performance Specifications	**Main unit** Oxygen concentrator capable of producing medical-grade oxygen from atmospheric air. The unit should be mobile on castors and capable of supplying oxygen to two patients at a time. It should incorporate an oxygen monitor facility complete with patient tubings. **Detailed Specifications** • Oxygen production technology: Pressure Swing Adsorption (PSA) • Oxygen outlets: Dual flow complete with flow meter • Oxygen flow rate: 2 – 10 LPM • Oxygen concentration: 95%±5% at maximum flow range. • Oxygen purity: Medical grade, oil free oxygen, 95%±5% oxygen concentration at maximum flow rate. • Alarm system alert: Audible/Visible light alarm on low oxygen concentration and power failure. • Safety: Shutdown with power failure, high or low pressure and high temperature. • Dimensions (mm): Approx. 800(H) x 500(W) x 400(D) • Power supply: 100 - 240VAC, 50Hz with an inbuilt suitable surge Protector or Stabiliser suitable for running oxygen concentrator. **Standard Accessories** • Power supply cable: 1No. x 3m length with 3-pin top plug (British Standard) • Nosal cannulas, peaditric
	• Humidifier bottles, 150ml x 2No.

	• Voltage Stabilizer with surge protection with 240VAC/50Hz output.	
Recommended Spare parts	• Air intake/gross particle bacterial filter: 1No. x Pack of 5Pcs	
	• Fine/final bacterial filter: 1No. x Pack of 5Pcs	
	• Fuses: Set	
Recommended Consumables	Oxygen tubings – Adult and Neonatal sizes	
Standards and Certifications	CE Mark, ISO9001, ISO13485	
Device Name	**Infusion Pump**	**Quantity**
General item description	Device used in clinical settings to administer set amount of drug or fluid to a patient over a set duration of time including continuous infusion of very small amounts.	2 Pc
Composition (Per set):	• Main unit: 1No. • Standard accessories: 1 set	
Features/Performance Specifications	**Main unit** • Microprocessor controlled infusion pump with in-built battery. • Flow rate: 0.1 to 999 ml/h or wider (in 1 ml/h increments) • Flow rate accuracy: ±5% or better • Volume increament limits range: 0.1-100ml, in 0.1 step. • Keep vein open (KVO) rate: 1 to 5ml/h • Syringe size range: 2 to 500ml • Safety/Alarm: Door open, occlusion, Air in line, improper set loading, low battery charge. • Anti-bolus system: Free flow protection, air-trapping capability, needle-less protection. • Syringe size detection: equipped	

	• Self check: Equipped • Giving set: Compatible with available giving sets on the local market [bidder should consult with National Medical Stores (NMS)] • Power supply: AC 100-240VAC/50Hz. • Battery capacity: At least 4 hours of continuous operation. • Net Weight: 2.1 kg Approx. **Standard Accessories** • Stand: 1No. • Power supply cable: 1No. • Giving set: 1No. set	
Recommended Spare parts	• Clamps: Set	
	• Fuse: Set	
Recommended Consumables	Giving set: 20No.	
Standards and Certifications	CE Mark, ISO9001, ISO13485, IEC60601-1	
Device Name	**Suction Machine**	**Quantity**
General item description	A device that removes body secretions (e.g. mucus, saliva, blood, pus).	9 Pc
Composition (Per set):	Main unit: 1	
	Standard Accessories: 1 Set	
Features/Performance Specifications	**Main unit** Suction machine suitable for use in theatre, for both adult and pediatric patients, constructed from non-corrosive material for use in a hospital environment and with a top push handle. **Performance Specifications** • Suction bottle: 2No. x 300ml polycarbonate autoclaveable jars with overflow valve.	

	• Flow rate: 40LPM or higher. • Suction vacuum pressure : 670mmHg, minimum. • Operation: Equipped with foot-operated switch and automatic stop function. • Castors: Equipped with electrically insulated and antistatic castors, Ø60mm, with 2 No. being lockable • Vacuum gauge: Graduated in mmHg and kPa; with suction control valve and marked vacuum gauge. • Anti-bacterial filters: Equipped. • Safety: Overflow pump protection. • Noise level: 50db Maximum • Power requirements 100-240VAC/50 Hz **Standard Accessories** • Power supply cable: 1No. • Suction tubing: Antistatic neoprene, with appropriate sizes for Adults and Neonates	
Recommended Spare parts	• Bacterial filters: 20 No.	
	• Fuse: Set	
	• Bottles/Jar: Pair	
Recommended Consumables	Suction tubing, antistatic neoprene: 25 sets of each size.	
Standards and Certifications	CE Mark, MDD93/42/EEC	
Device Name	**Phototherapy Unit**	**Quantity**
General description	Device for treatment of neonatal jaundice through exposure of the skin to UV light to convert unconjugated bilirubin molecules into water-soluble isomers that can be excreted by the usual pathways.	9 Pc
Composition (Per set):	• Main unit with UV light source: 1 No. • Standard Accessories: 1 Set	

Features/Performance Specifications	**Main unit and UV light source** • Stand: Height Adjustable - Approx. 1,100 - 1,500 mm • Treatment distance range: 25 to 45cm. • Irradiation angle: Adjustable (horizontal to vertical position) • Light Shading Plate: Equipped. • Cooling Fan: Equipped • UV Light source type: 4No. x LED tubes, medical grade. • UV light Wave length: 420 – 470 nm • Light source tilt angle: Continuous up to ±90 degree covering the entire treatment area. • Timer: In-built non-resettable timer. • Baby bed: Transparent with up/down Tiltable mechanism. • Power supply: 100 – 240 VAC/50Hz. • Safety: Should be supplied with safety test certificate from a competent authority. **Standard Accessories** • Eye-mask size M • Power supply cable.	
Recommended Spare parts	• UV LED tubes: 1 set • Fuses: 1 set	
Recommended Consumables	• Eye-mask size M: 10pcs	
Standards and Certifications	CE Mark, FDA (US) /STQC CB certified, STQCS certified or supplied with a functional safety test report from ERTL.	
Device Name	**Infant Incubator**	**Quantity**
Device description	A device used to maintain environmental conditions suitable for a neonate (newborn baby) especially for preterm births and for some ill full-term babies.	14 Pc
	• Main unit	

Composition (Per set):	• Standard accessories	
Features/Performance Specifications	**Main Unit** • Construction: Mobile baby incubator on an adjustable stand with 4No. x Ø100mm conductive castors; with at least 2 lockable. • Drawers: 2 Lockable drawers for storage.	
	• Control system: Servo and manual temperature control. • Temperature control range: Skin – 34 to 38 degrees centigrade; Air - 25 to 38 degrees centigrade. • Hood: Front door drop-down front access panel. • Access Ports: 3 entry access doors or more. • Humidity measuring range: 15 to 90% or wider. • Baby tray: Slide out baby tray with 65x37cm mattress. • Infant tilt angle: ±10 Degrees • Temperature fluctuation range: ±0.5 degrees Celsius • Warm up time: Less than 30 min. from 25 degrees Celsius • Alarms: Visible/Optical and acoustic alarm for pre-set skin or air temperature failure, fan motor failure and power failure. • Internal noise levels: <50db • Power: 100 - 240VAC/50 Hz, • Energy Consumption: 650VA **Standard Accessories** • Drip stand, 2 hooks minimum: 1No. • Medicine tray: 1No. • Mattress: 1No. • Skin temperature detection probe: 1No.	

	• Power supply cable: 1 No.	
Recommended Spare parts	• Skin temperature detection probe: 1No.	
	• Coarse particle filter: 10-pack box	
	• Fuses: 1 set	
Standards and Certifications	CE Mark, ISO9001, ISO13485, IEC60601-1	
Device Name	**Infant warmer**	**Quantity**
General item description	**Purpose of use** Used to provide radiant warmth for infants and neonates suffering from hypothermia (severe heat loss) thereby maintaining the body temperature of the baby and limit the metabolism rate at 37 degrees Celsius. **General item description** Servo-controlled infant warmer for providing radiant warmth for infants and neonates suffering from hypothermia (severe heat loss).	15 Pc
Composition (Per set):	Composition (Per set): • Main unit • Stand Accessories	
Features/ Performance Specifications	**Features/Performance Specifications** **Main Unit** • Construction: T-shaped base, with a height adjustment mechanism for the infrared (IR) unit to be used on beds at different heights. • Control system: Servo control, manual control • Modes: Pre-warm, baby and manual modes • Temperature setting range (skin): 34.0 to 38.0℃ • Skin temperature sensor range: 30.0 - 42.0℃ • Baby bed: Equipped with an x-ray cassette tray under the bed. • Alarm system: Skin temperature overheat and skin temperature sensor exfoliation.	

	• Lighting: LED lamps • Heat irradiation system: IR radiation quartz heater with parabolic reflector. • Capacity of Heater: 500W Maximum. • Temperature measurement and display system: Thermistor probe and LED display. • Control panel: Excellent ergonomic design with soft-touch keys. • Power supply: 100 – 240vVAC/50Hz. **Accessories** • Power supply cable • IV pole and Medicine trays • Skin temperature sensor/probe	
Recommended Spare parts	Skin temepererutre sensor 2	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**CPAP**	**Quantity**
General item description	**GENERAL DESCRIPTION** Continuous Positive Airway Pressure (Bubble CPAP) system for the non-invasive respiratory support of normal, premature or low-birth-weight neonates with spontaneous breathing. The unit has an integrated pulse oximeter to measure the level of arterial oxygen saturation in blood. **Intended Use** An electrically-powered device designed to deliver high-flow (exceeding peak inspiratory flow) heated and humidified ambient air or air/oxygen to a neonatal patient as part of non-invasive ventilation (NIV); it may additionally be used with a water tank to produce bubble continuous positive airway pressure (bubble CPAP). It consists of a gas flow generator that entrains room air, an oxygen input, a heating element,	6 Pc

	and humidification chamber; it does not include CPAP controlling pressure sensors (i.e., not a full CPAP unit). It is intended for use by a healthcare provider on a spontaneously breathing patient in hospital settings.	
Composition (Per set):	Main Unit Accessories	
Features/ Performance Specifications	**Technical specifications** • The unit supports bubble CPAP Mode. • Specifically, for support of neonates and newborns. • The unit has an integrated air compressor. • The pressure/flow of the CPAP generator is regulated electronically. • The unit has an integrated humidifier. • The unit is equipped with an electronic air/oxygen mixer. • The unit has an integrated FiO_2 analyser. • The unit accepts inlet gas supply pressures between 1.7 – 4.1 bar (25 to 60 psi). • DISS O_2 connection. (Other type connectors available indicate when ordering). • Supplied with pole mounting system, wheeled and with brakes. • The pole mounting system is equipped with 4 antistatic swivel castors, of which two castors. have been equipped with brakes. • Equipped with an air filter and a water trap. • Suitable for heated and non-heated closed patient circuits. • The unit accepts other than the manufacturer's patient circuits.	

	• All components of the system (single-use filters and patient circuits excluded) are suitable for disinfection with hospital-grade products. • CPAP pressure range adjustable from 0 to 10 cm H2O. • Maintains constant CPAP at outlet. • A minimum output flow range adjustable between: 4 – 9 L/min. • Oxygen concentration adjustable between: 21 - 100 %. • Pressure indicator in cmH2O. • Noise emission less than 50 db. • Including a clamp for a rail and/or pole. • External electric heating and humidifying chamber. • Relative humidity output up to 100%. • Adjustable output temperature up to 36°C. • Power requirements: 100 - 240 Volts - 50/60 Hz (not necessarily in a single unit). **Alarms** • Alarms are audible as well as visual. • The unit has alarms for the following: - High temperature. - System failure. - High/low oxygen concentration (FiO_2). - High airway pressure. - Failure in either air or oxygen supply. - Mains power failure. - Low battery. **Safety provisions** • Overheating protection. • The unit is equipped with a rechargeable battery which provides for a minimum of 1 hour of autonomous operation.	

