

Box 3592, Dar es Salaam, Tanzania Tel. 2127597, Fax: 255-22-2118704



CONTRACTUAL AGREEMENT

TENDER NO. PA-010/2021-2022/G/33

Between

OCEAN ROAD CANCER INSTITUTE, TANZANIA

And

ANUDHA LIMITED, TANZANIA

FOR

SUPPLY OF ICU EQUIPMENTS AND CONSUMABLES FOR ORCI, MTWARA ZONAL HOSPITAL, LIGULA RRH AND MOROGORO RRH

Executive Director,
Ocean Road Cancer Institute
Barack Obama road/Luthuli Road,
P.O. Box 3592
Dar es Salaam, Tanzania

DECEMBER 2021



Form of Contract

THIS AGREEMENT made the FRIDAY 24. day of December 2021 between OCEAN ROAD CANCER INSTITUTE, junction of Barack Obama road/ Luthuli road P.O Box 3592 Dar es Salaam of Tanzania (hereinafter called "the PE") of the one part and ANUDHA LIMITED of P. o. Box 5982. Plot 2169/82& 8170/82, Morogoro Road, Opp. DART Kisutu Bus stand, DAR ES SALAAM TANZANIA (hereinafter called "the Supplier") of the other part:

WHEREAS the PE invited Tenders for certain goods and ancillary services, viz SUPPLY OF ICU EQUIPMENTS AND CONSUMABLES FOR ORCI, MTWARA ZONAL HOSPITAL, LIGULA RRH AND MOROGORO RRH and has accepted a Tender by the Supplier for the supply of those goods and services in the sum of Tanzania shillings Three Billion Seven Hundred Sixty Seven Million, One Hundred Fifty Three Thousand and Five Hundred Only. TSH 3,767,153,500.00 (VAT inclusive) (hereinafter called "the Contract Price").

NOW THIS AGREEMENT WITNESSETH AS FOLLOWS:

- 1. In this Agreement words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to.
- 2. The following documents shall be deemed to form and be read and construed as part of this Agreement, In the event of any ambiguity or conflict between the Contract Documents listed below, the order of precedence shall be the order in which the Contract Documents are listed below:-
 - (a) This form of agreement;
 - (b) the Form of Tender and the Price Schedule submitted by the Tenderer;
 - (c) the Schedule of Requirements;
 - (d) the Technical Specifications;
 - (e) the Special Conditions of Contract;



- (f) the GCC;
- (g) the Purchaser's Letter of Acceptance; and
- (h) [add here: any other documents]
- 3. In consideration of the payments to be made by the PE to the Supplier as hereinafter mentioned, the Supplier hereby covenants with the PE to provide the goods and services and to remedy defects therein in conformity in all respects with the provisions of the Contract
- 4. The PE hereby covenants to pay the Supplier Tanzania shillings Three Billion Seven Hundred Sixty Seven Million, One Hundred Fifty Three Thousand and Five Hundred Only. TSH 3,767,153,500.00 (VAT inclusive), in consideration of the provision of the goods and services and the remedying of defects therein, the Contract Price sum as may become payable under the provisions of the contract at the times and in the manner prescribed by the contract.

IN WITNESS whereof the parties hereto have caused this Agreement to be executed in accordance with their respective laws the day and year first above written.

Signed by, for and on behalf of the Employer: *OCEAN ROAD CANCER INSTITUTE* DAR ES SALAAM TANZANIA

NAME DR JULIUS MWASELAGE

DESIGNATION: ERECUTIVE DIRECTOR

SIGNATURE: JUMAN SIGNATURE:

IN PRESENCE OF:

NAME ELIPENDO KOZIMOTO

DESIGNATION: HEAD OF LEURL SERVICES

CANCER TANS

EXECUTIVE DIRECTOR

24 DEC 2021

SIGNATURE: Primoto
Signed by, for and on behalf of the Supplier: ANUDHA LIMITED P.O BOX 5982 DAR ES SALAAM
NAME: Anurag Hassija
DESIGNATION: Director
SIGNATURE: ANUDHA LIMITED P.O.Box 5082 DAR-E8-SALAAM TEL: 2122745, 2122747 FAX: 2126490
IN PRESENCE OF: NAME: NIShuta Patel
DESIGNATION: Manages
SIGNATURE: ASPatel

Col

THE FORM OF TENDER AND THE PRICE SCHEDULE SUBMITTED BY THE TENDERER



THE SCHEDULE OF REQUIREMENTS





anudha limited

QUOTATION SUBMISSION FORM

DATE: 24th November, 2021

PURCHASER: OCEAN ROAD CARDIAC INSTITUTE TENDER NO. PA-010/2021-22/G/33

To, Executive Director Ocean Road Cancer Institute P.O Box 3592 Dar Es Salaam.

Dear Sir or Madam:

We agree to supply the goods specified in the Schedule of Requirement and prices of the in accordance with the Conditions of Contract accompanying this Bid for the Contract Price Tshs 3,767,153,500/-, Three Billion, Seven Hundred Sixty Seven Million, One Hundred, Fifty Three Thousand and Five Hundred Only in Tanzanian Shillings

We also offer to deliver the said goods within the period of 1 to 16 weeks after the LPO as specified in the schedule of requirement.

This quotation and your written acceptance of it shall constitute a binding Contract between us. We understand that you are not bound to accept the lowest or any quotation you receive.

We agree to abide by this Tender for the Tender Validity Period specified in ITT 6, and it shall remain binding upon us and may be accepted at any time before the expiration of that period.

We are not participating, as Tenderers, in more than one Tender in this tendering process, other than alternative offers in accordance with the Tendering Documents.

We declares that our quoted price did not involve agreement with other tenderers for the purpose of tender suppression.

We hereby confirm that this quotation complies with the conditions required by the invitation for quotations.

Dated on 23rd day of November, 2021

In the capacity of **DIRECTOR** Name: ANURAG HASSIJA

Signed:

Duly authorized to sign Cender for and on behalf of Anacha Limited

ANUDHA LTD. P. O. Box 5982 DAR-ES-SALAAM TEL: 2122745, 2122747

P. O. BOX 5982, DAR ES SALAAM, TANZANIA Plot 2169/82 & 8170/82, Morogoro Road, Opp. DART-KISUTU Bus stand Tel: +255 22 2125746/ 2121188/ 2122745/ 2122747/ 2122746; Fax: +255 22 2126490; Cell: +255 783 523 777 Email: sales@anudha.com, sales@anudha.com, anudha@ctvsatcom.net, service@anudha.com

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35	35 Hemophomometer Model: Hemocentrol	FKF - Germans	1 to 16 works	ei	00000051			•	•		3,000,000
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3	Examination Serventing Model F-35	Heber - Chine	1 to 16 weeks	0	450,000					,	
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7	7 UV Lamp		1 to 16 weeks	2	-0			•			



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		1		GRAND TOTAL FOR LOT 1, 2, 34

Director O. Box 5982 LTD.
DAR-ES-SALAAM

OTEL: 2122745, 2122747

TECHNICAL SPECIFICATIONS





TENDER NO. PA-010/2021-2022/G/33

FOR

SUPPLY OF ICU EQUIPMENTS AND CONSUMABLES FOR ORCI, MTWARA ZONAL, LIGULA AND

	COMPLY INOT COMPLY	5 section, epoxy coated steel & X-ray board	tops	Central control box under the bed only 1pcs battery bed end can extension 20cm	inside the side rail & head/foot board		H-13 10cm mattress, but with wave shape on one side
H		YES	YES YES YES	NO	YES YES YES	YES YES YES	YES
MOROGORO RHH	l echnical specifications	Fully Motorized ICU Bed) 5 Section PP Tops,	125mm Twin Wheel Castors. Central Locking Mech., Rubber Buffers at 4 corners Handset, ACP Box,	Battery Back Up, SMPS Bed Extension,	Linen Holder, Night Lamp, Angle Indicator.	Polymer Moulded Head and Foot Panels (Removable) Polymer Moulded Split Safety Rails (2 sets) Urine Bag Holder (Single)	S.S Heavy Duty I.V. Pole (2 hooks) Premium mattress, single section, 4", with wave shape on
Equipment Description		ICU BED MODEL: DA-2B	With 10cm mattressH-13 +I.v rod				
N/S	-	-			union de la companya		Colonia de