	• In case of power failure, the unit switches automatically to battery power. • The unit automatically recharges on reconnection to mains. **Integrated pulse oximeter** • The unit is equipped with an integrated pulse oximeter. • Saturation accuracy between 70% and 100% is ±3%. • Measurement range SpO_2: 1% to 100% saturation. • Perfusion index: 0.02% to 20%. • Pulse rate range: 25 to 240 bpm. • Pulse rate accuracy is ±5%	
Recommended Spare parts	**Supplied with** • Instructions for assembly, use and maintenance in English. • 1 x Plastic protective dustcover. • 4 x reusable patient circuits. • 2 sets of extra small size reusable CPAP head bonnets. • 2 sets of small size reusable CPAP head bonnets. • 2 sets of medium size reusable CPAP head bonnets. • 10 x nasal cannulas, soft and atraumatic, for premature neonates. • 10 x nasal cannulas, soft and atraumatic, for term neonates and older infants. • 1 x set of hoses for connecting external oxygen and medical air supply, length > 2 m.	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Vein Finder**	**Quantity**
General item description	Purpose of use A vein finder is used to locate and visualize the location of veins on a patient's body in real-time when they are unable to easily be located with the naked eye.	18 Pc

Composition (Per set):	Composition (Per set): • Main unit • Standard Accessories	
Features/ Performance Specifications	**Specifications** • Type: Handheld. • Function: Determining vein location and Visualization of those locations on the patient's body in real-time. • Light Type: Near-infrared light. • Infrared Wavelengths: 850 nm. • Image Resolution: 720 x 480. • Visible Vein Size: 1 mm or more. • Accuracy: 0.25 mm. • Depth of Visible Vein: 12 mm or less. • Projection Distance: 200 ± 20 mm. • Power Source: Rechargeable Li-ion Battery • Power Supply: 100-240VAC/50Hz.	
Standards and Certifications	CE Mark, ISO 13485, ISO 9001, ISO60601	

Lot 3: Supply and delivery of emergency services equipment

Device Name	**Patient Trolley**	**Quantity**
General item description	**Purpose of use** Designed specifically for patient transport, 2 sections **General item description** Patient trolley with side rails and stretcher	18 Pc
Composition (Per set):	One Main unit	
Features/ Performance Specifications	• Features/Performance Specifications • Heavy duty carriage mounted on 4 swivel castors, two with brake and four with anti-static wheel.	

	• Backrest angle adjustable via secured pawl and gear ratchet, safe for patient and operator • Transfer bars connect all lower distal portions of the 4 castors, providing maximal structural strength • With fold-away side rails • Head-end side has a removable height adjustable IV-pole; height is set with a robust clamp with a heavy knob • Fixings of the fold-away side rails and IV-pole are solid steel and welded to the frame of the stretcher • Material: • Frame: epoxy-coated tubular steel • Cover: plastic, flexible highly tear resistant, anti-static, flame retardant, disinfectant- and liquid proof, washable • Caster frame/bracket: steel or nylon • Dimensions: • Stretcher, two sections extended, including upholstery: 160-200x50-60x72-88cm (l x w x h) • Fold away side rails: 50-100 x 30-45cm (l x h) • Frame: 2.7-3.3cm (outside, across), 1.8-2.2mm (thickness) • Swivel castor wheel: 2.3-3 x11-15cm (w x diameter) • Upholstery: 4.5-55cm (h) • Carrying capacity: minimum 160 kg	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Emergency Unit Patient Monitor 5 Parameter with Stand**	**Quantity**
General item description	Portable 5-parameter patient monitor. Used to measure and monitor patients' basic physiologic parameters (“vitals”) and display them on screen.	23 Pc

Composition (Per set):	Composition (Per set): • Main unit • Standard accessories	
Features/ Performance Specifications	Purpose of use Features/Performance Specifications Main unit • The monitor should be equipped with appropriate software. • Display: At least 15 inches' touch screen TFT display with a minimum of 6 waveforms • Measured parameters: SpO2, ECG, NIBP, Temperature, Respiratory rate, Heart rate. • Resolution of screen: 1200x700, minimum. • NIBP: (Manual/Auto/STAT); • Measurement type: (adult, pediatric, neonatal); • Measurement range: Systolic, Diastolic, Mean; Mean Values - (2.6 - 35.0 kPa); Resolution - (0.1 kPa); Accuracy - (± 0.4 kPa or 5 %); • Spo2: Measurement range: 0 - 100 % • Pulse measurement range: 30- 250 bpm; • BPM Accuracy: ± 2 %; • Temperature: surface and rectal; Measurement range -Minus 1 to 45C; Accuracy: ± 0.1 • Alarms: Alarm sound (Crisis/warning/Advisory); • Alarm silence; Alarm suspend; Alarm Volume adjustable; Vital sign alarm; Arrhythmia; Technical alarms. • Up to five hours of battery backup. • rend recorder storage of at least 48 hours. • Power Supply: 100-240VAC/50Hz.	
Recommended Spare parts	**Standard Accessories** • Trolley on castor wheels with brakes.	

	• ECG/Resp: 5 lead ECG cables with Clip – 2 sets per monitor • NIBP; Adult cuff – 2No • NIBP Pediatric Cuffs – I set • SpO2: Adult SpO2 sensors – 2 No. • Temperature Probe: Central temperature Probe- 2 No. • Skin temperature probe – 2 No. • Recommended Consumables • NIBP; Adult cuff – 1 set • NIBP Pediatric Cuffs – 1 set • SpO2: Adult SpO2 sensors – 2 No. • Temperature Probe: Central temperature Probe- 2 No • Skin temperature probe – 2 No. • ECG electrodes – 50Pc pack	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Resuscitation Bed 3 function**	**Quantity**
General item description	**Purpose of use** To provide safety, comfort and mobility for hospitalized patients in need of health care.	10 Pc
Composition (Per set):	Main unit	
Features/ Performance Specifications	**Main Feature** • Backrest, Footrest and Trendelenburg movements by 3 cranks • Epoxy-coated mild steel bed frame • 3 section bed board • ABS side rails with angle indicator • ABS head and footboard with holes for moving the bed easily	

	• Dia125mm noiseless castors, all with individual brakes • Bumper castors on 4 corners for protection **Accessories** • IV pole with adjustable height • Drainage hooks at the side of the bed **Specification** • Overall size（L*W*H） 2160*960*500mm • Back section adjustable 0-75° • Foot section adjustable 0-40° • Trendelenburg 0-12° • Caster Dia with break 125mm (5") • Safe working load of 140-240 kg	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Emergency Crush trolley**	**Quantity**
General item description	Emergency response trolley with work surface and storage space Heavy duty carriage mounted on 4 swivel castors, two with brake and four with anti-static wheel.	19 Pc
Composition (Per set):	Main unit 1	
Features/ Performance Specifications	Emergency trolley • General Description Trolley, emergency, with drawers **Technical Specifications:** • Work surface with elevated edges, along one length and both widths, is finished with an anti-slip layer • Long edge has 5 side-to-side drawers, for storage of medicine, renewable and small equipment.	

	• Front of each drawer fit with prefixed holder for content identification strip • Drawers have recessed finger grip-hold, leaving the front entirely flat • One central lock secures all drawer • The key is unique for every single trolley, key-bow folds away from the key-blade avoiding breakage when inside the lock • The inside of the drawers is customizable, with organizer dividers • Integrated fitting with lid for 2 waste baskets (4.5-6L each, the total capacity is 9-12L) and for 2 sharps containers (1.0-2.0L each, the total capacity is 2.0-4.0L). • A flat worktop slides in/out from underneath the work surface, extending the workspace • One short side is fit with a push-bar handle • Both castors with brakes are mounted at the push-bar handle side • With protective bumpers at all four corners **Materials:** • High resistance to corrosion (tropical environment) • Frame, side panels, base and drawers: epoxy-coated steel plate, ABS or equivalent polymer • Push handle: Austenitic stainless steel 18/10 • Worktop extension: ABS or equivalent polymer • Caster frame/bracket: steel or nylon • Caster brake: total-lock type (wheel and rotational lock) • Caster wheel: single wheel, mold-on type, non-hooded (for easy maintenance), anti-static (for 4)	

	• Wheel bearing: sealed bearing in the swivel and the wheel • Swivel is ball-bearing **Dimensions:** • Trolley, overall: 80-86 x 51-60 x 94-105 cm (l x w x h) • Frame, drawers and panels 1.5- 2.0mm thickness • Elevated edges, work surface: 1.2-5cm (h) • Worktop extension: 37-40 x 39-40 cm (l x W). • Upper drawers: 7-10cm (h) • Second drawers: 12-15 cm (h) • Third drawers: 12-15 cm (h) • Base drawers: 21-27 cm (h) • Swivel castor wheel: 2.3-3x10-12.5 cm (w by diameter) • Carrying capacity: 100-120kg • Knockdown construction: No. **Items supplied with:** • 1 x complete set of tools required for assembly • 1 x set of organizers for each drawer • 1 x set of 2 keys, unique per trolley **List of accessories and parts** Detailed step-by-step instructions for assembly and safe use, text-free pictorial based (i.e. line-drawings only).	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Wheel Chair, Adult**	**Quantity**
General item description	For moving patients who are unable to walk on their own or persons with disability. General item description Foldable wheel chair with foot rest.	27 Pc
Composition (Per set):	Composition (Per set): • Main unit	

	• IV Pole	
Features/ Performance Specifications	**Features and specifications:** Main Unit • Type: Foldable wheelchair for adults • Front wheels: Free rolling, 360 degrees swivel. • Rear Wheels: Equipped with hand-operated brakes. • push-handles push push-handles push handles at the rear. • plated chrome-plate-coated epoxy-coated tubular steel. • IV Pole: Equipped. • Frame, diameter: 2.2cm. • Frame: 2.2cm (outside, across), 1.2mm thickness • Side-to-side legs support: Yes • Swing-away foot and arm supports: Equipped • Upholstery for Armrests, seat and back: Washable plastic, flexible, tear resistant, anti-static, flame retardant, disinfectant and liquid proof. • Tyres: heavy-duty solid rubber • Carrying capacity: 150 Kg. **Dimensions** • Overall Dimension (D x W x H).: 45 x 60 x 89 cm • Seat Depth: 42-43cm • Back Rest: 43.5-49 x4 0.5-42 cm (W X H). • Side-to-side legs support: 47-50 x 8.5-23 cm. • Front Wheel diameter: 18-22cm, • Rear Wheel diameter: 58-60cm.	
Standards and Certifications	CE Mark, MDD93/42/EEC, ISO13485	
Device Name	**Nebulizer, Ultrasonic**	**Quantity**
	Purpose of use to administer medication in the form of a mist inhaled into the lungs during the treatment of asthma, cystic	20 Pc

General item description	fibrosis, Chronic obstructive pulmonary disease and other respiratory diseases or disorders. Should be a heavy-duty ultrasonic Nebuliser suitable for hospital use.	
Composition (Per set):	Composition (Per set): • Main unit • Standard Accessories	
Features/ Performance Specifications	**Features/Performance Specifications** • Type: Ultrasonic Nebulizer • Design: Compact, lightweight, low noise, and easy to handle. • Mist particle size: 1-5 microns • maximum pressure 2-2.5 bars • Mist flow rate: 15 - 20 LPM • Operational Frequency: 1.7 MHz • Water tank capacity: 250 cc or more • Medical Cup capacity: 50 cc or more • Minimum continuous run time: 1 hour or more. • Power consumption: 60W or less. • Power supply: 100-240VAC/50Hz. • Supplied with ; • Adult mask: 2pc • Child mask: 2pc • Air tube: 2pcs • Mouthpiece with exhaust valve: 2pcs • T-piece: 2pcs • Spare air filters: Pack of 5pcs • Medical cup: 5pcs	
Standards and Certifications	CE Mark, ISO 9001, IEC60601, ISO13485	

1.2 Requirements for the ancillary services

Place of delivery

The equipment shall be delivered to the address (es) within Rwenzori and Busoga region below in accordance with the delivery schedule

Rwenzori region

S/N	Health facilities	Distance in Kilometres
1.	Fort Portal Regional Referral Hospital	295 Km from Kampala
2.	Ruteete HCIV	15.1 Km from Fortportal
3.	Kataraka HCIV	5 Km from Fortportal
4.	Kyegegwa GH	105 Km from Fortportal
5.	Bwera GH	131 Km from Fortportal
6.	Bujubuli HCIV	123 Km from Fortportal
7.	Rwesande HCIV	55Km from Fortportal
8.	RUKOKI HCIV	76 Km from Fortportal
9.	Bukuku HCIV	11 Km from Fortportal
10	Nyamirami HCIV	20 Km from Kasese

Busoga region

S/N	Health facilities	Distance in Kilometres
1.	Jinja Regional Referral Hospital	96 km from Jinja
2.	Walukuba HCIV	6 km within Jinja City
3.	Mpumudde HCIV	5.9 Km from Jinja City
4.	Bugembe HCIV	9 Km from Jinja
5.	Budondo HCIV	24 Km from Jinja
6.	Buwenge HCIV	37 Km from Jinja
7.	Buwenge GH	37 Km from Jinja
8.	Kamuli GH	63 Km from Jinja
9.	Namwendwa HCIV	16.4 Km from Kamuli town
10.	Nankandulo HCIV	34.8 Km from Kamuli town

Delivery Schedule

Lot 1: Supply, delivery, installation, testing and user training of Cemonc medical equipment.

No.	Equipment	Namwendwa	Nankandulo	Buwenge	Budondo	Bugembe	Mpumudde	Walukuba	Kamuli GH	Buwenge GH	Jinja RRH	Fortportal RRH	Kyegegwa GH	Bwera GH	Bujubuli HCIV	Nyamirami HCIV	Rukoki HCIV	Rwesande HCIV	Ruteete HCIV	Bukuku HCIV	Kataraka HCIV	Total
1.	Oxygen concentrator 10LPM	0	1	1	1	1	0	0	2	1	0	0	1	0	0	0	0	0	0	0	0	8
2.	Vacuum extractor	1	0	1	1	1	0	0	1	1	0	1	1	1	1	0	0	1	1	1	1	14
3.	Dilation and Curettage set	1	0	2	2	0	0	0	2	2	0	2	0	0	0	0	0	0	1	1	1	14
4.	Portable Ultrasound scanner	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1
5.	Universal Anaesthesia machine	0	0	0	0	0	0	0	1	0	1	0	0	0	0	1	1	1	1	0	0	6

6.	Autoclave 75Ls. (Sterilizer)	1	1	0	0	1	0	0	0	1	0	0	1	0	0	0	0	0	0	0	0	5
7.	operation table, hydraulic /Theatre bed	1	1	0	1	0	0	0	0	0	0	0	0	0	0	0	0	1	0	1	0	5
8.	Operating Light mobile [LED]	1	1	1	1	1	0	0	0	0	0	0	1	0	0	0	0	1	0	1	0	8
9.	Delivery beds	0	0	0	0	2	0	0	1	0	2	2	0	0	0	0	0	1	0	0	0	8
10.	Wheel Chair Adult	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1
11.	Patient stretcher/ trolley	1	1	1	1	1	0	1	0	0	1	2	2	2	2	2	2	2	0	1	2	24
12.	The patient monitors 5 Parameters with a stand	0	1	1	1	1	0	0	2	1	0	0	1	0	0	0	0	0	0	0	0	8
13.	Foetal Doppler	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	20
14.	Patient bed	0	1	1	1	1	0	0	2	1	0	0	1	0	0	0	0	0	0	0		8

15.	Blood Refrigerator	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	1
16.	HemoCue	1	0	1	0	0	0	1	0	0	0	0	0	0	0	0	0	1	1	1	1	7

Lot 2: Supply, delivery, installation, testing and user training of NICU medical equipment.

No.	Equipment	Namwendwa	Nankandulo	Buwenge	Budondo	Bugembe	Mpumudde	Walukuba	Kamuli GH	Buwenge GH	Jinja RRH	Fortportal RRH	Kyegegwa GH	Bwera GH	Bujubuli HVIV	Nyamirami HCIV	Rukoki HCIV	Rwesande HCIV	Ruteete HCIV	Bukuku HCIV	Total
1.	Oxygen concentrator 10LPM Dual flow	2	0	2	2	2	2	2	0	0	0	0	0	0	0	0	0	0	0	1	13
2.	Infusion pump	0	0	0	0	0	0	1	1	0	0	0	0	0	0	0	0	0	0	0	2
3.	Suction machine	0	0	1	1	1	1	1	1	1	1	0	0	0	0	0	0	0	0	1	9
4.	Phototherapy Unit	1	1	1	1	1	0	1	0	0	1	0	0	0	0	1	1	0	0	0	9
5.	Infant Incubator	1	0	2	2	2	0	2	0	0	0	0	1	1	0	1	1	0	0	1	14
6.	Infant Warmer	1	2	2	2	2	0	2	1	0	0	0	0	0	0	1	1	0	0	1	15
7.	CPAP	0	1	0	0	0	0	0	1	1	2	0	1	0	0	0	0	0	0	0	6
8.	Vein Finders	1	1	1	1	1	1	1	1	1	2	2	1	1	0	1	1	0	0	1	18

Lot 3: Supply, delivery, installation, testing and user training of Emergency services medical equipment.

No.	Equipment	Fortportal RRH	Kyegegwa GH	Bwera GH	Bujubuli HCIV	Nyamirami HCIV	Rukoki HCIV	Rwesande HCIV	Ruteete HCIV	Bukuku HCIV	Kataraka HCIV	Namwendwa HCIV	Nankandulo HCIV	Buwenge HCIV	Budondo HCIV	Bugembe HCIV	Mpumudde HCIV	Walukuba HCIV	Kamuli GH	Buwenge GH	Jinja RRH	Total
1.	Patient trolley	0	0	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	18
2.	Emergency Unit patient monitor 5 parameter	2	2	2	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	23
3.	Resuscitation Bed 3 function	0	0	0	0	1	1	0	0	1	0	1	1	1	1	1	1	1	0	0	0	10
4.	Emergency crash trolley	0	1	1	1	1	1	0	1	1	1	1	1	1	1	1	1	1	1	1	2	19
5.	Wheelchair Adult	2	2	2	1	1	1	1	1	1	1	1	1	2	1	1	1	1	2	2	2	27
6.	Nebulizer Ultrasonic	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	20

Installation and User training

At the time of delivery, the contractor shall carry out user training and testing that shall come with installation of the equipment. The contractor shall provide user manuals for the equipment.

After sales services

The contractor shall all times have in stock spare parts for the equipment above to fulfil the contracting authority's order as and when needed.

The contractor shall also engrave the equipment before delivery to the various health centres. The contracting authority shall provide the codes for the engravement.

2 General provisions

2.1 Contracting authority

The contracting authority of this public contract is Enabel, the Belgian development agency, public-law company with social purposes, with its registered office at Rue Haute 147, 1000 Brussels in Belgium (enterprise number 0264.814.354, RPM/RPR Brussels). Enabel has the exclusive competence for the execution, in Belgium and abroad, of public service tasks of direct bilateral cooperation with partner countries. Moreover, it may also perform other development cooperation tasks at the request of public interest organisations, and it can develop its own activities to contribute towards realisation of its objectives.

For this procurement contract, Enabel is represented by person(s) who shall sign the award letter and are mandated to represent the organisation towards third parties.

2.2 Institutional framework of Enabel

- The general framework of reference in which Enabel operates is:
  - The Belgian Law on Development Cooperation of 19 March 2013[1];
  - The Belgian Law of 21 December 1998 establishing the Belgian Technical Cooperation as a public-law company[2];
  - The Belgian Law of 23 November 2017 changing the name of the Belgian Technical Cooperation and defining the missions and functioning of Enabel, the Belgian development agency, published in the Belgian Official Gazette on 11 December 2017.

The following initiatives are also guiding Enabel in its operations and are given as main examples:

- In the field of international cooperation: the United Nations Sustainable Development Goals and the Paris Declaration on the harmonisation and alignment of aid;
- In the field of the fight against corruption: the Law of 8 May 2007 approving the United Nations Convention against Corruption, adopted in New York on 31 October 2003[3], as well as the Law of 10 February 1999 on the Suppression of

[1] Belgian Official Gazette of 30 December 1998, of 17 November 2001, of 6 July 2012, of 15 January 2013 and of 26 March 2013.

[2] Belgian Official Gazette of 1 July 1999.

[3] Belgian Official Gazette of 18 November 2008

Corruption transposing the Convention on Combating Bribery of Foreign Public Officials in International Business Transactions;

- In the field of Human Rights: the United Nations' Universal Declaration of Human Rights (1948) as well as the 8 basic conventions of the International Labour Organization[4] on Freedom of Association (C. n°87), on the Right to Organise and Collective Bargaining (C. n°98), on Forced Labour (C. n°29 and 105), on Equal Remuneration and on Discrimination in Respect of Employment (C. n°100 and 111), on Minimum Age for Admission to Employment (C. n°138), on the Prohibition of the Worst Forms of Child Labour (C. n°182);
- In the field of environmental protection: The Climate Change Framework Convention of Paris, of 12 December 2015;
- The first Management Contract contracting Enabel and the Belgian federal State (approved by the Royal Decree of 17 December 2017, Belgian Official Gazette of 22 December 2017) that sets out the rules and the special conditions for the execution of public service tasks by Enabel on behalf of the Belgian State.
- Considering Enabel's Code of Conduct of January 2019, Enabel's Policy regarding sexual exploitation and abuse of June 2019 and Enabel's Policy regarding fraud and corruption risk management of June 2019;

2.3 Rules governing the public contract

- The following, among other things, apply to this public contract:
- The Law of 17 June 2016 on public procurement[5];
- The Law of 17 June 2013 on justifications, notification and legal remedies for public contracts and certain contracts for works, supplies and services[6];
- The Royal Decree of 18 April 2017 on the awarding of public contracts in the classic sectors[7];

[4] https://www.ilo.org/global/standards/lang--en/index.htm

[5] Belgian Official Gazette 14 July 2016.

[6] Belgian Official Gazette of 21 June 2013.

[7] Belgian Official Gazette 9 May 2017.

- The Royal Decree of 14 January 2013 establishing the General Implementing Rules for public procurement and for concessions for public works[8];
- Circulars of the Prime Minister with regards to public procurement.
- All Belgian regulations on public contracts can be consulted on www.publicprocurement.be.
- Enabel's Policy regarding sexual exploitation and abuse – June 2019
- Enabel's Policy regarding fraud and corruption risk management – June 2019
- Regulation (EU) 2016/679 of the European Parliament and of the Council of 27 April 2016 on the protection of natural persons with regard to the processing of personal data and on the free movement of such data (General Data Protection Regulation, hereinafter referred to as 'the GDPR'), and repealing Directive 95/46/EC;
- The Law of 30 July 2018 on the protection of natural persons with regard to the processing of personal data;

All Belgian regulations on public contracts can be consulted on www.publicprocurement.be

Enabel's Code of Conduct and the policies mentioned above can be consulted on Enabel's website via: https://www.enabel.be/content/integrity-desk

2.4 Definitions

The following definitions apply to this contract:

The tenderer: An economic operator submitting a tender;

The contractor/ service provider: The tenderer to whom the public contract is awarded;

The contracting authority: Enabel, represented by the Resident Representative of Enabel in Uganda.

The tender: The commitment of the tenderer to perform the public contract under the conditions that he has submitted;

Days: In the absence of any indication in this regard in the Tender Specifications and the applicable regulations, all days should be interpreted as calendar days;

Procurement documents: Contract notice and Tender Specifications including the annexes and the documents they refer to;

[8] Belgian Official Gazette 27 June 2017.

Technical specifications: A specification in a document defining the characteristics of a product or a service, such as the quality levels, the environmental and climate performance levels, the design for all needs, including accessibility for people with disabilities, and the evaluation of conformity, of product performance, of the use of the product, safety or dimensions, as well as requirements applicable to the product as regards the name by which it is sold, terminology, symbols, testing and test methods, packaging, marking or labelling, instructions for use, the production processes and methods at every stage in the life cycle of the supply or service, as well as the evaluation and conformity procedures;

Variant: An alternative method for the design or the performance that is introduced either at the demand of the contracting authority, or at the initiative of the tenderer;

Option: A minor and not strictly necessary element for the performance of the contract, which is introduced either at the demand of the contracting authority, or at the initiative of the tenderer;

Inventory: The procurement document which splits up the performance in different items and specifies the quantity or the method to determine the price for each of them;

General Implementing Rules (GIR): Rules laid down in the Royal Decree of 14 January 2013 establishing the General Implementing Rules for public procurement and for concessions for public works;

The Tender Specifications (Cahier spécial des charges/CSC): This document and its annexes and the documents it refers to;

BDA: Belgian Public Tender bulletin;

OJEU: Official Journal of the European Union;

OECD: Organisation for Economic Cooperation and Development;

Corrupt practices: The offer of a bribe, gift, gratuity or commission to a person as an inducement or reward for performing or refraining from an act relating to the award of a contract or performance of a contract already concluded with the contracting authority;

Litigation: Court action;

Subcontractor in the meaning of public procurement regulations: The economic operator proposed by a tenderer or contractor to perform part of the contract;

Controller in the meaning of the GDPR: The natural or legal person, public authority, agency or other body which, alone or jointly with others, determines the purposes and means of the processing of personal data.

Processor (subcontractor) in the meaning of the GDPR: A natural or legal person, public authority, agency or other body which processes personal data on behalf of the controller;

Recipient in the meaning of the GDPR: A natural or legal person, public authority, agency or another body, to which the personal data are disclosed, whether a third party or not;

Personal data: Any information relating to an identified or identifiable natural person ('data subject'); an identifiable natural person is one who can be identified, directly or indirectly, in particular by reference to an identifier such as a name, an identification number, location data, an online identifier or to one or more factors specific to the physical, physiological, genetic, mental, economic, cultural or social identity of that natural person.

2.5 Confidentiality

2.5.1 Processing of personal data

The contracting authority undertakes to process the personal data that are communicated to it under the framework of this procedure with the greatest care, in accordance with legislation on the protection of personal data (General Data Protection Regulation, GDPR). Where the Belgian law of 30 July 2018 on the protection of natural persons with regard to the processing of personal data contains stricter provisions, the contracting authority will act in accordance with said law.

2.5.2 Confidentiality

The tenderer or contractor and Enabel are bound to secrecy vis-à-vis third parties with regards to any confidential information obtained within the framework of this public contract and will only divulge such information to third parties after receiving the prior written consent of the other party. They will disclose this confidential information only among appointed parties involved in the assignment. They guarantee that said appointed parties will be adequately informed of their obligations in respect of the confidential nature of the information and that they shall comply therewith.