Pethnical specifications both sides, High quality, high density. Flame retardant foam and cover fabric having flame retardant, antiskid properties Matteres for Bed Extension Should have facility for Invasive and Non-Invasive ventilation Microprocessor Control suitable for Pediatric and Adult ventilation for ventilation Microprocessor Control suitable for Pediatric and Adult ventila
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sure High and Low, woxygen, High ave in-built exhalation life coclavable Nebulization rat least 24 hours skup. rat least 24 hours skup. ay screen. rat least 24 hours skup. rat least 24 hours skup. rat least 24 hours rat least 24 hours skup. rat least 25 hours rat least 24 hours rat least 24 hours rat least 24 hours rat least 24 hours rat least 25 hours rat least 26 hours rat least 27 hours rat least 28 hours rat least 29 hours rat least 29 hours rat least 27 hours rat least 28 hours rat least 28 hours rat least 29 hours rat least 20 ho	COMPLY / NOT COMPLY		V I MV MV illation	Smart mode AMV (optional) For Apnea Ventilation, we are using pressure control type									with 15 inches capacitive touch screen		5 years warranty from the date of B/L -hill of			
ist ont tion Y Y Y Y Y Y Y Y Y Y Y Y Y Y Y Y Y Y Y			V-A/C, P-A/C V-SIMV, P-SIMY APRV, DuoLevel CPAP, PSV PRVC, PRVC-SII NIV, Apnea Vent	For Apnea Ventile									with 15 inch		5 years warranty	lading		
ist ort tion tion		YES		Z Z	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	YES	2007
	Technical specifications		Modes of Ventilation equipped with newer modes of ventilation: Assist Control, Volume Control, Pressure Control SIMV with pressure support (Pressure and Volume control) PEEP Inverse ratio Ventilation Non invasive ventilation - BIPAP, CPAP. Apnea Ventilator, user selectable, volume and pressure control	Should have built-in safety alarms for Airway Pressure High and Low, Minute volume, High and Low, Power Failure, Low oxvoen High	Respiratory Rate, Air Source in-operable. Should have in-built exhalation filter	Humidifier: Servo controlled heated Respiratory Humidifier Temperature of delivered Gas on LED display	Temperature should be adjustable Jar should be autoclavable Nebulization assembly compatible with ventilator and circuit.	Should have interface facility Flow sensor	should have life more than I year Expiratory Unit-life	should be more than 3 years Data storage facility for at least 24 hours Internal rechargeable battery at least 30 minutes backup.	Supplied complete with compatible UPS and all standard accessories.	Power Supply: 220 - 240V, 50Hz	Modular & Suntable for Adult/Pediatric/ Patients monitoring	Minimum 15 inches multi colour TET display screen.	Five years comprehensive Maintenance inclusive.	Eight Channel digital and waveforms/traces display.	Capability of storage of patient data and printing of patient report. :	T

		Multichannel (up to 12 lead) ST segment analysis.	VEG	COMPLY / NOT COMPLY
		3 or 5 lead with cascade waveform facility	res	With 12 lead ECG
		Monitoring Diagnostic & OT 1. 6	YES	
		Simultaneous Multi - lead ECG monitoring of 7 ECG lead. 48 range 20-	YES	Better, we have 3/5 12 lead monitoring
		HR/PR Source selection facility from Automatic, SpO2 IBP and NIBP PULSE OXYMETRY: Display of Plethsmograph with Pulse strength indicator & SpO2 values and perfusion index.	YES	
		SPUZ range 1 - 100% and PR Range 20 - 230 BPM	YES	
		Lectrocardiogram (ECG) digital monitor and recorder, 12-leads detection, multi-channel recording, portable, AC and battery powered, with printer and accessories.	YES	
		Five years comprehensive Maintenance inclusive.	YES	5 years warranty from the date of B/L -bill of
		ECG diagnostic monitor and recorder with mintain DCG		lading
	ECG	analysis and full interpretation (rhythm and events), real-time and manual.6-channels recording.	YES	Yes, only auto measurement provide measurement results and diagnoses manual and rhythm no analysis) 6 channels and rhythm no analysis) 6 channels
		III, and VI - V6	YES	de la control de
		and visualization modes ECG gain 2.5, 5, 10, 20 mm/Mv and Auto, accuracy +-5%	YES	2.5, 5, 10, 20, L=10 C=5, L=20 C=10 mm/mV, Auto
			YES	30 to 300 BPM
		ranteed) 0.05 - 159 Hz, on +/-5%, calibration ce > 2.5 MOhm or >	YES	Frequency 0.05 to 150 Hz Input Impedance > 50MOhm.
-		Internal noise level <15 mV (p-p).	YES	



	lechnical specifications		
	Leakage current to patient <10 micro A With integrated keyboard, built-in printer with thermal printing head.		COMPLY / NOT COMPLY
		YES	More than 3.5 hours of continuous operation
	47	YES	
	chest electrodes, suction ball type.	YES	
	folded sheets 300ml of ECG conductive gel	YES	Thermal Z-fold A4 paper (210 mm x 295 mm)
	50Hz	YES	
	Complete with vascular, cardiac and curvelinear probes and Jelly for use in Ambulance and in the Emergency Room.	YES	machine come with vascular, cardiac and curvelinear probes and Jelly for use in
Dortshingly	Five years comprehensive Maintenance inclusive.	YES	Ambulance and in the Emergency Room. 5 years warranty
Diagonal association	heart, abdomen, pelvis, blood vessels, musculoskeletal and soft tissue.	YES	Capable of generating imaging procedures involving lungs, heart, abdomen, pelvis, blood
	CONSOLE: Laptop style console design, optional touch screen combined with conventional user control panel	ON	Laptop design, No touch screen
	conditions Supplied complete with clear protective control panel cover for infection control. Imaging Focusing:	YES	endurance >120 min
	selected scanning depth, synchronization of focal zone to the Zooming Canability with	YES	Adjustable focal depth, synchronization of
	Image Orientation:	YES	and a second as an a second a seanning depth.
	Capable of lateral and vertical Inversion (in B-mode) Image Modes: 2D Imaging, M-Mode, B/M mode, Dual 2D/Colour Imaging mode with Cine loop.	YES	lateral and vertical Inversion (in B-mode) Image Modes: 2D Imaging, M-Mode, B/M mode, Dual 2D/Colour Imaging mode with Cine loop.

6	racibilitati pescubilon	Technical specifications		
Do		Doppler, Colour Doppler Imaging (CDI), Power Doppler Imaging (PDI), Duplex, Continuous wave Doppler, Triple mode (optional)	YES	Doppler, Colour Doppler Imaging (CDI), Power Doppler Imaging (PDI), Duplex. Continuous wave
		Software Application that include at least: Obstetrics/Gynecology measurements and calculations, including gestational sac mean, mean sac diameter, femur length, crown-rump length, biparietal diameter and abdominal circumference, enabling estimation of gestational age.	YES	Doppler, Triple mode (standardl) Software Application that include at least: Obstetrics/Gynecology measurements and calculations, including gestational sac mean, mean sac diameter, femur length, crown-rump length, biparietal diameter and abdominal circumference, enabling sestimation, of
		Vascular/basic cardiac quantification Measurements capability (distance, area and circumference by ellipse and trace method) Equipment with write-zoom function available Screen annotations capture patient data, date and time, scanning	YES	gestational age. Vascular/basic cardiac quantification Measurements capability (distance, area and circumference by ellipse and trace method) Equipment with write-zoom function available Screen annotations capture patient data, date
		protocols, probes. Monitor and Display: High Definition (HD) digital black and white and colour liquid crystal display (LCD) monitor of at least 25 cm diagonal (across), with reflection filter.	YES	15" LCD screen, 38cm diagonal(across)
		broadband Curvilinear transducer of at least 5-2 MHz and 12-5 MHz transducer Power Supply: 220 - 240V, 50 Hz	YES	1.8~6 MHz convex probe 3.4~12.9MHz linear probe 200~240V, 50/60MHz
	***	Portable		
	Blood gas analyzer Model: ST-200CC	 Sample volume 110µl Result in one min after sample aspiration Power supply -220v with rechargeable battery at least 42mAh At least 10 parameter PH, pO2, pCO2, Na. K. Ca. Cl. Glu. Lac and Hot 		YES 140 μl YES 220V without reachargable battery YES
	Suction machine	le for ragm	YES	
	-{	Must have maintenance free pumps of international design for continuous use Motor shall be of Class "F" insulation to operate in ambient temperature to withstand up to 50oC, with thermal cut-outs Able to produce minimum vacuum of 700mm Hg and which must be adjustable and monitored by vacuum gauge of suitable range.		

N/S	Equipment Description	Technical specifications		COMPLY/NOT COMPLY
		The suction capacity must be 25 liters per minute and can be regulated It must have two bottles of 2l each. Each made of unbreakable polycarbonate with ABS Lid with float (overflow control device) The jars must be graduated in cc levels.	YES	40L/M
		The suction bottles shall be autoclavable On/Off Switch and power indicator must be available. Shall provide foot switch.	YES	
		Base, top and panel made of rust proof and corrosion resistant moulded ABS.	YES	
		Accessories:		
		Spare bottles: 02nos, Lid: 02nos, Rubber Seals: 02nos, Blades: 02nos, Suction Tubing set at least 5metres: 02nos,	YES	
		Spare fuse: 01 set and Bacterial filter: 05 nos.	YES	
		Supplied complete with all standard accessories Power Supply: 220 - 240 V, 50Hz		
		Made of aluminium and high density polyethylene	YES	
		5" castors standard with five drawers Drawers made of aluminium composite and high density polyethylene Built-in Security Rail Top corrosion resistant, high density polyethylene 5	YES	
	Crash Cart trolley	4 qt Sharps Dispenser with mount which can be mounted on left or right along with waste container	YES	
		Mount made of high density polyethylene Waste Container with Mount which can be mounted on left or right side along with sharps dispenser	YES	
		Mount made of high density polyethylene	YES	
		Crash Cart Handle: Made of high density polyethylene 360 degree Locking Defibrillator Mount with Security Straps: Holds up to 151bs.	YES	
	10	Made of high density polyethylene Crash Cart Cardiac Board: Made of high density polyethylene: Made of high density polyethylene Oxygen Tank Mount with Security Straps: Made of high density polyethylene.	YES	Not include the oxygen tank
	J.	Holds sizes D & E oxygen tanks Telescoping I.V. Pole with Mount: Two loop design.	YES	Only can hold the 4l oxygen tanks and include the I.V Pole
-		Extends to a maximum of 72 inches Breakaway Drawer Locks: Individual drawers or gang lock for all drawers available	YES	



Ī		lechnical specifications	Widelico TON V IGMOD
the effect of the second		Features Delivered fully assembled, no tools required Top constructed of solid high density polyethylene 100% Total Replacement Lifetime Guarantee 31 x 42 x 24 inches (width, height depth), 88 pounds Lightweight aluminium composite construction Multiple accessories available for any application 5" enclosed high compressive nylon rubber castors Adjustable and customizable built in track system	Size: L850 x W 520 x H 1045mm 4 pcs dia. 125mm castors with brakes Yes, for the drawer
		3 Set for Adults and 2 Sets for Children (Marc and Miller Type) Large hollow, cylindrical, slightly ribbed handle. Handle made of either Chromium plated or Stainless Steel	Comply – set meaning Blades
	Laryngoscope set	Can be opened to insert two batteries (type LR 14, size C, 1.5V) Stud contact, fitting various sizes and types of depressors With a set of four stainless steel depressors, with halogen bulb Marc. Type Curved Nr 2, length approx. 110mm Curved Nr 3, length approx. 135 mm Miller Type Straight Nr 1, length approx. 155 mm Miller Type Straight Nr 1, length approx. 100 mm Supplied with: -1 x Durable protective plastic box or padded vinyl case - 4 x Spare halogen bulb (one for each depressor) - 2 x Dry cell, alkaline, "C" 1.5V	S
AND ASSESSMENT OF THE PARTY OF	Nebulizer	Electrical powered with accessories Portable, compressor driven nebulizer pump Easy to operate and sturdy design for use in demanding environment Cup capacity minimum 3ml to max 10ml Nebulization approximate 0.3 to 0.8 ml/min Residual Volume maximum 2ml Flow delivery range (under load) 0.5 to max 10 L/min Operating temperature +5 to +50° C Humidity up to 65% Power Supply 220 - 240V, 50Hz, 50W Supplied with: 1 atomizer body piece 1 pediatric reusable mouth piece 1 adult reusable mouth piece 2 air tubing 9mm diameter and 1 m length Dosage cup Reusable pediatric and adult masks	professional piston compressor Nebulizer for Aerosoltherapy. Designed for intensive treatments in clinics or hospitals. Standard consumables include the adult and child masks with elastic strings, mouthpiece, nosepiece, air tube, and air filter. Compressor – Oilless and maintenance-free piston pump Power feeding – 230V-50Hz Power consumption – 180VA Max Pressure – 2.5bar 36psi 250kPa

N/S	Equipment Description	Tochnical case iff		
		recuired specifications		COMPLY / NOT COMPLY
	Blood and fluid	Fluid Temperature Range: 33 - 37°C Heat Transfer Method: Dry heat Time to heat from 23°C to usable range (33 - 37°C): 30 - 40 minutes	COMPLY	
	warmer	High Temperature protection set point: 37°C Temperature: Operation Mode: +5°C to +40°C Humidity: Operation Mode: -30% to 75% non-condensing Supply Voltage: Warmer, 15 VDC Power Supply: 220 - 240V, 50Hz		
		One Channel at least Capable of accepting any kind o fluids (solutions and medications) Pump Capabilities: Flow Range: 0		
		1 to > 999 ml/hr Increments: 0		
		1 - 100 ml/hr Keep Vein Open (KVO) Rate: 1 - 5 ml/hr Volume to be Infused Selector (VTBI): 1 - 9999 ml Flow Rate Accuracy of +/- 5% When multiple channel automatic piggybacking Ingress protection not less than IPX2 Front Panel lockout Self-check carried out on powering on Events Stored System: Log book Pause Infusion Facility required Antibolus System to reduce pressure on sudden release of occlusion IV Set Free-Flow Protection and Air Trapping Capability Needleless IV connection "Dose Error" reduction system preferable Drug library software available, including updates (free during warranty)	YES Better Flow Rate 0.1-20 Better.KVO 0.1-5ml/hr IP23 not supply administratic please use common univ Open system please use Sets	Better Flow Rate 0.1-2000ml/hr Better.KVO 0.1-5ml/hr IP23 not supply administration set. Open system please use common universal Sets Open system please use common universal Sets
		Air Bubble Detector with single and cumulative functions	VEC	ac
	Intusion Pump		Q Q	
		oecification of at least	YES	
	l	1.00	YES	
		Occlusion upstream and downstream Air in line and System malfunction Set loaded improperly and Door Open Infusion Complete and Loss of mains power Low Battery and Clinical Advisory Messages Consumables labeled "single use" Compatible Administration Set: 100	YES	



N/S	Industry meaning and in the control of the control	Technical specifications		TO LOW MINISTER
		Compatible Administration Sets micro-bore and macro-bore: 100 Compatible long Administration Sets both micro-bore/small and macro-bore or long extension sets: 100 Accessories Clamp for mounting pump on IV stand Clamp for external transportation preferable Portability Data port required, at least RS 232 and/or USB interface Long analysis software and updates provided	YES	COMPLYTON COMPLY
- * -		Wireless connectivity	YES	
of the latest the late		Event log required and recording Software to diagnose and calibrate the equipment withaccess to calibration settings Power Supply Operates from AC mains power: 220 - 240V, 50Hz In-built rechargeable battery	YES	
		Battery with operating time at least 4 hours at 25 ml/hr Automatic switch from AC mains power mode to battery operating mode and vice versa	YES	5 hour working
+		Total re-charging time not greater than 6 hours	YES	
***		Mucroprocessor controlled programmed syringe pump Programming: Up to 99 steps of speed and time, Time Resolution: 0 to 999 minutes in 1 minute steps Time	YES	
		Accuracy: +- 1% Reproducibility: +/-0.2%Syringes: Glass,		
		plastic. Metal syringes from 5 micro-liter to over 150ml Flow Rate Range: Depends on the inner syringe diameter 0.4nl/min		
	Syringe pump	With a 5ml syringe to 110ml/min with a 150ml syringe	YES	
		Maximum Force: 300N reducible by a switch to 80N Motor: Microprocessor controlled brushless long life BLDC motor with Neodymium magnets Transmission: Efficient force transmission by a ball screw with highest mechanical load capacity of 12,000N Pusher Travel: 120mm Pusher Travel Rate: Minimum: 0	YES	
		volatile Memory: Storage of all settings Power Supply: 220 - 240V, 50Hz Interface: RS-485 or RS-232 automatic valve control Remote Control: 0 - 10V Operation Temperature: 0 - 40oC Operation Humidity: 0 - 90% RH not condensing	YES	
щ а.	Portable Hand held Pulse oximeter	Patient Range: Adult, Pediatrics and Neonatal patients High Resolution, 2.4" colour display Visual and sound alarms Uses AA size alkaline or rechargeable batteries Digital SpO2	YES	4.3" TOUCH LCD Display range 0% ~ 100% SpO2 display resolution 1%