PRIVACY NOTICE OF ENABEL Enabel takes your privacy serious. We undertake to protect and process your personal data with due care, transparently and in strict compliance with privacy protection legislation.

See also: https://www.enabel.be/content/privacy-notice-enabel

2.6 Deontological obligations

Any failure to comply with one or more of the deontological clauses may lead to the exclusion of the candidate, tenderer or contractor from other public contracts for Enabel.

For the duration of the contract, the contractor and his staff respect human rights and undertake not to go against political, cultural or religious customs of the beneficiary country. The tenderer or contractor is bound to respect fundamental labour standards, which are internationally agreed upon by the International Labour Organization (ILO), namely the conventions on union freedom and collective bargaining, on the elimination of forced and obligatory labour, on the elimination of employment and professional discrimination and on the abolition of child labour.

In accordance with Enabel's Policy regarding sexual exploitation and abuse, the contractor and his staff have the duty to behave in an irreproachable manner towards the beneficiaries of the projects and towards the local population in general. They must abstain from any acts that could be considered a form of sexual exploitation or abuse and they must abide by the basic principles and guidelines laid down in this policy.

Any attempt of a candidate or a tenderer to obtain confidential information, to proceed to illicit arrangements with competitors or to influence the evaluation committee or the contracting authority during the investigation, clarification, evaluation and comparison of tenders and candidates procedure will lead to the rejection of the application or the tender.

Moreover, in order to avoid any impression of risk of partiality or connivance in the follow-up and control of the performance of the public contract, it is strictly forbidden to the contractor to offer, directly or indirectly, gifts, meals or any other material or immaterial advantage, of whatever value, to appointees of the contracting authority who are concerned, directly or indirectly, by the follow-up and/or control of the performance of the contract, regardless of their hierarchical rank.

The public contractor commits to supply, upon the demand of the contracting authority, any supporting documents related to the performance conditions of the contract. The contracting authority will be allowed to proceed to any desk review or on-the-spot check which it considers necessary to collect evidence to support the presumption of unusual commercial expenditure. Depending on the gravity of the facts observed, the contractor having paid unusual commercial expenditure is liable to have his contract cancelled or to be permanently excluded from receiving funds.

In accordance with Enabel's Policy regarding sexual exploitation and abuse of June 2019 and Enabel's Policy regarding fraud and corruption risk management complaints relating to issues of

integrity (fraud, corruption, etc.) must be sent to the Integrity desk through the website www.enabelintegrity.be

2.7 Applicable law and competent courts

The contract must be performed and interpreted according to Belgian law.

The parties commit to sincerely perform their engagements to ensure the good performance of the public contract.

In case of litigation or divergence of opinion between the contracting authority and the contractor, the parties will consult each other to find a solution.

If agreement is lacking, the Brussels courts are the only courts competent to resolve the matter.

3 Modalities of the contract

3.1 Type of contract

This is a direct contract for the supply of goods by means of purchase.

3.2 Scope of contract

3.2.1 Subject-matter

This public supplies contract consists of the supply and delivery of medical equipment for health facilities, in conjunction with the provision of the ancillary services of installation, testing, user training and commissioning of the equipment in conformity with the conditions of these Tender Specifications.

3.2.2 Lots

The public contract has 3 lots, each of which is indivisible. The tenderer may submit a tender for one lot or all the lots. A tender for part of a lot is inadmissible.

The description of each lot is included in Part 1 of these Tender Specifications.

The lots are:

Lots	Description of the lots
Lot 1	Supply, delivery, installation, testing and user training of Cemonc medical equipment.
Lot 2	Supply, delivery, installation, testing and user training of NICU medical equipment.
Lot 3	Supply, delivery, installation, testing and user training of emergency services medical equipment.

3.2.3 Items

Each lot of this contract consists of the items mentioned in part 1 of the technical specification. These items are pooled and form one single contract per lot. It is not possible to tender for one or several items and the tenderer must submit price quotations for all items of the same lot.

3.2.4 Variants

Each tenderer may submit only one tender. Variants are forbidden.

3.3 Duration of the contract

For each of the lots, the contract shall commence upon award notification and last for a duration of 180 calendar days. Thereafter there shall be a warrant period of 1 year for all the equipment.

4 Special contractual provisions

This chapter of these Tender Specifications holds the specific provisions that apply to this public contract by way of derogation from the 'General Implementing Rules for public procurement and for concessions for public works' of the Royal Decree of 14 January 2013, hereinafter referred to as 'GIR', or as a complement or an elaboration thereof. The numbering of the articles below (between brackets) follows the numbering of the GIR articles. Unless indicated, the relevant provisions of the General Implementing Rules (GIR) apply in full.

4.1 Managing official (Art. 11)

The managing official is Ms Benedict Briot, e-mail: benedicte.briot@enabel.be

Once the public contract is concluded the managing official is the main contact point for the supplier. Any correspondence or any questions with regards to the performance of the contract will be addressed to him or her, unless explicitly mentioned otherwise in these Tender Specifications.

The managing official is responsible for the follow-up of the performance of the contract.

- The managing official is fully competent for the follow-up of the satisfactory performance of the contract, including issuing service orders, drawing up reports and states of affairs, approving the services, progress reports and reviews. He or she may order any modifications to the contract with regards to its subject-matter provided that they remain within its scope.
- However, the signing of amendments or any other decision or agreement implying derogation from the essential terms and conditions of the contract are not part of the competence of the managing official. For such decisions the contracting authority is represented as stipulated under the point Contracting authority.
- Under no circumstances is the managing official allowed to modify modalities (e.g. delivery deadlines) of the contract, even if the financial impact is nil or negative. Any commitment, change or agreement derogating the conditions in the Tender Specifications and that has not been notified by the contracting authority, will be considered null and void.

4.2 Subcontractors (Art. 12 to 15)

The fact that the contractor entrusts all or part of his commitments to subcontractors does not relieve him of liability to the contracting authority. The latter does not recognise any contractual relation with third parties.

The contractor remains, in any case, solely liable to the contracting authority. The contractor may not subcontract the contract or a part of the contract to other subcontractors than those presented at the time of submission; subcontracting to subcontractors presented in the tender is allowed only after preliminary approval by the contracting authority of these subcontractors. When the contractor uses a subcontractor to carry out specific processing activities on behalf of the contracting authority, the same data protection obligations as those of the contractor are imposed on that subcontractor by contract or any other legal act.

In the same way, the contractor will respect and enforce to his subcontractors, the provisions of Regulation (EU) 2016/679 of the European Parliament and of the Council of 27 April 2016 on the protection of natural persons with regard to the processing of personal data and on the free movement of such data (General Data Protection Regulation, GDPR). The contracting authority may conduct an audit of the processing carried out in order to validate compliance with this legislation.

4.3 Confidentiality (Art. 18)

Knowledge and information obtained by the contractor, including any persons responsible for the mission and any other person involved in this public contact, are strictly confidential.

Under no circumstances can the information collected, regardless of its origin and nature, be transferred to third parties in any form.

All parties directly or indirectly involved are therefore bound by the duty of discretion.

In accordance with Article 18 of the Royal Decree of 14 January 2013 establishing the general rules for public procurement, the tenderer or contractor undertakes to consider and process in a strictly confidential manner any information, all facts, any documents and/or any data, whatever their nature and support, which have been communicated to him, in any form and by any means, or to which he has access, directly or indirectly, in the context or on the occasion of this public contract. Confidential information covers, in particular, the very existence of this public contract, without this list being limited.

Therefore, he undertakes to:

•Respect and enforce the strict confidentiality of these elements and to take all necessary precautions in order to preserve their secrecy (these precautions cannot in any case be inferior to those taken by the tenderer for the protection of his own confidential information);

• Consult, use and/or exploit, directly or indirectly, all of the above elements only to the extent strictly necessary to prepare and, where applicable, to carry out this public contract (particularly regarding the privacy legislation with respect to personal data processing);

• Not reproduce, distribute, disclose, transmit or otherwise make available to third parties the above elements, in whole or in part, and in any form, unless having obtained prior and written consent of the contracting authority;

• Return, at the first request of the contracting authority, the above elements;

• In general, not disclose directly or indirectly to third parties, whether for advertising or any other reason, the content of this public contract, or the fact that the tenderer or contractor performs this public contract for the contracting authority, or, where applicable, the results obtained in this context, unless having obtained prior and written consent of the contracting authority.

4.4 Personal data protection

4.4.1 Processing of personal data by the contracting authority

The contracting authority undertakes to process the personal data that are communicated to it in response to the Call for Tenders with the greatest care, in accordance with legislation on the protection of personal data (General Data Protection Regulation, GDPR). Where the Belgian law of 30 July 2018 on the protection of natural persons with regard to the processing of personal data contains stricter provisions, the contracting authority will act in accordance with said law.

4.4.2 Processing of personal data by the contractor

PROCESSING OF PERSONAL DATA BY A CONTROLLER (RECIPIENT)

Where during contract performance, the contractor processes personal data of the contracting authority or in execution of a legal obligation, the following provisions apply:

For any processing of personal data carried out in connection with this public contract, the contractor is required to comply with Regulation (EU) 2016/679 of the European Parliament and of the Council of 27 April 2016 on the protection of natural persons with regard to the processing of personal data and on the free movement of such data (GDPR) and the Belgian law of 30 July 2018 on the protection of natural persons with regard to the processing of personal data.

By simply participating in the contracting process, the tenderer certifies that he will strictly comply with the obligations of the GDPR for any processing of personal data conducted in connection with that public contract.

Given the public contract, it is to be considered that the contracting authority and the contractor will each be responsible, individually, for the processing.

4.5 Intellectual property (Art. 19 to 23)

The contracting authority does not acquire the intellectual property rights created, developed or used during performance of the public contract.

Without prejudice to clause 1 and unless otherwise stipulated in the procurement documents, when the subject-matter of the public contract consists of the creation, manufacture or the development of designs or of logos, the contracting authority acquires the intellectual property thereof, as well as the right to trademark them, to have them registered and to have them protected.

For domain names created under the contract, the contracting authority also acquires the right to register and protect them, unless otherwise stipulated in the procurement documents.

Where the contracting authority does not acquire the intellectual property rights, it obtains a patent licence of the results protected by intellectual property law for the exploitation modes that are mentioned in the procurement documents.

The contracting authority lists the exploitation modes for which it intends to obtain a licence in the procurement documents.

4.6 Performance bond (Art. 25 to 33)

For each lot, the performance bond is set at 5% of the total value, excluding VAT, of procurement. The value thus obtained is rounded up to the nearest 10 euros.

In accordance with the legal and regulatory provisions, the performance bond may be constituted either of cash or of public funds or may take the form of a joint performance bond.

The performance bond may also take the form of a surety bond issued by a credit institution meeting the requirements of the law on the statute and control of credit institutions.

By way of derogation from Article 26 of the GIR the performance bond may be posted through an establishment that has its registered office in one of the countries of destination of the services. The contracting authority reserves the right to accept or refuse the posting of the bond through that institution. The tenderer mentions the name and address of this institution in the tender.

This derogation is founded on the idea of providing possible local tenderers with an opportunity to submit a tender. This measure is made essential by the specific requirements of the contract. The contractor must, within 30 calendar days from the day of contract conclusion, furnish proof that he or a third party has posted the bond in one of the ways set out below:

1° in the case of cash, by transfer of the amount to the bpost bank account number of the Deposit and Consignment Office. Fill out the following form as completely as possible: https://finances.belgium.be/sites/default/files/01_marche_public.pdf (PDF, 1.34 Mo), and return it to the e-mail address: info.cdcdck@minfin.fed.be

2° in the case of public funds, by depositing such funds, for the account of the Deposit and Consignment Office, with the State Cashier at the head office of the National Bank in Brussels or at one of its provincial agencies or with a public institution with an equivalent function

3° in the case of a joint surety, by deposit via an institution that lawfully carries out this activity of a deed of joint surety with the Deposit and Consignment Office or with a public institution with an equivalent function

4° in the case of a guaranty, by the deed of undertaking of the credit institution.

Proof is provided, as appropriate, by submission to the contracting authority of:

1° the deposit receipt of the Deposit and Consignment Office or of a public institution with an equivalent function; or

2° a debit notice issued by the credit institution; or

3° the deposit certificate issued by the State Cashier or public institution with an equivalent function; or

4° the original copy of the deed of joint surety stamped by the Depot and Consignment Office or by a public institution with an equivalent function; or

5° the original copy of the deed of undertaking issued by the credit institution granting a guaranty.

These documents, signed by the depositor, must state why the performance bond was posted and its precise usage, consisting of a concise indication of the subject-matter of the contract and a reference to the procurement documents, as well as the name, first names and full address of the contractor and, where relevant, that of the third party that made the deposit on the contractor’s account, bearing the statement 'lender' or 'mandatary', as appropriate.