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COMPLY / NOT COMPLY	Spo2 Accuray ±2% Neonate unspecified	YES Measuring range $25 \sim 250 \text{bpm}$ Resolution $\pm 1 \text{bpm}$ Accuracy $\pm 2\%$ or $\pm 2 \text{bpm}$, whichever is greater		ch YES SPO2 alarms and various system			data	Ve min				3 01:	mg of		L L	This is not CMS specification, it's the nation monitors		,			ean	
Technical specifications	Range: 0 - 100% Resolution: 1%, Accuracy: 70% to 100% +/-2% Refreshing Rate: < 13 seconds Pitch Tone: Yes	Pulse Rate Range: 25 - 250 bpm Resolution: 1 bpm, Accuracy: +-2% or +/-1 bpm Refreshing Rate: < 13 seconds	Display: Type: 2.4" colour display 320 x 240 pixels Parameter Digital SpO2, Pulse Rate, Pleth bar and SpO2 waveform	Alarm Audible alarm, audible button tone Supports Pitch Tone and multi-level volume	General Description Modular and Suitable for	Adult/Paediatric/ Patient monitoring Minimum 15" multi	coloui 11.1 uispiay seleeti Etgir Channel digital and waveforms/traces display Capability of storage of natient data	and printing of patient reports. Five years comprehensive	Maintenance inclusive.	Parameters: Eight digital and waveforms/traces display Facilities to	monitor and display ECG, Respiration, NIBP, Et CO2, SpO2 and	5 lead with cascade waveform facility Monitoring. Diagnostic and OT	modes of monitoring of ECG Simultaneous Multi-lead ECG monitoring of 7 ECG lead	Heart Rate (HR) Range: 2 - 350 bpm HR/PR Source selection facility	from Automatic, SpO2 IBP and NIBP Automatic arrhythmia detection	alarm for standard and lethal arrhythmia Pulse Oximetry Nellcor or	indicator and SpO2 values and perfusion index Sp07 Range: 1 - 100% PR	Range: 20 to 230bpm ETCO2 Should be Main stream cannography with	display of CO2 and digital Values of ErCO2, FiCO2 and RR	EtCO2 Range: 0 - 99 numHg FiCO2 Range 0 - 20 mmHg Flow Rate:	50ml/min NIBP Measurement and display of systolic, diastolic and mean	pressure values of NIBP measurement for Adult Child and Noonata
Equipment Description												Central Monitoring	Station 8 Channel									
N/S			1000														-		-			

N/S	Equipment Description	Technical specifications	COMPLY / NOT COMPLY
		Use selectable alarm settings: Mode: Manual, STAT (continuous 5 minute operation) and automatic (selectable time interval 2 - 90 minutes) Range: 20 - 250 mnHg Temperature Two channel and with two units (oC and o F) selectable Temperature Range: 0 - 50oC	
		Option for differential temperature Should be provided Respiration: RR range 1 - 150 bpm Apnea alarms should be provided Trends and Alarms 72 hours	
		Non volatile graphical tabular trends with zoom facility and separate dedicated trend for storing minimum200 NIBP readings Should have Alarm recall facility for last 24 Alarm events with date, time and message Should have facility to print Graphical trend, tabular trend and alarm recall	
8.41	•	Recorder In-built dual channel thermal array recorder	
		Include Laser Printer and dual channel strip chart recorder Others Defibrillator and Cautery protection should be provided	
		Should work on Mains as well as Battery (backup for 2 hours) Automatic zoom in facility in the monitor display	
		Should have facility to download trend data on USB and SD Card Supplied complete with all standard accessories and consumables that will enable the machine to operate	
		Power Supply: 220 - 240V.50 Hz	
		CENTRAL STATION Central Station should have facility to display up to 20 real time waves at a time and upgradable	S better, if display resolution 1920×1080, support display 36 Patient Sectors with max 3 waves ner natient
		Central Station should have separate patient window for viewing detailed YES real-time or stored data for individual patient	
		It should have 24 hr stored patient data monitoring trends and 24 hr event review facility	8
		The Central Nurse Station should also have the following: - Multi-lead arrhythmia and ST review facility - 50 alarms strips storage per bed - Offer wave review with 24 hr full disclosure - Optional facility for dual display for detailed analysis of individual bed without compromising on full ICU monitoring	
3770			parameter name and 10-second wavetorm before and after an alarm is triggered
		Facility for interfacing Holter data for analysis, in case of the Holter from YES the same brand is available	



COMPLY / NOT COMPLY							Display: 1 page maximum: 3, turn the page to view all waves the monitor have	Windows® 7 Professional SP1	■ Support Windows® 10	■ Support Windows® Server 2008	■ Support Windows® Server 2012 R2	Support Windows® Server 2016	24" Monitor			Central Venous Catheter Triple Lumen 5FR 200mm	Central Venous Catheter Triple Lumen 7FR	Central venous Cameter Triple Lumen 4.5 FR 80mm	COMPLY	COMPLY	PLY	Better, from 1J and up to 360J
									nS ■	nS ■	nS ■	nS ■	24			Comply ALL Central	Cent	FR 8	CON	CON	COMPLY	
	YES	YES	YES	YES	YES	YES	YES	YES					YES		YES	Соп						YES
Technical specifications	Export the ICU Patient data to Holter for analysis	Remote display (Slave) facility should be available	Real time recording thru Dual Channel Recorder	Facility for interfacing a laser printer for printing patient information and trend formats	Advanced arrthythmia analysis package of at least 20 arrthythmia analysis	12 lead ECG Monitoring and view possible at Central Nurse Station	Continuous full disclosure of up to 4 configurable waves per patient					Operate on Microsoft Windows NT workstation operating system	Supplied with 19" Flat Screen TFT display, Laser Printer and Recorder, UPS	- The entire networking and cabling witrh hardware Wall mounts	Power Supply: 220 - 240 V, 50Hz	Set of Sizes 3, 4, 5, 6, 7, 8, 9, 10			Set of Sizes 0,1, 2, 3, 4, 5 for Adult and Pediatrics	Set of Sizes 3, 4, 5, 6 for Adults and Pediatrics	Set of Sizes 3, 3.5, 4, 5, 6, 6.5, 7, 7.5, 8	Should be a Low Energy Biphasic defibrillator monitor with Recorder, having capability to deliver shocks from 2 Joules to 200Joules.
Equipment Description																	Central Vernous	catheters (CVC)	Oropharyngeal Air way (OPA)	Nasalpharyngeal Air Way (NAP)	Endotrachea	defilbrillator
S/N																						



		Tochairal engolifications		COMPLY / NOT COMPLY
N/S	Equipment Description			
		Five years comprehensive Maintenance inclusive.	YES	5 years warranty from the date of B/L -bill of lading
		Should monitor ECG through paddles, pads and monitoring electrodes and defibrillate through pads and paddles	YES	
		Should compensate for body impedance for a range of 25 to	YES	better, 25 - 300 ohms
		Should have charging time of less than 5 seconds for	YES	better, less than 3s charge to 200J
		Should have High resolution more than 8 inch colour display for viewing monitoring parameters like ECG, SpO2, NIBP	YES	8.4 inches screen for ECG, SPO2, NIBP, Et CO2
		t and Pediatric pad t summary facility	YES	better, up to 1000 event
		55 events Should have a battery capable of usage for at least 5 hours of	YES	better, 6 hours monitoring
		monitoring Should be capable of printing Reports on event summary, configuration, self test, battery capacity etc Should have facility for self test/check before	YES	better, up to 1000 evenT
		usage and set up function. Should have facility to monitor parameters like SpO2, NIBP and etCO2 along with non invasive pacing (Demand and Fixed mode) facility	YES	with SPO2, NIBP, CO2 monitoring
		Should be able to upgrade the defibrillator for 12 lead ECG monitoring and ECG transmission System	YES	Could be upgrade to 12 leads ECG
		Configuration Accessories, spares and consumances. Defibrillator with AED and external Pacemaker - 1pc Adult with built in Pediatric External Paddles - 1pc Patient Cables - 1pc ECG Rolls 50 Adult Pediatric External Paddles - 1pc Patient Cables - 1pc EtCO2	YES	
		SpUZ reusable Sensor - 100 Adultifunction Pads for Adults - 10 pairs Tubing (box of 20) - 1box AED Multifunction Pads for Adults - 10 pairs with Fach unit	YES	
		The unit shall be capable of operating continuously in ambient temperature of 5 to 45oC and relative humidity of up to 95%	YES	



13.0	Carrimont Decription	Technical specifications		COMPLY / NOT COMPLY
N/O	Equipment pescription	f safety for Electromagnetic	YES	
			YES	
		Fowel Input. 240 VAC, Solite		COMPIV. Bonnies
	Endotrachea Tube Introducers (Stylate and Bougie)	A disposable, latex free, sterile and individually wrapped device. Markings: at 20 cm, 30 cm, and 40 cm intervals. It consists of a 50 to 60 cm stylet with the distal tip bent at a 30 degree angle Endotracheal Tube Introducer bougie: - size 15 FR, OD 0.5mm, 700 mm - size 10 FR, OD 3.3mm, 700 mm -		
		SIZO U.N, OD Z.O mini, 550 mini		ASS. COV. Y.
	Magills Forceps	Made from stainless steel and can be disinfected and sterilized Twin-bladed tong-like forceps Handles for gripping by the user Rounded ends for grasping Oblique angle between handles and blades to prevent obstruction of the view of the airway during use. Sizes for Adult and Child use		COMPLY
		Nasal cannula (nasal prongs), device designed for easy administration of oxygen into the patient nose through two small prongs placed in the nostrils. It consists of soft twin		COMPLY
	Nasal Cannula (prongs)	prongs nasal tips to ensure equal oxygen flow to both. Adjustable, smoothly finished nasal tips for maximum patient comfort Star lumen main tube to avoid accidental blockage. Made from soft and kink resistant polyvinyl chloride (PVC)		
	Wall mounted otoscope	material For Adult and Child use Illumination: Pressurized halogen lamp Image Resolution: 1280 x 1024 pixels Connection/Interface: USB 2.0 with 3m cable System Requirements: Windows XP Service Pack Electrical Rating: Lamp 3.5V DC, 810mA USB: 5V DC, 150mA Focal length: 114mm and Magnification: 2.2X		COMPLY
		Constructed from ABS and Acrylic Plastic or Stainless Steel Automatic Image Brightness and white balancing for optimal		
		Illiage		



olley Colley	Equipment Description	quality Supplied complete with Charger Range of lenses not smaller than -20D to +29D with steps not smaller than lens Magnification: 12 - 15X		COMPLY
type only) Sufficient light bulbs for 3 years uso. Stainless steel shelves and railings Available in full stainless YES Stainless steel shelves and railings Available in full stainless YES Stainless steel bottom and top shelves with railing on top shelf Frame Mounted on 100 mm diameter wheels Overall Size: L YES 710 mm x W460 mm x H830 mm	wall mounted Opthalmoscope	Apertures: Small, Large and Semi Circle, Fixation Star Apertures: Small, Large and Semi Circle, Fixation Star Colour Temperature: Cool White in the range 3100 - 5500K Colour Temperature: Cool White in the range 3100 - 5500K Light Intensity: 8,000 - 12,000 lux. Dust free sealed optics and aspherical optical system Red-free, Blue, Green and and aspherical optical system Red-free, Blue, Green and polarization filters Handle with On/Off switch. Scratchproof polarization filters Handle with On/Off switch. Scratchproof lens: Glass or Plastic Batteries: AA or rechargeable. Battery Charger (as applicable to rechargeable Accessories: Battery Charger (as applicable to rechargeable		
Stainless steet sheres and tame. Steel/powder coated frame. Stainless steel bottom and top shelves with railing on top Stainless steel bottom and top shelves with railing on top Shelf Frame Mounted on 100 mm diameter wheels Overall Size: L 710 mm x W460 mm x H830 mm		type only) Sufficient light builts and years years. Battery 1.5V. Battery 1.5V.	YES	Stainless steel surface without powder coated
shelf Frame Mounted on 100 mm diameter wheels Overall Size: L YES 710 mm x W460 mm x H830 mm		Stainless steel sucryes and top shelves with railing on top Stainless steel bottom and top shelves with railing on top	YES	OFORTAGE
Man Man	=	shelf Frame Mounted on 100 mm diameter wheels Overall Size: L	YES	Size: L730*W4/0" II970
	nent trolley			



Made of stainless steel with removable top tray Height adjustable by foot 75 mm swivel castors Dimensions: YES Width 60 cm, Depth 40 cm and Height 120 - 130 cm Width 60 cm, Depth 40 cm Width 6	Equipment Description	Technical specifications		
Five-claw wheel Adjustable IV Pole • Material: Cast iron base 5 Yes Wheels.• Down Tube: \$\phi 25.• Top Tube: \$\phi 16.• Adjustment mode: Rotary Knob / Jackscrew.• Height: 2m, 2 hooks.	Adjustable Mayo table	Made of stainless steel with removable top tray Height adjustable by foot 75 mm swivel castors Dimensions: Width 60 cm, Depth 40 cm and Height 120 - 130 cm	YES	Size: L600*W400*H850-1000mm
	IV Pole	Five-claw wheel Adjustable IV Pole • Material: Cast iron base 5 Wheels.• Down Tube: φ 25.• Top Tube: φ 16.• Adjustment mode: Rotary Knob / Jackscrew.• Height: 2m, 2 hooks.	Yes	IV stand (type C) main feature: 1. stainless steel round tube and ABS plastic base; 2. height can be adjusted from 135cm to 240cm; 3. top with 4 chrome hooks, base with castors



N/U	Fortinment Description	Technical specifications		COMPLY / NO! COMPLY	
5					
		• Dimensions(mm): 850x500x39mm	YES	Frame: 857 x 502 x 25mm	
	x-ray viewer	• Film: 14 x 17" • View area: 730x442mm	YES	View area: 714 x 442mm	
		• Lamp: CCFL 12pcs	YES	40117	
	MST-4001	• Luminance: 12,000lux or more		48 W A C90-240V 50/60Hz	
	-	• Power consumption: 60W	VES		
		• Power: 220VAC 50/60HZ	I LO		
		• Life Time: over 80,000hr	IES		
		Film Clipping: Spring Roller or similar			



This Machine should be able to provide a minimum of 300 images per day. Should have a high frequency generator of 50-150k.W., Automatic exposure device, Anatomical programming radiography, overloading protection feature, digital display of KV and mAs. X-Ray tube should be Floor stand mounted. Axis should rotate 360 degree Five years comprehensive Maintenance inclusive. High speed rotating anode and exposure should be 50 – 150k.V and 0-600mA. Heat strength capacity of the anode at least 150,000HU. The digital detector (Two fixed Detectors) should be flat panel of latest technology. The digital workstation should have high speed processors, preview time of 5s or less. The workstation should provide basic functions for image processing and be compatible to other HIS Displayed Parameters: The console monitor should display patient Displayed Parameters: The console monitor should display patient awith brakes to allow Longitudinal and lateral movement and better patient position. Floating table with up and down movement Table Size from 1800x800mm to 200x800mm Patient weight: at least 150Kg Bucky wall stand: Height: 1900mm Center Height Stroke Range from 400mm to 1600mm Source to image detectors should include the range of 90cm to 125cm Detectors: Fixed Flat Panel Detectors Fixed Flat Panel Detectors Fixed Flat Panel Pates 17 more 18 more 17	
Pixel Size at least 17 men of Pixel Size at least 140 micron Dimension: 460 x 460 x 15mm	This Machine should be able to provide a minimum of 300 images or day. Should have a high frequency generator of 50-150KW, Automatic exposure device, Anatomical programming radiography, Automatic exposure device, Anatomical programming radiography, Automatic exposure device, Anatomical programming radiography, Averloading protection feature, digital display of KV and mAs. X-axy tube should be Floor stand mounted. Axis should loate 36 of degree Axis should be Floor stand mounted. Light speed rotating anode and exposure should be 50 – 150KV and 1500nHU. Lodollan-A. Heat strength capacity of the anode at least 150,000HU. The digital workstation should have high speed processors, preview intensity and better high speed rotating should be best functions for image processing and be compatible to other HIS. Components: Patient table: Mounted on heavy duty four castors with bracks to allow to console monitor should display patient better stories, warning sign and other important parameters. The console monitor should important parameters. Components: Patient table: Mounted on heavy duty four castors with bracks to allow to congulate in 1900mm of 200x800mm to 1250m. Conter Height Stroke Range from 400mm to 1600mm Center Height Stroke Range from 400mm to 1600mm Detectors: Active Image Size at least 17 inch x 17 inch or 43cm x 43cm Pixel Size at least 140 micron Dimension: 460 x 460 x 15mm
Dust cover for control unit to be supplied. Protection against insect and rodent ingress to be incorporated.	if to be supplied.



Z	Equipment Description	Technical specifications	COMPLY / NOT COMPLY
2/0	Edulphient Describer	Total Control of the	
		Electrical Requirements: Should meet Tanzania Electrical	
		Standards (Voltage of between 220-240V with the standard	
		Standards Country of the Country Assessment	
		frequency of 50Hz) with type G adaptor system. Accessories.	
		Multi-tray printer for different X-Ray film sizes (10 x 12, 14 x 14,	
		14 v 17 10 x 8) Protective sear (lead apron minimum of 2 small, 2	
		11 A 11, 12 A J. 12 L. 1	
		medium and 2 large), Googles (Millinnin 1 sinan, 1 modium	
		large). Gonad shield (minimum 2 small, 2 medium, 2 large), neck	
		collar chiefd (minimum of 1small, 1 medium, 1 large), Gloves	
		County Smooth (managed of the state of the s	
		(Minimum 2 small, 2 medium and 2 large). National mazarus	
		warning signs to be supplied with the machine. It should come with	
		de Production Contain with canacity of nower storage for not	
		the Fowel Dackup System with Capacity of power	
		less than 15 Minutes. Radiation protective gears: A-ray should	
		with (i) two (2) nairs of lead anrons with back protection	
		Collic With, (1) two (2) puns of rock of the Model Classic	
		0.35/0.25mmPb, (11) two (2) pairs of Inyrold Silled Model Classic	
		0.35mmPh (iii) two (2) pairs of Patient Apron with belt 0.50mmPb	
		r and M. 201 20 Printer Multi-tray printer for different X-Ray	
		Leadille W/ U-30 L-301 Illicitions and form	
	1	film sizes (10 x 12, 14 x 14, 14 x 17, 10 x 8). Training, installation	
		and UtilizationRequirements for commissioning:	
		Manufactures/countries should nerform installation. safety and	
		Manuacunier/supplier success percent meaning	
		operation checks before handover. Acceptance lesis to be specified	
		and local clinical and technical staff to verify proper and full	
		functioning of device. Training of user/s: Application specialist shall	
		provide training of users in operation and basic	
		provide damas or asset in a commence commence of the commence	
		maintenance. Warranty and Maintenance: rive years compromisive	
		maintenance should include service, spare parts and labour starting	
		from the day of the acceptance testing of the machine. Lifetime	
		support: spare parts consumables should be available throughout	
		outport, open parcy comments	
		The menuic period of the machine.	