The period of 30 calendar days specified above is suspended during the period of closure of the contractor's business for paid annual holidays and the days off in lieu stipulated by regulation or by a collective binding labour agreement.

Proof that the required performance bond has been posted must be sent to the address that will be mentioned in the contract conclusion notification.

Request by the contractor for the acceptance procedure to be carried out:

1° For provisional acceptance: This is equal to a request to release the first half of the performance bond;

2° For final acceptance: This is equal to a request to release the second half of the performance bond, or, in case no provisional acceptance applied, to release the whole of the performance bond.

4.7 Conformity of performance (Art. 34)

The supplies must comply in all respects with the procurement documents. Even in the absence of technical specifications in the procurement documents, the supplies must comply in all aspects with good practice.

4.8 Changes to the public contract (Art. 37 to 38/19)

4.8.1 Replacement of the contractor (Art. 38/3)

Provided that he meets the selection and exclusion criteria set out in this document, a new contractor may replace the contractor with whom the initial contract was agreed in cases other than those provided for in Art. 38/3 of the General Implementing Rules (GIR).

The contractor submits his request as quickly as possible by registered post, stating the reasons for this replacement and providing a detailed inventory of the state of the supplies and services already delivered, the new contractor's contact details and the documents and certificates which the contracting authority cannot access free of charge.

The replacement will be recorded in an amendment dated and signed by all three parties. The initial contractor remains liable to the contracting authority for the performance of the remainder of the contract.

4.8.2 Revision of prices (Art. 38/7)

For this contract, price revisions are not permitted.

4.8.3 Indemnities following the suspensions ordered by the contracting authority during performance (Art. 38/12)

- The contracting authority reserves the right to suspend the performance of the contract for a given period, mainly when it considers that the contract cannot be performed without inconvenience at that time.
- The performance period is extended by the period of delay caused by this suspension, provided that the contractual performance period has not expired. If it has expired, the return of fines for late performance will be agreed.
- When activities are suspended, based on this clause, the contractor is required to take all necessary precautions, at his expense, to protect the services already performed and the materials from potential damage caused by unfavourable weather conditions, theft or other malicious acts.

The contractor has a right to damages for suspensions ordered by the contracting authority when:

- The suspension lasts in total longer than one twentieth of the performance period and at least ten working days or two calendar weeks, depending on whether the performance period is expressed in working days or calendar days;
- The suspension is not owing to unfavourable weather conditions;
- The suspension occurred during the contract performance period.

- Within thirty days of their occurrence or the date on which the contractor or the contracting authority would normally have become aware of them, the contractor reports the facts or circumstances succinctly to the contracting authority and describes precisely their impact on the progress and cost of the contract.

4.8.4 Unforeseeable circumstances

As a rule, the contractor is not entitled to any modification of the contractual terms due to circumstances of which the contracting authority was unaware.

A decision of the Belgian State to suspend cooperation with a partner country is deemed to be unforeseeable circumstances within the meaning of this article. Should the Belgian State break

off or cease activities which implies therefore the financing of this public contract, Enabel will do everything reasonable to agree a maximum compensation figure.

4.9 Preliminary technical acceptance (Art. 42)

Products may not be used if they have not been accepted by the managing official or his or her representative.

Products that at a given stage do not satisfy the technical acceptance tests imposed will be declared unfit for technical acceptance. Upon the request of the contractor, the contracting authority in accordance with the procurement documents verifies whether the products have the required qualities or at the very least comply with good practice and satisfy the conditions of the contract. If certain products are destroyed during verification, the contractor replaces these at its own expense. The procurement documents specify the quantity of products to be destroyed.

Where the contracting authority declares that the product presented is not in the required condition for examination, the acceptance request by the building contractor will be considered not having been made. A new request is made when the product is fit for acceptance.

4.10 Performance modalities (Art. 115 et seq.)

4.10.1 Deadlines and terms (Art. 116)

For each lot, the supplies must be delivered within 180 calendar days as from the day following the date on which the supplier received the contract conclusion notification letter. The closure of the supplier's business for annual holidays is not included in this calculation.

There after a defects liability period of 365 calendar days shall be observed.

The contract conclusion notification letter is addressed to the supplier either by registered letter, fax or any other means through which the date of dispatch can be determined unambiguously.

4.10.2 Quantities to be supplied (Art. 117)

For each lot, the public contract quantities are stated in section 1 of this tender specifications document.

Without prejudice to the possibility for the contracting authority to terminate the contract if the supplies delivered do not meet the requirements imposed or if they are not delivered by the deadlines asked.

4.10.3 Place where the supplies must be delivered and formalities (Art. 149)

The supplies shall be delivered at the addresses mentioned in section 1 of the tender specifications document.

4.10.4 Packaging (Art. 119)

Packaging will become the property of the contracting authority, without the supplier having any claim to compensation in this regard.

4.10.5 Inspection of the supplies delivered (Art. 120)

The supplier delivers only goods that have no apparent and/or hidden defects and that correspond strictly to the order (in kind, quantity, quality…) and, if necessary, to the prescriptions of related documents as well as applicable regulations, in compliance with good practice, the state of the art, the highest standards of usage, of reliability and of longevity, and for the purposes that the contracting authority has in mind, which the supplier knows or at least should know.

The products are stored for delivery in the supplier's warehouses. Delivery cannot occur prior to the contracting authority's verification of the goods stored for delivery. This verification does not amount as acceptance.

Acceptance (provisional acceptance) only takes place after the complete inspection by the contracting authority of the conformity of the goods and services delivered. The contracting authority disposes of a period for verification of thirty days starting on the date of delivery. This period will begin on the day after arrival of the supplies at the place of delivery, provided that the contracting authority is in possession of the delivery note or invoice.

The signature of (a staff member of) the contracting authority, in particular in electronic reception devices, upon delivery of the goods, does consequently only count as evidence of taking possession and does not concern the acceptance of the goods.

Acceptance on the premises of each of the health facilities will serve as complete provisional acceptance for that health facility.

Acceptance implies the transfer of ownership and of risks of damage and loss.

In case of full or partial refusal of a delivery, the supplier is bound to take back, at his own costs and risks, the products refused. The contracting authority may ask the supplier to deliver goods that comply as soon as possible, either cancel the order and get supplied by another supplier.

4.10.6 Liability of the supplier (Art. 122)

The supplier shall be liable for his supplies up to the time when the inspection and notification formalities referred to in Article 120 are carried out, unless losses or damage sustained in the warehouses of the consignee are due to the events or circumstances referred to in Articles 54 and 56.

Moreover, the supplier indemnifies the contracting authority against damages for which it is liable towards third parties due to late performance of the contract or due to failure of the supplier.

4.11 Zero tolerance Sexual exploitation and abuse

In application of Enabel's Policy regarding sexual exploitation and abuse of June 2019 there will be zero tolerance towards any misconduct that could impact the professional credibility of the tenderer.

4.12 Means of action of the contracting authority (Art. 44–51 and 123–126)

The service provider's default is not solely related to services as such but also to the whole of the service provider's obligations.

In order to avoid any impression of risk of partiality or connivance in the follow-up and control of the performance of the public contract, it is strictly forbidden to the contractor to offer, directly or indirectly, gifts, meals or any other material or immaterial advantage, of whatever value, to appointees of the contracting authority who are concerned, directly or indirectly, by the follow-up and/or control of the performance of the contract, regardless of their hierarchical rank.

In case of violation, the contracting authority may impose a lump-sum fine to the service provider for each violation, which can be to up to three times the amount obtained by adding up the (estimated) values of the advantage offered to the employee and of the advantage that

the contractor hoped to obtain by offering the advantage to the employee. The contracting authority will decide independently about the application and the amount of this fine.
This clause is without prejudice to the possible application of other measures as of right provided in the GIR, namely the unilateral termination of the contract and/or the exclusion from procurement by the contracting authority for a determined duration.

4.12.1 Failure of performance (Art. 44)

§1The contractor is considered to be in failure of performance under the public contract:
1° when performance is not carried out in accordance with the conditions specified in the procurement documents;
2° at any time, when performance has not progressed in such a way that it can be fully completed on the due dates;
3° when he does not observe written orders, which have been given in due form by the contracting authority.
§2. Any failure to comply with the provisions of the public contract, including the non-observance of orders of the contracting authority, is recorded in a report ('process verbal'), a copy of which will be sent immediately to the contractor by registered mail.
The contractor must repair the defects without any delay. He may assert his right of defence by registered letter addressed to the contracting authority within fifteen days from the date of dispatch of the report (process verbal). Silence on his part after this period shall be deemed acknowledgement of the reported facts.
Any defects detected that can be attributed to the contractor render him liable to one or more of the measures provided for in Articles 45 to 49, 154 and 155.

4.12.2 Fines for delay (Art. 46 and 123)

The fines for delay differ from the penalties referred to in Article 45. They are due, without the need for notice, by the mere lapse of the performance period without the issuing of a report and they are automatically applied for the total number of days of delay.
Regardless of the application of any fines for delay, the contractor indemnifies the contracting authority against damages for which it is liable towards third parties due to late performance of the contract.

4.12.3 Measures as of right (Art. 47 and 124)

§1 When, upon expiry of the term given in Article 44, §2, the contractor has not taken action or has presented means deemed unjustified by the contracting authority, the contracting authority may apply the measures as of right described in paragraph 2.

However, the contracting authority may apply measures as of right without waiting for the expiry of the term given in Article 44, §2, when the contractor has explicitly recognised the defects detected.

§2. The measures as of right are:

1° Unilateral termination of the contract. In this case the entire performance bond, or if no bond has been posted an equivalent amount, is acquired as of right by the contracting authority as lump sum damages. This measure excludes the application of any fine for delay in performance in respect of the terminated part;

2° Performance under regie of all or part of the non-performed contract;

3° Conclusion of one or more replacement contracts with one or more third parties for all or part of the contract remaining to be performed.

The measures referred to in 1°, 2° and 3° will be taken at the expense and risk of the defaulting contractor. However, any fines or penalties imposed during the performance of a replacement contract will be borne by the new contractor.

4.13 End of the public contract

4.13.1 Acceptance of the products delivered (Art. 64-65 and 128)

The managing official will closely follow up the delivery.

The products are stored for delivery in the supplier's warehouses. Delivery cannot occur prior to the contracting authority's accepting the goods stored for delivery. The managing official who will carry out acceptance is named in the contract award notification if his/her name has not yet been mentioned in the procurement documents.

Provisional acceptance

Upon expiry of the thirty-day period specified in Article 120, §2, as appropriate, a provisional acceptance report or acceptance refusal report will be drawn up.

Partial acceptance at the place of manufacture requires a request in writing by the supplier to the contracting authority.

Provisional acceptance is carried out in full at the place of delivery. To investigate and test the supplies as well as to notify its decision to accept or reject the delivery, the contracting authority disposes of a period of thirty days

This period will begin on the day after the date of arrival of the supplies at the place of delivery, provided that the contracting authority is in possession of the delivery note or invoice. It comprises the 30-day period stipulated in Article 120.

4.13.2 Transfer of ownership (Art. 132)

The contracting authority automatically becomes the owner of the supplies as soon as they have been accepted for payment pursuant to Article 127 of GIR.

4.13.3 Guarantee period (Art. 134)

The warranty period commences on the date on which provisional acceptance is given. It lasts for 365 calendar days.

4.13.4 Final acceptance (Art. 135)

Final acceptance occurs upon expiry of the warranty period. It is implicit when the delivery has not led to any claims during said period.

If delivery has led to complaints during the warranty period, a final acceptance or refusal of acceptance report will be issued within 15 days prior to the expiry of said period.

4.14 Invoicing and payment of services (Art. 66 to 72 and 127)

The contractor sends (one copy only of) the invoices and the contract acceptance report (original copy) to the following address:

Mr. Omona Francis

francis.omona@enabel.be

Financial controller- We care project

Kakiza road, plot No. 9 Booma, Fort portal City

Only delivery that has been performed correctly may be invoiced.

The contracting authority disposes of a period for verification of thirty days starting on the end date of the delivery, set in conformity with the modalities in the procurement documents, to carry out the technical acceptance and provisional acceptance formalities and to notify the result to the supplier.

The amount owed to the supplier must be paid within thirty days with effect from the expiry of the verification term or with effect from the day after the last day of the verification term, if this is less than thirty days. And provided that the contracting authority possesses, at the same time, the duly established invoice and any other documents that may be required.

When the procurement documents do not provide for any separate debt claim, the invoice will constitute the debt claim.
The invoice must be in **EUROS.**
Payment corresponding to 100% shall be made after delivery and acceptance of equipment at each of the health facilities

Advance payment:

By way of derogation from the foregoing, and in accordance with Articles 12/1 to 5 of the Law of 17 June 2016, inserted by the Law of 22 December 2023 amending the regulations relating to public contracts with a view to promoting access by SMEs to the said contracts, the contracting authority shall pay an advance when the successful tenderer proves to be an SME within the meaning of Article 163, § 3, subparagraph 2, of the Law of 17 June 2016.
The amount of the advance payment is calculated by applying the following percentages to a reference value determined in accordance with Article 12/5 of the Law of 17 June 2016:
1° if the successful tenderer is a micro-enterprise, i.e. an enterprise that employs fewer than ten (10) people and whose annual turnover or annual balance sheet total does not exceed two million euros (2M euro), the percentage to be taken into account is twenty per cent (20%);
2° if the successful tenderer is a small business, i.e. a business that employs fewer than fifty (50) people and whose annual turnover or annual balance sheet total does not exceed ten million euros (10M euro), the percentage to be taken into account is ten per cent (10%);
3° where the successful tenderer is a medium-sized company, i.e. a company employing fewer than two hundred and fifty (250) people and whose annual turnover does not exceed fifty million euros (50M euro) or whose annual balance sheet total does not exceed forty-three million euros (43M euro), the percentage to be taken into account is five per cent (5%).
According to Article 12/5 of the Law of 17 June 2016, the reference value relevant for calculating the advance in a framework agreement is equal to the amount of each order, including all taxes.
The first half of the advance shall be set off against the sums due to the contractor when the value of the services performed reaches thirty per cent of the original order amount and the

second half of the advance shall be set off against the sums due to the contractor when the value of the services performed reaches sixty per cent of the original order amount. The aforementioned amounts shall be understood as amounts inclusive of value-added tax.

The supplier must provide an **advance bank guarantee** prior to any advance payment.

The amount of the advance will be deducted from the final invoice of each order.

No advance will be paid when implementation duration of an order is less than 60 days.

4.15 Litigation (Art. 73)

The competent courts of Brussels have exclusive jurisdiction over any dispute arising from the performance of this public contract. French or Dutch are the languages of proceedings.

The contracting authority will in no case be held liable for any damage caused to persons or property as a direct or indirect consequence of the activities required for the performance of this contract. The contractor indemnifies the contracting authority against any claims for compensation by third parties in this respect.

In case of 'litigation', i.e. court action, correspondence must (also) be sent to the following address:

Belgian development agency - Enabel

Legal unit of the Logistics and Acquisitions service (L&A)

To the attention of Ms Inge Janssens

rue Haute 147

1000 Brussels

Belgium

4.16 Obligations of the contracting authority (Art. 136)

- The contracting authority shall:
- 1° use the goods delivered for the needs stipulated under the public contract and in accordance with technical user guidance provided by the supplier;
- 2° make not changes to the goods delivered without the written preliminary approval of the supplier.

4.17 Obligations of the supplier (Art. 137 and 138)

- The supplier shall:

- 1° put the supplies at the disposal of the contracting authority within the deadline set in the procurement documents;
- 2° unless otherwise stipulated in the procurement documents, ensure their maintenance and make all necessary repairs within the timing imposed to keep the goods in good state during the public contract term.
- Where the supplies are completely or partially destroyed during the contact term without the contracting authority being liable, the supplier shall replace these or repair them at his costs within the deadline set.

5 Procurement Procedure

5.1 Type of procedure

This contract is awarded in accordance with Article 36 of the Law of 17 June 2016 via an Open Procedure.

5.2 Publication

Official notification

This contract is officially advertised in the Belgian Public Tender bulletin and in the Official Journal of the European Union.

Further publication

This Tender Specifications are posted on the Enabel website

https://www.enabel.be/public-procurement/

5.3 Information

The awarding of this procurement contract is coordinated by the Contract Service Centre of Enabel in Uganda. Throughout this procedure all contacts between the contracting authority and the (prospective) tenderers about this procurement contract will exclusively pass through this service. (Prospective) tenderers are prohibited to contact the contracting authority in any other way with regards to this contract, unless otherwise stipulated in these Tender Specifications.

Until 10 calendar days before the time for the receipt of tenders, candidate-tenderers may ask questions about these Tender Specifications and the procurement contract. Questions will be in writing to UGA_CSC_CONTRACTS@enabel.be with copy to sandra.adero@enabel.be with a clear indication in the subject of the e-mail of the procedure reference and the contract title, as stated on the cover page of the tender specifications. They will be answered in the order received. The complete overview of questions asked will be available at the address mentioned above as soon as available.

Until the notification of the award decision no information will be given about the evolution of the procedure.

The contracting Authority shall organize **an optional information (pre-bid) meeting** at the time and location specified below.

Public contract for the supply of supply and delivery of medical equipment for health facilities	**27th August, 2024 at 11:00 am Kampala time** at Enabel Representation Office Plot 1B Lower Kololo Terrace **+ Online Meeting** Microsoft teams meeting Meeting ID: **371 002 510 826** Passcode: **cDGbSB**

The tenderer is supposed to submit his tender after reading and taking into account any corrections made to the contract notice or the Tender Specifications that are published in the Belgian Public Tender bulletin or that are sent to him by e-mail. To do so, when the tenderer has downloaded the Tender Specifications, it is strongly advised that he gives his coordinates to the public procurement administrator mentioned above and requests information on any modifications or additional information.

In accordance with Article 81 of the Royal Decree of 18 April 2017, the tenderer is required to report immediately any gap, error or omission in the procurement documents that precludes him from establishing his price or compare tenders, within ten days at the latest before the deadline for receipt of tenders.

5.4 Preparation and submission of tenders

5.4.1 Preparation of tenders

The tenderer shall prepare separately, the administrative, technical and financial proposals as explained below;

5.4.1.1 Content of tenders

The tenderer must use the tender form in annexe. In case he does not use this form, he is fully responsible for the perfect concordance between the documents he has used and the form.

The tender and the annexes to the tender form are drawn up in English.

By submitting a tender, the tenderer automatically renounces to his own general or specific sales conditions, even if these are mentioned in any of the annexes to his tender.
The tenderer clearly designates in his tender which information is confidential and/or relates to technical or business secrets and may therefore not be divulged by the contracting authority.
The tender shall contain the following parts:

1. Administrative Proposal

The tenderer shall use the tender forms included in the corresponding section of the Annex.
The Administrative proposal shall respect the following structure:

- Identity form
- Legal identification form
- Financial Identification Form **(along with an account confirmation letter from the bank. This account shall not change throughout the contract duration and implementation)**
- Subcontractor form
- Exclusion Criteria Form
- Integrity form
- Technical capacity form
- Financial capacity form
- European Single Procurement Document (ESPD)

The successful tenderer shall be required to provide the following documents before award

- Articles of Association
- Tax Clearance Certificate (e.g; URA, as applicable)
- Social Security Contribution Clearance (e.g. NSFF as applicable)
- An extract from the criminal record in the name of the tenderer (legal person) or his representative (natural person) if there is no criminal record for legal persons (ex. certificate of good conduct from Interpol);

2. Technical Proposal

The technical proposal may be presented in the following format:
The technical proposal shall be presented in a free format but it is mandatory to include the original product brochure for all the medical equipment. These brochures must clearly mention

the brand name, model and technical specification in a way that allows comparability of the offers. The contractor shall provide Manufacturers' authorization for all the equipment.

3. Financial Proposal

The tenderer shall use the tender forms included in the corresponding section of the Annex.

Determination of prices

All prices given in the tender form must obligatorily be quoted in EUROS.

This procurement contract is a lump sum contract, meaning a contract in which a flat rate price covers the whole performance of the contract or each of the items of the inventory.

In accordance with Article 37 of the Royal Decree of 18 April 2017, the contracting authority may for the purpose of verifying the prices carry out an audit of any and all accounting documents and an on-site audit to check the correctness of the indications supplied.

Elements included in the price

The tenderer is to include in his unit and global prices any charges and taxes generally inherent to the performance of the contract, with the exception of the value-added tax.

The following are in particular included in the prices:

1° packaging (except if these remain the property of the tenderer), loading, trans-shipment and intermediate unloading, transportation, insurance and customs clearance;

2° unloading, unpacking and deployment at the place of delivery, provided that the procurement documents state the exact place of delivery and the means of access;

3° documentation pertaining to the delivery of supplies and any documentation required by the contracting authority;

4° assembly and taking into operation;

5° training required for operation;

6° where applicable, the measures imposed by occupational safety and worker health legislation;

87° customs and excise duties;

All prices are DDP (INCOTERMS 2020)

5.4.1.2 Validity of tenders

Tenders shall remain valid for 120 calendar days from the final date for receiving tenders.

5.4.2 Submission of tenders

Without prejudice to any variants, the tenderer may only submit one tender per lot.

The tenderer submits his tender as follows:

The tenderer shall submit separately (in separate envelopes), the administrative, technical and financial proposals. The sealed envelopes containing the different proposals shall then be put together and sealed in one big envelope to be submitted to the contracting authority.

One original copy of the completed tender shall be submitted on paper (hard copy). Electronic copies shall be submitted in one or more PDF files on a USB stick. the USB stick shall be inserted into the envelop containing the hard copy tender.

The tender submitted in a properly sealed envelope bearing the following information: Name of tenderer, the title of the contract and the reference number of the procurement, as stated on the cover page of the tender specifications.

It shall be submitted:

a) By mail (standard mail or registered mail)

In this case, the sealed envelope is put in a second closed envelope addressed to:

Enabel Uganda

Contract Service Center

Lower Kololo Terrace, Plot 1B

PO Box 40131 Kampala – Uganda

OR

b) Delivered by hand with acknowledgement of receipt.

The service can be reached on working days during office hours: from 9:00 am to 12:00 pm and from 2:00 pm to 4:00 pm (see the address given under point a) above).

The tender shall be received by the Contracting Authority before **23rd September, 2024, 11:00 am, Kampala time.** Tenders that arrive late will not be accepted. (Article 83 of the Royal Decree on Awarding)

5.4.3 Modification or withdrawal of a tender that has already been submitted

When a tenderer wants to change or withdraw a tender already sent or submitted this must be done in accordance with the provisions of Articles 43 and 85 of the Royal Decree of 18 April 2017.

To change or withdraw a tender already sent or submitted a written statement is required, which will be correctly signed by the tenderer or his representative. The subject-matter and the scope of the changes must be indicated in detail. Any withdrawal must be unconditional.

The withdrawal may also be communicated by fax or electronic means, provided that it is confirmed by registered letter deposited at the post office or against acknowledgement of receipt at the latest the day before the tender acceptance deadline.

The subject-matter and the scope of the changes must be indicated in detail.

The withdrawal must be pure and simple.

5.5 Opening and evaluation of Tenders

5.5.1 Opening of tenderers

The opening of tenders will take place on the same day of the final date for receiving tenders indicated above. Tenders not received before 11:00 am will be rejected. The opening shall be a public opening at 11:30 am Kampala time at the address below.

Enabel Uganda

Lower Kololo Terrace, Plot 1B

PO Box 40131 Kampala – Uganda

5.5.2 Evaluation of Tenders

5.5.2.1 Selection of tenderers

Exclusion grounds

The mandatory and optional exclusion grounds are given in the Declaration on Honour enclosed to these Tender Specifications.

By submitting his tender together with the completed European Single Procurement Document (ESPD) the tenderer declares officially on his honour that:

- he is not in one of the mandatory or facultative exclusion cases, which must or may lead to his exclusion;

- he fulfils the selection criteria established by the contracting authority in this public contract

The European Single Procurement Document (ESPD) is a self-declaration by economic operators providing preliminary evidence replacing the certificates issued by public authorities or third parties. As provided in Article 73 of the Law of 17 June 2016, it is a formal statement by the economic operator that it is not in one of the situations in which economic operators shall or may be excluded; that it meets the relevant selection criteria.

The tenderer can either complete the ESDP given in attachment, or generate his document via the website: https://ec.europa.eu/tools/espd/filter

Where the tender is submitted by a group of economic operators, it must include an ESPD for each of the participants in the group.

In accordance with Article 38 §2 of Article 73 of the Royal Decree of 18 April 2017, regarding part IV of the ESPD on the selection criteria, the contracting authority has decided to limit the information to be filled out to one single question, namely whether the economic operator fulfils the required selection criteria, in accordance with the section "Global indication for all selections criteria". So, only this section must be completed.

The contracting authority will verify the accuracy of this Declaration on honour Based on the supporting documents.

Conflicts of interest - Revolving doors (Art. 51 Royal Decree 18/04/2017).

Without prejudice to Articles 6 and 69, paragraph 1, 5° of the Law, a conflict of interest is also considered any (‘revolving doors’) situation in which a natural person who has worked for a contracting authority as an internal staff member, whether in a hierarchy relation or not, as a concerned civil servant, public officer or any other person linked whatsoever to the contracting authority, would later intervene under a public contract awarded by this contracting authority and where a relation exists between the former activities that the above person conducted for the contracting authority and the activities he or she conducts under the contract.

The application of above-mentioned provision is limited however to a two-year term from the resignation of said person or any other type of termination of the former activities.

Selection criteria

Moreover, by means of the documents requested in the Annexes - Administrative Proposal, the tenderer must prove that he is sufficiently capable, from an economic and financial as well as from a technical point of view, to successfully perform this public procurement contract.