Edulpment pessenbusen	Technical space regions	
	Uptake time should be a minimum of 90%. Proof of locally available technical support personnel, including	
	CVs and work permit for foreign personnel.	
	Availability of technical personnel within the County should be stated; this should include CVs, work permits for foreign personnel.	
	Software should be flexible and provide the room for upgrade to add new parameters to be measured by the Machine and report	
	format.	
	Documentation: Operating and service manuals (In English)	
	including lists of important spares and accessories - with their part	
	numbers and list of equipment and procedures required for	
	calibration and routine maintenance should be provided.	
	Documentation must also show recommended procedures for	
	disposal and any probable hazards to the environment and/or	
	community. Tife and of the machine should be not less than 10	
	Luc apan. Luc apan of the machine strong of received the received to the recei	
	Risk Classification: As per ISO 14971:2007- Application of risk	
	management to medical devices.	
	Regulatory Approval / Certification: TAEC, TBS and TFDA	



COMPLY / NOT COMPLY	
	COMPLY
Technical specifications	
Equipment Description	Neonatal ventiliator MODEL: SV300PR0
No	



Equipment Description 1.E ratio 1.0 to 1:10 Inspired Time 0.1 to 2 sec Expired Time 0.2 to 30 sec Frequency Up to 200 BPM Base Flow (VIVE) 1 to 30 LPM Synchronization Patient synchronization Integrated blender for Oxygen 21 % to 100% Integrated mebulization facility Integrated screen for display of Pressure-Time, Flow-Time and Volume-Time curves Integrated monitoring of FiO2 Monitoring of flow: At the Y piece with facility to activate it Audiovisual alarms with advisory on-screen message: MV high/low, Apnea, tube obstruction, FiO2, high/low high PIP, low PEEP/CPAP, fail to cycle, gas supply low, power failure, ventilator inoperative, alarm log book. The ventilator should have automatic compensation for leakage and should monitor and display leakages: The ventilator should show trends of important parameters viz. C.R, FiO2, MAP, etc for evaluation of patient improvement Power Supply: 220 - 240V, 50Hz. SCOPE OF SUPPLY Ventilator on trollesy with wheels and brake facility Integrated medical air compressor Humidifier with autoclavable chamber and complete with patient circuit Circuit support arm Nebulizzation accessories (2 sets) Bacterial filters 50 sets Flow sensors 20 sets Oxygen cell and Oxygen connecting hose Air connecting hose Battery back-up (at least 30 minutes) Hand held bype Glucometer Statery operated "Memory up to 10 measurements Sicks method measurements One box of sticks in available: Indication of high and low measurements and punctures is included Operating instructions is included			Comply
Technical 1:E ratio 1:0 to 1:10 lb Expired Time 0.2 to 30 see Base Flow (VIVE) 1 to 30 L synchronization Integrated ble Integrated nebulization facilit of Pressure-Time, Flow-Time Integrated monitoring of FiO: piece with facility to activate advisory on-screen message: obstruction, FiO2, high/low h cycle, gas supply low, power alarm log book. The ventilato compensation for leakage and leakages The ventilator shoul parameters viz. C,R, FiO2, M improvement Power Supply: SUPPLY Ventilator on trolle Integrated medical air comprautoclavable chamber and co Circuit support arm Nebuliza Bacterial filters 50 sets Flow Oxygen connecting hose Air (at least 30 minutes) Hand held type Glucometer Batt measurements sicks method is available Indication of high and and punctures is included Operati			
Glucometer	Technical specifications	1:0 to 1:10 lb 0.2 to 30 sec VE) 1 to 30 L n Integrated blo ulization facilit me, Flow-Time intoring of FiO. Lity to activate reen message: O2, high/low high low, power K. The ventilator on trolle for leakage and ventilator on trolle dical air compribations. Can be supply: Itilator on trolle dical air compribations hose Air for So sets Flow cetting hose Air fintes)	Hand held type Glucometer ·Battery operated ·Memory up to 10 ·Hand held type Glucometer ·Battery operated ·Memory up to 10 measurements ·Sticks method measurements ·Code of sticks interring is available ·Indication of high and low measurements One box of sticks and punctures is included Operating instructions is included
В	Faminant Description		Glucometer

COMPLY / NOT COMPLY Hemoglobin an Hemotrocrit result from one sample available after 25-60 seconds available after 26-60 seconds sul venous, arterial or capillary blood sample Stores upto 4000 patient reults Measuring range: 0-25.6g/dL: 0-15.9mmol/L	Manable semi-fowler bed with ABS headboards B-21-1	Moverage 2740mm(L)*940mm(W)*500mm(r) Oversize: 2140mm(L)*940mm(W)*500mm(r) Main features: 1. Epox coated bed frame, with Collapsible aluminum alloy side rails, and alloy side rails, and detachable ABS engineering plastic headboards, detachable ABS engineering plastic headboards, cation and single function, which can be operated by crank, the 2. single function, which can be operated by crank, the details are as follows: (1)back rest:0-70±2° (1)	5. Freight saving knock-down consumers.	ABS Bedside cabinet ABS bedside cabinet With one foot board, one shelf board)	Movable over bed table F-32 Size: L795*W400*H700~960mm	main feature: 1. epoxy coated steel frame with wooden top board 2. the height can be adjusted by gas-spring from 700 to 2. the height can be adjusted by gas-spring from 700 to 960mm 2. the height can be adjusted by gas-spring from 700 to		M 2125748
Technical specifications Parameter: Hemoglobin, HCT(Hematocrit) Principle: Optical reflectance · Sample: Capillary or venous whole blood (13~15ul) · Strip: H12 Hemoglobin Test Strip · Speed: ≤ 10 seconds · Measuring Range: H12 Hemoglobin Test Strip · Speed: ≤ 10 seconds · Measuring Range: A · Og/dL~25.6g/dL(g/L optional) · Memory: 1200 test results A upwatization: Self-checking, auto-judging and displaying malfunction		• Heavy duty bed Size (L×W×H): 2150×950×500mm• High quality section steel welding• The surface electrostatic sprayed• Corrosion resistant• Easy to clean• Side guard made of well plastic• I.V pole• Four casters, two caster with brakes		• dimension 520*480*740 • made of high density wood • one drawer and one door with handles, top with rail • two tower hanger, four wheels with cross breaks	o. ooo*4<0*900-1200mm	Size:900° 4-50° 50° 50° 50° 50° 50° 50° 50° 50° 50°	castors, \$55mm ANUDHA LIMIT	DAR ES SALAAM (TEL: 2121188, 2125748 (FAX: 2128490
Equipment Description Hemoglobinometer		Patient Beds with matress, Model:B-21-1	7	Bedside lockers		Cardiac table		





[Text of Technical Specification to be inserted in the Tendering Documents by the Procurement Entity, as applicable

S/N	Equipment Description	Technical specifications
	ICU Beds	Fully Motorized ICU Bed) 5 Section PP Tops, 125mm Twin Wheel Castors. Central Locking Mech., Rubber Buffers at 4 corners Handset, ACP Box, Battery Back Up, SMPS Bed Extension, Linen Holder, Night Lamp, Angle Indicator.Polymer -Moulded Head and Foot Panels (Removable) - Polymer Moulded Split Safety Rails (2 sets) Urine Bag Holder (Single) S.S Heavy Duty I.V. Pole (2 hooks) Premium mattress, single section, 4", with wave shape on both sides, High quality, high density, flame retardant foam and cover fabric having flame retardant, antibacterial, antiskid properties
		Mattress for Bed Extension Should have facility for Invasive and Non-Invasive ventilation
		Microprocessor Control suitable for Pediatric and Adult ventilation Electromagnetic Compatible Hinged arm holder for holding the circuit Should have built-in touch colour screen TFT display of minimum 10 inches or more for display of waveforms and Monitored value. Five years comprehensive Maintenance inclusive. Should have in-built facility to upgrade with EtcO2 Facility to Measure and Display: Status Indicator for ventilate and Display: Status Indicator for ventilate and Display.
		Setting Automatic compliance and leakage compensation for circuit and ET Tube Should have facility of log book, for events and alarms with date and time Should have the following settings: Tidal volume (Minimum at least 50ml, Maximum up to 2000ml Inspiratory Pressure (up to 80cm of water) Respiratory rate 1 to 80 bpm
	Ventilator	Apnea back up rate CPAP/PEEP Pressure support FiO2 Pause Time Pressure and Flow Trigger Inspiratory Flow up to 120bpm Monitoring and Display of the following Parameters: Airway Pressure (Peak and Mean) Tidal Volume (Inspired and Expired) Minute Volume (Inspired and Expired) Respiratory mechanics, Spontaneous Minute Volume Total Frequency, FIO2 dynamic, Intrinsic PEEP, Plateau Pressure, Resistance and Compliance.
		Use selector Alarm for all measured and monitored parameters, Occlusion Pressure and Pressure Flow and Volume curves.
		Modes of Ventilation equipped with newer modes of ventilation: Assist Control, Volume Control, Pressure Control SIMV with pressure support (Pressure and Volume control) PEEP Inverse ratio Ventilation Non invasive ventilation - BIPAP, CPAP. Apnea Ventilator, user selectable, volume and pressure control Should have built-in safety alarms for Airway Pressure High and Low, Minute volume, High and Low, Power Failure, Low oxygen, High
		Respiratory Rate, Air Source in-operable. Should have in-built exhalation filter Humidifier: Servo controlled heated Respiratory Humidifier Temperature of delivered Controlled.
		Temperature should be adjustable Jar should be autoclavable Nebulization assembly compatible with ventilator and circuit.
		Should have interface facility Flow sensor