Only tenders from tenderers who meet the selection criteria are taken into consideration in order to participate in the comparison of tenders on the basis of the award criteria set out below, subject to the regularity of these tenders.

1	**Sufficient Economic and Financial Capacity**
1.1	Sufficient turn-over
Minimum Standard	Lot 1: Minimum average annual turnover of 165,000 Euros during the past three financial years Lot 2: Minimum average annual turnover of 110,000 Euros during the past three financial years Lot 3: Minimum average annual turnover of 64,000 Euros during the past three financial years (If a contractor submits for more than one lot, the amount above shall be summed up for the lots tendered.)
2	**Sufficient Technical and Professional Capacity**
2.1	Sufficient experience in supply and delivery of medical equipment
Minimum Standard	For each lot, minimum of 2 assignments within the scope of the lot, which was totally and successfully completed in the last 3 years, each of the assignments shall be at least Lot 1: 120,000 Euros Lot 2: 80,000 Euros Lot 3: 50,000 Euros
	Manufacturer's authorization for all the equipment
	Proof of manufacturer's training of at least one technical person for anaesthesia machine (MUST) and or ultra sound machine (training certificate to be provided)

A tenderer may, where appropriate and for a particular contract, rely on the capacities of other entities, regardless of the legal nature of the links which he has with these entities. In that case, the following rules apply:

- Where an economic operator wants to rely on the capacities of other entities, it shall prove to the contracting authority that it will have at its disposal the resources necessary, for example, by producing a commitment by those entities to that effect.
- The contracting authority shall verify whether the entities on whose capacity the economic operator intends to rely fulfil the relevant selection criteria and whether there are grounds for exclusion.
- Where an economic operator relies on the capacities of other entities with regard to criteria relating to economic and financial standing, the contracting authority may require that the economic operator and those entities be jointly liable for the execution of the contract.
- The contracting authority may require certain essential tasks to be carried out directly by the tenderer himself or, if the tender is submitted by a group of economic operators, by a member of the said group.

Under the same conditions, a group of candidates or tenderers may submit the capacities of the group's participants or of other entities.

Where a candidate or tenderer relies on the capacity of other entities (particularly subcontractors or independent subsidiaries) for economic and financial capacity criteria and technical and vocational capacity criteria, the candidate or tenderer, as appropriate, answers the question in part II, C, of the ESPD. He also mentions for which part of the public contract he will rely on such capacity and which other entities he proposes.

The tender also comprises a separate ESPD for these entities.

Regularity of tenders

The tenders submitted by the selected tenderers will be evaluated as to formal and material regularity.

The tenders must be drawn up in such a way that the contracting authority can make a selection without starting negotiations with the tenderer. For this reason, and in order to be able to assess the tenders fairly, it is essential that the tenders be completely in conformity with the provisions of the Tender Specifications, both formally and materially.

The substantially irregular tenders are rejected.

A substantial irregularity is such as to give a discriminatory advantage to the tenderer, to distort competition, to prevent the evaluation of the tenderer's tender or its comparison with the other tenders, or to render non-existent, incomplete or uncertain the commitment of the tenderer to perform the contract under the conditions laid down.

The following irregularities are deemed substantial:

1° if applicable, failure to comply with environmental, social or labour law, provided that such non-compliance is punishable by law;

2° failure to comply with the requirements of Articles 38, 42, 43, §1, 44, 48, §2, clause 1, 1alinéa 1er, 54, §2, 55, 83 and 92 of the Royal Decree of 18 April 2017 and of Article 14 of the Law, insofar as they contain obligations vis-à-vis the tenderers;

3° failure to comply with the minimum requirements and the requirements that are indicated in the technical specifications;

4° tenders that do not bear an original handwritten signature on the tender form.

The contracting authority will also declare void any tender that is affected by several non-substantial irregularities which, by reason of their accumulation or combination, are capable of having the same effect as described above (in accordance with Article 76 of the Royal Decree of 18 April 2017).

5.5.2.2 Financial evaluation of tenders

Award Criteria

The contracting authority selects the regular tender that it finds to be the least expensive, taking account of the following criteria:

- Price: 100 %;

With regards to the 'price' criterion, the following formula will be used:

Points tender A = amount of lowest tender * 100

amount of tender A

Final score

For each lot, the procurement contract will be awarded to the tenderer with the highest final score, after the contracting authority has verified the accuracy of the Declaration on honour of

this tenderer and provided the control shows that the Declaration on honour corresponds with reality.

5.6 Award and Conclusion of Contract

5.6.1 Awarding the contract

For each lot, the contract will be awarded to the tenderer who has submitted the least expensive tender.

It is to be noted that in accordance with Art. 85 of the Law of 17 June 2016, there is no obligation for the contracting authority to award the procurement contract.

The contracting authority may either decide not to award the procurement contract; either redo the procedure, if necessary through another award procedure.

The contracting authority maintains the right to award only a certain lot or certain lots.

5.6.2 Concluding the contract

In accordance with Art. 88 of the Royal Decree of 18 April 2017, the procurement contract occurs through the notification to the selected tenderer of the approval of his tender.

Notification is via e-mail.

So, the full contract agreement consists of a procurement contract awarded by Enabel to the chosen tenderer in accordance with:

- These Tender Specifications and its annexes;
- The registered letter of notification of the award decision;
- Any later documents that are accepted and signed by both parties, as appropriate.

In an objective of transparency, Enabel undertakes to publish each year a list of recipients of its contracts. By introducing his tender, the successful tenderer declares that he agrees with the publication of the title of the contract, the nature and object of the contract, its name and location, and the amount of the contract.

6 Annexes

6.1 Contractual Documents

Model Performance Bond

Only for the successful tenderer:

Bank X

Address

Performance bond n° X

This performance bond is posted in the context of the Law of 17 June 2016 on public contracts and on certain works, supply and service contracts and in conformity with the General Implementing Rules (GIR) provided in the Royal Decree of 14 January 2013 establishing the general implementing rules of public contracts and the award of public works.

X, address (the "Bank")

hereby declares posting security for a maximum amount of X € (X euros) for the Belgian Development Agency (Enabel) for the obligations of X, address for the contract:

"X, tender documents Enabel < UGAX, lot X" (the "Contract").

Consequently, the Bank commits, under condition of the beneficiary waiving any right to contest or divide liability, to pay up to the maximum amount, any amount which X may owe to Enabel in case X defaults on the performance of the "Contract".

This performance bond shall be released in accordance with the provisions of the tender documents Enabel < UGAX and of Art. 25-33 of the Royal Decree of 22 June 2017, and at the latest at the expiry of 18 months after the provisional acceptance of the Contract.

Any appeal made to this performance bond must be addressed by registered mail to the Bank X, address, with mention of the reference of the procurement procedure.

Any payment made from this performance bond will ipso jure reduce the amount secured by the Bank.The performance bond is governed by the Belgian Law and only Belgian courts are competent in case of litigation.

Done in X on X Signature: Name:

6.2 Procedural Documents – Tender Forms

6.2.1 ADMINISTRATIVE PROPOSAL

Legal Identification forms

I. PERSONAL DATA **FAMILY NAME(S)**① **FIRST NAME(S)**① **DATE OF BIRTH** **JJ MM YYYY** **PLACE OF BIRTH** **COUNTRY OF BIRTH** **(CITY, VILLAGE)** **TYPE OF IDENTITY DOCUMENT** **IDENTITY CARD** **PASSPORT** **DRIVING LICENCE**② **OTHER**③ **ISSUING COUNTRY** **IDENTITY DOCUMENT NUMBER** **PERSONAL IDENTIFICATION NUMBER**④ **PERMANENT** **PRIVATE ADRESS** **POSTCODE** **P.O. BOX** **CITY** **REGION** ⑤ **COUNTRY** **PRIVATE PHONE** **PRIVATE E-MAIL**

II. BUSINESS DATA	If YES, please provide business data and attach copies of official supporting documents
Do you run your own business without a separate legal personality (e.g. sole traders, self-employed etc.) **YES** **NO**	**BUSINESS NAME (if applicable)** **VAT NUMBER** **REGISTRATION NUMBER** **PLACE OF REGISTRATION** **CITY** **COUNTRY**

DATE	**SIGNATURE**

① As indicated on the official document.

② Accepted only for Great Britain, Ireland, Denmark, Sweden, Finland, Norway, Iceland, Canada, United States and Australia.

③ Failing other identity documents: residence permit or diplomatic passport.

④ See table with corresponding denominations by country. ⑤ To be completed with Region, State or Province by non EU countries only, excluding EFTA and candidate countries.

Legal person entity private/public legal body

OFFICIAL NAME ②			
ABREVIATION			
MAIN REGISTRATION NUMBER③			
SECONDARY REGISTRATION NUMBER (if applicable)			
PLACE OF MAIN REGISTRATION	**CITY**		**COUNTRY**
DATE OF MAIN REGISTRATION		**DD MM YYYY**	
VAT NUMBER			
OFFICIAL ADDRESS			
POSTCODE	**P.O. BOX**		**CITY**
COUNTRY			**PHONE**
E-MAIL			

DATE	**STAMP**
SIGNATURE OF AUTHORISED REPRESENTATIVE	

① **Public law body WITH LEGAL PERSONALITY, meaning a public entity being able to represent itself and act in its own name, i.e. being capable of suing or being sued, acquiring and disposing of property, entering into contracts. This legal status is confirmed by the official legal act establishing the entity (a law, a decree, etc.).**

② **National denomination and its translation in EN or FR if existing.**

③ **Registration number in the national register of the entity.**

Public law entity

OFFICIAL NAME①			
BUSINESS NAME			
(if different)			
ABREVIATION			
LEGAL FORM			
ORGANISATION TYPE	**FOR PROFIT**		
	NOT FOR PROFIT	**NGO② YES**	**NO**
MAIN REGISTRATION NUMBER③			
SECONDARY REGISTRATION NUMBER			
(if applicable)			
PLACE OF MAIN REGISTRATION	**CITY**		**COUNTRY**
DATE OF MAIN REGISTRATION			
	DD MM YYYY		
VAT NUMBER			
ADDRESS OF			
HEAD OFFICE			
POSTCODE	**P.O. BOX**		**CITY**
COUNTRY			**PHONE**
E-MAIL			

DATE	**STAMP**
SIGNATURE OF AUTHORISED REPRESENTATIVE	

① National denomination and its translation in EN or FR if existing.

② NGO = Non Governmental Organisation, to be completed if NFPO is indicated.

③ Registration number in the national register of companies. See table with corresponding field denomination by country.

Financial identification form

BANKING DETAILS		
ACCOUNT NAME [9]		
IBAN/ACCOUNT NUMBER[10]		
CURRENCY		
BIC/SWIFT CODE		
BANK NAME		
ADDRESS Of BANK BRANCH		
STREET & NUMBER		
TOWN/CITY	POST CODE	
COUNTRY		

ACCOUNT HOLDER'S DATA (AS DECLARED TO THE BANK)		
ACCOUNT HOLDER		
STREET & NUMBER		
TOWN/CITY	POST CODE	
COUNTRY		

SIGNATURE OF ACCOUNT HOLDER (Obligatory) NAME: TITLE:	DATE (Obligatory)

9 This does not refer to the type of account. The account name is usually the one of the account holder. However, the account holder may have chosen a different name to its bank account.

10 Qa1Fill in the IBAN Code (International Bank Account Number) if it exists in the country where your bank is established.

Subcontractors

Name and legal form	Address / Registered office	Object

Declaration on honour – exclusion criteria

Hereby, I / we, acting as legal representative(s) of above-mentioned tenderer, declare that the tenderer does not find himself in one of the following situations :

1) The tenderer or one of its 'directors[1]' was found guilty following a conviction by final judgement for one of the following offences:

1° involvement in a criminal organisation

2° corruption

3° fraud

4° terrorist offences, offences linked related to terrorist activities or incitement to commit such offence, collusion or attempt to commit such an offence

5° money laundering or terrorist financing

6° child labour and other trafficking in human beings

7° employment of foreign citizens under illegal status

8° creating a shell company.

2) The counterparty which fails to fulfil his obligations relating to the payment of taxes or social security contributions for an amount in excess of EUR 3 000, except if the counterparty can demonstrate that a contracting authority owes him one or more unquestionable and due debts which are free of all foreseeable liabilities. These debts are at least of an amount equal to the one for which he is late in paying outstanding tax or social charges.

3) The counterparty who is in a state of bankruptcy, liquidation, cessation of activities, judicial reorganisation or has admitted bankruptcy or is the subject of a liquidation procedure or judicial reorganisation, or in any similar situation resulting from a procedure of the same kind existing under other national regulations;

4) When Enabel can demonstrate by any appropriate means that the counterparty or any of its directors has committed serious professional misconduct which calls into question his integrity.

Are also considered such serious professional misconduct:

a. A breach of Enabel's Policy regarding sexual exploitation and abuse – June 2019

b. A breach of Enabel's Policy regarding fraud and corruption risk management – June 2019

c. A breach of a regulatory provision in applicable local legislation regarding sexual harassment in the workplace

d. The counterparty was seriously guilty of misrepresentation or false documents when providing the information required for verification of the absence of grounds for exclusion or the satisfaction of the selection criteria, or concealed this information

e. Where Enabel has sufficient plausible evidence to conclude that the counterparty has committed acts, entered into agreements or entered into arrangements to distort competition

The presence of this counterparty on one of Enabel's exclusion lists as a result of such an act/agreement/arrangement is considered to be sufficiently plausible an element.

5) When a conflict of interest cannot be remedied by other, less intrusive measures;

6) When significant or persistent failures by the counterparty were detected during the execution of an essential obligation incumbent on him in the framework of a previous contract, a previous contract placed with another contracting authority, when these failures have given rise to measures as of right, damages or another comparable sanction.

Also failures to respect applicable obligations regarding environmental, social and labour rights, national law, labour agreements or international provisions on environmental, social and labour rights are considered 'significant'.

The presence of the counterparty on the exclusion list of Enabel because of such a failure serves as evidence.

7) Restrictive measures have been taken vis-à-vis the counterparty with a view of ending violations of international peace and security such as terrorism, humanrights violations, the destabilisation of sovereign states and de proliferation of weapons of mass destruction.

The counterparty or one of its directors are on the lists of persons, groups or entities submitted by the United Nations, the European Union and Belgium for financial sanctions:

For the United Nations, the lists can be consulted at the following address:

https://finances.belgium.be/fr/tresorerie/sanctions-financieres/sanctionsinternationales-nations-unies

For the European Union, the lists can be consulted at the following address: https://finances.belgium.be/fr/tresorerie/sanctions-financieres/sanctionseurop%C3%A9ennes-ue https://eeas.europa.eu/headquarters/headquarters-homepage/8442/consolidatedlist-sanctions_en https://eeas.europa.eu/sites/eeas/files/restrictive_measures-2017-01-17-clean.pdf For Belgium: https://finances.belgium.be/fr/sur_le_spf/structure_et_services/administrations_generales/tr%C3%A9sorerie/contr%C3%B4le-des-instruments-1-2

8) If Enabel executes a project for another funder or donor, other grounds for exclusion may be added.

Signature preceded by 'read and approved', in writing, and indication of name and function of the person signing: ……..

Place, date

Integrity statement for the tenderers

Hereby, I / we, acting as legal representative(s) of above-mentioned tenderer, declare the following:

- Neither members of administration or employees, or any person or legal person with whom the tenderer has concluded an agreement in view of performing the public contract, may obtain or accept from a third party, for themselves of for any other person or legal person, an advantage appreciable in cash (for instance, gifts, bonuses or any other kind of benefits), directly or indirectly related to the activities of the person concerned for the account of Enabel.
- The board members, staff members or their partners have no financial or other interests in the businesses, organisations, etc. that have a direct or indirect link with Enabel (which could, for instance, bring about a conflict of interests).
- I have / we have read and understood the articles about deontology and anticorruption included in the Tender Documents (see 1.7.), as well as *Enabel's Policy regarding sexual exploitation and abuse* of June 2019 and *Enabel's Policy regarding fraud and corruption risk management* of June 2019 and I / we declare fully endorsing and respecting these articles.

If above-mentioned public contract is awarded to the tenderer, I/we declare, moreover, agreeing with the following provisions:

- In order to avoid any impression of risk of partiality or connivance in the follow-up and control of the performance of the public contract, it is strictly forbidden to the public contractor (i.e. members of the administration and workers) to offer, directly or indirectly, gifts, meals or any other material or immaterial advantage, of whatever value, to the employees of Enabel who are concerned, directly or indirectly, by the follow-up and/or control of the performance of the public contract, regardless of their hierarchical rank.
- Any (public) contract will be terminated, once it appears that contract awarding or contract performance would have involved the obtaining or the offering of the abovementioned advantages appreciable in cash.
- Any failure to comply with one or more of the deontological clauses will be considered as a serious professional misconduct which will lead to the exclusion of the contractor from this and other public contracts for Enabel.

- The public contractor commits to supply, upon the demand of the contracting authority, any supporting documents related to the performance conditions of the contract. The contracting authority will be allowed to proceed to any control, on paperwork or on site, which it considers necessary to collect evidence to support the presumption of unusual commercial expenditure.

Finally, the tenderer takes cognisance of the fact that Enabel reserves the right to lodge a complaint with the competent legal instances for all facts going against this statement and that all administrative and other costs resulting are borne by the tenderer.

Signature preceded by 'read and approved', in writing, and indication of name and function of the person signing: ……..

Place, date

Economic and financial capacity Form

Financial Statement

The tenderer must complete the following table of financial data based on his/her annual accounts.

Financial data	**Year- 3** **€ or NC**	**Year- 2** **€ or NC**	**Year -1** **€ or NC**	**Average** **€ or NC**
Annual turnover, excluding this public contract				

The tenderer must also provide his/her approved financial statements for the last three financial years or an appropriate supporting document, such as a document listing all assets and liabilities of the enterprise. In case the enterprise has not yet published its Financial Statements, an interim balance certified true by an accountant or by a registered auditor or by the person or body with this function in the country concerned will do

Technical and professional capacity form

List of main similar assignments

Lot 1: Supply, delivery, installation, testing and user training of Cemonc medical equipment.

Description of the main similar assignments totally performed	Location	Amount involved	Completion date in the last 3 years (only totally performed assignments)	Name of the public or private bodies

Certificates of completion

For each of the assignments listed, the tenderer must provide in the administrative proposal as annexes to this form the certificates of completion/acceptance (statement or certificate without major reservation) and / or any supporting documents (contracts, invoices…) approved by the entity which awarded the contract.

Lot 2: Supply, delivery, installation, testing and user training of NICU medical equipment.

Description of the main similar assignments totally performed	Location	Amount involved	Completion date in the last 3 years (only totally performed assignments)	Name of the public or private bodies

Certificates of completion

For each of the assignments listed, the tenderer must provide in the administrative proposal as annexes to this form the certificates of completion/acceptance (statement or certificate without major reservation) and / or any supporting documents (contracts, invoices…) approved by the entity which awarded the contract.

Lot 3: Supply, delivery, installation, testing and user training of Emergency services medical equipment.

Description of the main similar assignments totally performed	Location	Amount involved	Completion date in the last 3 years (only totally performed assignments)	Name of the public or private bodies

Certificates of completion

For each of the assignments listed, the tenderer must provide in the administrative proposal as annexes to this form the certificates of completion/acceptance (statement or certificate without major reservation) and / or any supporting documents (contracts, invoices…) approved by the entity which awarded the contract.

6.3.2 TECHNICAL PROPOSAL

The technical proposal shall be presented in free format:

The technical proposal shall be presented in a free format but it is mandatory to include the original product brochure for all the medical equipment. These brochures must clearly mention the brand name, model and technical specification in a way that allows comparability of the offers. The contractor shall provide Manufacturers' authorization for all the equipment.

6.3.3 FINANCIAL PROPOSAL

Lot 1: Supply, delivery, installation, testing and user training of Cemonc medical equipment.

Tender Forms – prices

By submitting this tender the tenderer commits to performing this public contract in conformity with the provisions of the Tender Specifications/ – and explicitly declares accepting all conditions listed in the Tender Specifications and renounces any derogatory provisions such as his own general sales conditions.

The unit prices and the global prices for each item in the inventory are established relative to the value of these items in relation to the total value of the tender. All general and financial costs as well as the profits are distributed between the various items in proportion to their weight.

The value-addedS tax is dealt with on a separate line in the summary bill of quantities or the inventory, to be added to the tender's value.

The tenderer commits to performing the public contract in accordance with the provisions of the Tender Specifications for the following prices, given in euros and exclusive of VAT:

Should this tender be approved, the performance bond will be constituted under the conditions and deadlines stipulated in the Tender Specifications.

The confidential information and/or the information relating to technical or business secrets is indicated clearly in the tender.

In order to correctly compare the tenders, the duly signed information or documents mentioned under Preparation of Tenders.

N°	Description	Unit of measure	Quantity	Unit price in Euros excl. VAT	Lump-sum unit prices in euros excl. VAT
1.	Oxygen concentrator 10LPM	Pc	8	€	€
2.	Vacuum extractor	Pc	14	€	€
3.	Dilation and Curettage set	Set	14	€	€
4.	Portable Ultrasound scanner	Pc	1	€	€

5.	Universal Anaesthesia machine	Unit	6	€	€
6.	Autoclave electric 75Ls. (sterilizer)	Pc	5	€	€
7.	Operation table, hydraulic/ Theatre bed	Pc	5	€	€
8.	Operating light mobile LED	Pc	8	€	€
9.	Delivery beds	Pc	8	€	€
10	Wheel Chair Adult	Pc	1	€	€
11.	Patient stretcher/trolley	Pc	24	€	€
12.	The patient monitors 5 Parameters with a stand	Pc	8	€	€
13.	Foetal Doppler	Pc	20	€	€
14.	Patient bed	Pc	8	€	€
15.	Blood Refrigerator	Pc	1	€	€
16.	HemoCue	Pc	7	€	€
				Total price in Euros	
				VAT percentage (if applicable):	18%

Name and first name: ……

Duly authorised to sign this tender on behalf of: ……

Place and date: ……

Signature: ……

Lot 2: Supply, delivery, installation, testing and user training of NICU medical equipment.

Tender Forms – prices

By submitting this tender, the tenderer commits to performing this public contract in conformity with the provisions of the Tender Specifications/ – and explicitly declares accepting all conditions listed in the Tender Specifications and renounces any derogatory provisions such as his own general sales conditions.

The unit prices and the global prices for each item in the inventory are established relative to the value of these items in relation to the total value of the tender. All general and financial costs as well as the profits are distributed between the various items in proportion to their weight.

The value added tax is dealt with on a separate line in the summary bill of quantities or the inventory, to be added to the tender's value.

The tenderer commits to performing the public contract in accordance with the provisions of the Tender Specifications for the following prices, given in euros and exclusive of VAT:

Should this tender be approved, the performance bond will be constituted under the conditions and deadlines stipulated in the Tender Specifications.

The confidential information and/or the information relating to technical or business secrets is indicated clearly in the tender.

In order to correctly compare the tenders, the duly signed information or documents mentioned under Preparation of Tenders.

N°	Description	Unit of measure	Quantity	Unit price in Euros excl. VAT	Lump-sum unit prices in euros excl. VAT*
1.	Oxygen concentrator 10LPM Dual flow	Pc	13	€	€
2.	Infusion pump	Pc	2	€	€
3.	Suction machine	Pc	9	€	€
4.	Phototherapy machine	Pc	9	€	€
5.	Infant Incubator	Pc	14	€	€
6.	Infant Warmers	Pc	15	€	€
7.	CPAP	Pc	6	€	€

8.	Vein Finders	Pc	18	€	€
				Total price in Euros	
				VAT percentage (if applicable):	18%

Name and first name: ……

Duly authorised to sign this tender on behalf of: ……

Place and date: ……

Signature: ……

Lot 3: Supply, delivery, installation, testing and user training of Emergency services medical equipment.

Tender Forms – prices

By submitting this tender, the tenderer commits to performing this public contract in conformity with the provisions of the Tender Specifications/ – and explicitly declares accepting all conditions listed in the Tender Specifications and renounces any derogatory provisions such as his own general sales conditions.

The unit prices and the global prices for each item in the inventory are established relative to the value of these items in relation to the total value of the tender. All general and financial costs as well as the profits are distributed between the various items in proportion to their weight.

The value added tax is dealt with on a separate line in the summary bill of quantities or the inventory, to be added to the tender's value.

The tenderer commits to performing the public contract in accordance with the provisions of the Tender Specifications for the following prices, given in **euros and exclusive of VAT:**

Should this tender be approved, the performance bond will be constituted under the conditions and deadlines stipulated in the Tender Specifications.

The confidential information and/or the information relating to technical or business secrets is indicated clearly in the tender.

In order to correctly compare the tenders, the duly signed information or documents mentioned under Preparation of Tenders.

N°	Description	Unit of measure	Quantity	Unit price in Euros excl. VAT	Lump-sum unit prices in euros excl. VAT*
1.	Patient trolley	Pc	18	€	€
2.	Emergency Unit patient monitor 5 parameter	Pc	23	€	€
3.	Resuscitation Bed 3 function	Pc	10	€	€
4.	Emergency crash trolley	Pc	19	€	€
5.	Wheelchair Adult	Pc	27	€	€
6.	Nebulizer, Ultra sonic	Pc	20	€	€

Total price in Euros	
VAT percentage (if applicable):	18%

Name and first name: ……

Duly authorised to sign this tender on behalf of: ……

Place and date: ……

Signature: ……