S/N	Equipment Description	Technical specifications
	Description	should have life more than 1 year Expiratory Unit- life
	+	should be more than 3 years Data storage facility for at least 2 in the
		least 30 minutes backup.
		least 30 minutes backup. Supplied complete with compatible UPS and all standard accessories.
		Power Supply: 220 - 240V, 50Hz
		Power Supply: 220 - 240V, 50Hz Modular & Suitable for Adult/Pediatric/ Patients monitoring Minimum 15 inches multi colour TET display screen. Five years comprehensive Maintenance
		MAN NEW CONTRACTOR CON
	1	Eight Channel digital and waveforms/traces display.
		Eight Channel digital and waveforms are properly and printing of patient report.: Capability of storage of patient data and printing of patient report.: Facility to monitor and display:-ECG, Respiration, NIBP, SpO2, EtCO2 and Temperature. ECG
		Facility to monitor and display:-ECG, Respiration, 1997, of
	Monitor	Multichannel (up to 12 lead) ST segment analysis.
		3 or 5 lead with cascade waveform facility. Monitoring, Diagnostic & OT modes of monitoring ECG lead Simultaneous Multi - lead ECG Monitoring, Diagnostic & OT modes of monitoring ECG lead Simultaneous Multi - lead ECG
		Monitoring, Diagnostic & C.1 Indeed Monitoring, Diagnostic & C.1 I
		HR/PR Source selection facility from Automatic, SpO2 IBF and NIBF 1 6 36 4. Display of Plethsmograph with Pulse strength indicator &SpO2 values and perfusion index.
		1 DD Dange 20 - 230 BPW
		Electrocardiogram (ECG) digital monitor and recorder, 12-leads detection, multi- channel recording, portable, AC and battery powered, with printer and accessories. Five years comprehensive Maintenance inclusive.
		Five years comprehensive Mantenance and Figure ECG analysis and full
		ECG diagnostic monitor and recorder with printer ECG analysis and full interpretation (rhythm and events), real-time and manual.6-channels recording.
		Tagg F 12 standard derivations aVR, aVL, aVF, I, II, III, and VI - VO
		Simultaneous 12-leads acquisition adjustable ECG acquisition and visualization modes ECG gain 2.5, 5, 10, 20 mm/Mv and Auto, accuracy +-5%
		Adjustable ECG sweep (trace speed in mm/s) HR range 30 - 300 bpm with rhythm analysis Common Mode Rejection Ratio (CMRR) >89 dB.
	ECG	Frequency response (minimum guaranteed) $0.05 - 159$ Hz, accuracy of input signal reproduction +/-5%, calibration signal 1 mV +/- 5% Input Impedance > 2.5 MOhm or > 50MOhm.
		Internal poise level $<15 \text{ mV} \text{ (p-p)}$.
		Leakage current to patient <10 micro A With integrated keyboard, built-in printer with thermal printing head.
		Built-in rechargeable battery, autonomy >2.5 hours or 800 examinations.
		Automatic switch to battery in case of power failure and automatic recharge on connection to main
		ITEMS SUPPLIED WITH: 1 x patient cable 6 x reusable chest electrodes, suction ball type.
		4 x reusable clamp electrodes 1 x supply of 960 thermal Z-folded sheets 300ml of ECG conductive gel
		1 x spare rechargeable battery pack Power Input: 240 VAC, 50Hz



S/N	Equipment	Technical specifications
SIN	Description	Complete with vascular, cardiac and curvelinear probes and Jelly for use in Ambulance and in the Emergency Room . Five years comprehensive
		Maintenance inclusive. A properties imaging procedures involving lungs, heart, abdomen, pelvis,
		blood vessels, musculoskeletal and soft tissue. CONSOLE: Laptop style console design, optional touch screen combined with
		conventional user control paner
		Battery Duration: minimum 2 nours under normal and complete with clear protective control panel cover for infection control. Imaging
		Focusing: Adjustable focal depth, synchronization of focal zone to the selected scanning
		depth. Zooming Capability with automated image optimization.
	Portable	Zooming Capability with
	ultrasoun d	Image Orientation: Capable of lateral and vertical Inversion (in B-mode) Image Modes: 2D Imaging, M-Mode, B/M mode, Dual 2D/Colour Imaging mode with Cine loop.
		Doppler, Colour Doppler Imaging (CDI), Power Doppler Imaging (193), 2 3
		Doppler, Triple mode (optional) Software Application that include at least: Obstetrics/Gynecology measurements and calculations, including gestational sac mean, mean sac diameter, femur length, crown-rump length, biparietal including gestational sac mean, mean sac diameter, femur length, crown-rump length, biparietal including gestational sac mean, mean sac diameter, enabling estimation of gestational age.
		Vascular/basic cardiac quantification Measurements capability (distance, area and circumference by ellipse and trace method) Equipment with write-zoom function available Screen annotations capture
		patient data, date and time, scalling
		protocols, probes. Monitor and Display: High Definition (HD) digital black and liquid crystal display (LCD) monitor of at least 25 cm diagonal (across), with reflection filter. Broadband Curvilinear transducer of at least 5-2 MHz and 12-5 MHz transducer Power Supply: 220
		• Portable
		• Sample volume 110µl
	Blood ga analyze	• Power supply -220v with recnargeable battery at reast 12
		• At least 10 parameter pH, pO2, pCO2, Na, K, Ca, Cl, Glu, Lac and Hct pH, pO2 and possible on four wheels and with a handle for transportation. The The machine shall be portable on four wheels and with a handle for transportation. The
		suction pump must be totally off-free diaphtagin type:
	Suction machin	Must have maintenance free pumps of international design for continuous use Motor shall be of Class "F" insulation to operate in ambient temperature to withstand up to 50oC, with thermal cut-outs Able to produce minimum vacuum of 700mm Hg and which must be
		adjustable and monitored by vacuum gauge of sumore range.
		The suction capacity must be 25 liters per minute and can be regulated It must have two bottles of each. Each made of unbreakable polycarbonate with ABS Lid with float (overflow control device)
		The jars must be graduated in cc levels. The suction bottles shall be autoclavable On/Off Switch and power indicator must be available. Shall be autoclavable on/Off Switch and power indicator must be available.



5/N	Equipment Description	Technical specifications
	Description	Base, top and panel made of rust proof and corrosion resistant moulded ABS.
		Accessories: Spare bottles: 02nos, Lid: 02nos, Rubber Seals: 02nos, Blades: 02nos, Suction Tubing set at least
		Spare fuse: 01 set and Bacterial filter: 05 nos. Supplied complete with all standard accessories Power Supply: 220 - 240 V, 50Hz
		a i i and bigh density horizonto
		5" castors standard with five drawers Drawers made of aluminium composite and high density
		d at Sharps Dispenser with mount which can be mounted of the
		Mount made of high density polyethylene Waste Container with Mount which can be mounted on left or right side along with sharps dispenser
		Mount made of high density polyethylene Crash Cart Handle: Made of high density polyethylene 360 degree Locking Crash Cart Handle: Made of high density polyethylene 360 tegree Locking Crash Cart Handle: Made of high density polyethylene 360 tegree Locking
	Crash Cart trolley	Made of high density polyethylene Crash Cart Cardiac Board. Made of high density polyethylene Oxygen Tank Mount with Security Straps: Made of high density
		polyethylene
		Extends to a maximum of 72 inches Breakaway Drawer Locks. Individual
		all drawers available Features Delivered fully assembled, no tools required Top constructed of solid high density polyethylene 100% Total Replacement Lifetime Guarantee 31 x 42 x 24 inches (width, height depth) 88 pounds Lightweight aluminium composite construction Multiple accessories available for any application 5" enclosed high compressive nylon rubber castors Adjustable and customizable built in
		track system
		3 Set for Adults and 2 Sets for Children (Marc and Miller Type) Large hollow, cylindrical, slightly ribbed handle. Handle made of either Chromium plated or
	Laryngos ope set	Stainless Steel Can be opened to insert two batteries (type LR 14, size C, 1.5V) Stud contact, fitting the contact of depressors with a set of four stainless steel depressors,

S/N	Equipment	Technical specifications
,,,,,	Description Nebulizer	Electrical powered with accessories Portable, compressor driven nebulizer pump Easy to operate and sturdy design for use in demanding environment Cup capacity minimum 3ml to max 10ml Nebulization approximate 0.3 to 0.8 ml/min Residual Volume maximum 2ml Flow delivery range (under load) 0.5 t0 max 10 L/min Operating temperature +5 to +50° C Humidity up to 65% Power Supply 220 - 240V, 50Hz, 50W Supplied with: 1 atomizer body piece 1 pediatric reusable mouth piece 1 adult reusable mouth piece 2 air tubing 9mm diameter and 1 m length Dosage cup Reusable pediatric and adult masks
	Blood and fluid warmer	Reusable pediatric and adult masks Fluid Temperature Range: 33 - 37°C Heat Transfer Method: Dry heat Time to heat from 23°C to usable range (33 - 37°C): 30 - 40 minutes High Temperature protection set point: 37°C Temperature: Operation Mode: +5°C to +40°C Humidity: Operation Mode: -30% to 75% non-condensing Supply Voltage: Warmer, 15 VDC Power Supply: 220 - 240V, 50Hz One Channel at least Capable of accepting any kind o fluids (solutions and medications) Pump
		Capabilities: Flow Range: 0 1 to > 999 ml/hr Increments: 0 1 - 100 ml/hr Keep Vein Open (KVO) Rate: 1 - 5 ml/hr Volume to be Infused Selector (VTBI): 1 - 9999 ml Flow Rate Accuracy of +/- 5% When multiple channel automatic piggybacking Ingress protection not less than IPX2 Front Panel lockout Self-check carried out on powering on Events protection not less than IPX2 Front Panel lockout Self-check carried out on powering on Events protection not less than IPX2 Front Panel lockout Self-check carried out on powering on Events protection and System: Log book Pause Infusion Facility required Anti-bolus System to reduce pressure on sudden release of occlusion IV Set Free-Flow Protection and Air Trapping Capability Needleless IV connection "Dose Error" reduction system preferable Drug library software available, including updates (free during warranty) Air Bubble Detector with single and cumulative functions preferable Clearly visible optical alarms Acoustic Alarm not less than 45Db Real Time Display Availability of a Nurse call system connectable to a staff alerting system, 24V/0
	Infusion Pump	2A static or dynamic preferable Continuous operation within specification in ambient temperature of at least 5 - 40oC, Relative humidity of at least 10 - 90% noncondensing Monitored and Displayed Parameters (colour and graphic preferable) Flow Pressure Dose Availability of software to monitor the delivery of drugs preferable Alarms Audible alarm required with volume control Momentary silence less than 2 min Occlusion upstream and downstream Air in line and System malfunction Set loaded improperly and Door Open Infusion Complete and Loss of mains power Low Battery and Clinical Advisory Messages Consumables labeled "single use" Compatible Administration Set: 100 Compatible Administration Sets micro-bore and macro-bore: 100 Compatible long Administration Sets both micro-bore/small and macro-bore or long extension sets: 100 Accessories Clamp for mounting pump on IV stand Clamp for external transportation preferable Portability Data port required, at least RS 232 and/or USB interface Long analysis software and updates provided
		Wireless connectivity Event log required and recording Software to diagnose and calibrate the equipment withaccess to calibration settings Power Supply Operates from AC mains power: 220 - 240V, 50Hz In-built rechargeable battery Battery with operating time at least 4 hours at 25 ml/hr Automatic switch from AC mains power mode to battery operating mode and vice versa

S/N	Equipment Description	Technical specifications
		Total re-charging time not greater than 6 hours
		Microprocessor controlled programmed syringe pump Programming: Up to 99 steps of speed and time' Time Resolution: 0 to 999 minutes in 1 minute steps Time Resolution can be selected individually for each program step Accuracy: +- 1% Reproducibility: +/-0.2%Syringes: Glass, plastic. Metal syringes from 5 micro-liter to over 150ml Flow Rate Range: Depends on the inner syringe diameter 0.4nl/min with a 5ml syringe to 110ml/min with a 150ml syringe Maximum Force: 300N reducible by a switch to 80N Motor: Microprocessor controlled brushless long life BLDC motor with Neodymium magnets Transmission: Efficient force transmission by a ball screw with highest mechanical load capacity of 12,000N Pusher Travel: 120mm Pusher Travel Rate:
		Minimum: 0 08 mm/min Maximum: 80 mm/min Speed Control Range: 0 to 999 Non-volatile Memory: Storage of all settings Power Supply: 220 - 240V, 50Hz Interface: RS-485 or RS-232 automatic valve control Remote Control: 0 - 10V Operation Temperature: 0 - 40oC Operation Humidity: 0 - 90% RH not
	Portable Hand held Pulse oximeter	Patient Range: Adult, Pediatrics and Neonatal patients High Resolution, 2.4" colour display Visual and sound alarms Uses AA size alkaline or rechargeable batteries Digital SpO2 Range: 0 - 100% Resolution: 1%, Accuracy: 70% to 100% +/-2% Refreshing Rate: < 13 seconds Pitch Tone: Yes Pulse Rate Range: 25 - 250 bpm Resolution: 1 bpm, Accuracy: +-2% or +/-1 bpm Refreshing Rate: < 13 seconds Display: Type: 2.4" colour display 320 x 240 pixels Parameter Digital SpO2, Pulse Rate, Pleth bar and SpO2 waveform
		Alarm Audible alarm, audible button tone Supports Pitch Tone and multi-level volume General Description Modular and Suitable for Adult/Paediatric/ Patient monitoring Minimum 15" multi colour TET display screen Eight Channel digital and waveforms/traces display Capability of storage of patient data and printing of patient reports. Five years comprehensive Maintenance inclusive.
	Central Monitorin g Station 8 Channel	Parameters: Eight digital and waveforms/traces display Facilities to monitor and display ECG, Respiration, NIBP, Et CO2, SpO2 and Temperature ECG Multichannel (uo to 12 lead) ST segment analysis 3 or 5 lead with cascade waveform facility Monitoring, Diagnostic and OT modes of monitoring of ECG Simultaneous Multi-lead ECG monitoring of 7 ECG lead Heart Rate (HR) Range: 2 - 350 bpm HR/PR Source selection facility from Automatic, SpO2 IBP and NIBP Automatic arrhythmia detection alarm for standard and lethal arrhythmia Pulse Oximetry Nellcor or Masimo technology Display of Plethysmograph with Pulse Strength indicator and SpO2 Nellcor or Masimo technology Display of Plethysmograph with Pulse Strength indicator and SpO2
		and Neonate Use selectable alarm settings: Mode: Manual, STAT (continuous 5 minute operation) and automati (selectable time interval 2 - 90 minutes) Range: 20 - 250 mmHg Temperature Two channel and with two units (oC and o F) selectable Temperature Range: 0 - 50oC
		Option for differential temperature

S/N	Equipment	Technical specifications
	Description	Should be provided Respiration: RR range 1 - 150 bpm Apnea alarms should be provided Trends and Alarms 72 hours
		Non volatile graphical/ tabular trends with zoom facility and separate dedicated trend for storing minimum200 NIBP readings Should have Alarm recall facility for last 24 Alarm events with date,
		Should have facility to print Graphical trend, tabular trend and alarm recall
		Should have facility to print Graphical trend, tabular trend and diameters
	1	Recorder In-built dual channel thermal array recorder
		Include Laser Printer and dual channel strip chart recorder Others Defibrillator and Cautery protection should be provided Should work on Mains as well as Battery (backup for 2 hours) Automatic zoom in facility in the monitor display
		Should have facility to download trend data on USB and SD Card Supplied complete with all standard accessories and consumables that will enable the machine to operate
		7 C 1 220 240V 50 Hz
		CENTRAL STATION Central Station should have facility to display up to 20 leaf time waves at a
		Central Station should have separate patient window for viewing detailed real-time of stored data to individual patient
		It should have 24 hr stored patient data monitoring trends and 24 hr event review facility
		The Central Nurse Station should also have the following: - Multi-lead arrhythmia and ST review facility - 50 alarms strips storage per bed - Offer wave review with 24 hr full disclosure - Optional facility for dual display for detailed analysis of individual bed without compromising on full ICU monitoring
		Facility for interfacing Holter data for analysis, in case of the Holter from the same brand is available
		Export the ICU Patient data to Holter for analysis
		Remote display (Slave) facility should be available
		Real time recording thru Dual Channel Recorder
		Facility for interfacing a laser printer for printing patient information and trend formats
		Advanced arrthythmia analysis package of at least 20 arrthythmia analysis
		12 lead ECG Monitoring and view possible at Central Nurse Station
		Continuous full disclosure of up to 4 configurable waves per patient
		Operate on Microsoft Windows NT workstation operating system
		Supplied with 19" Flat Screen TFT display, Laser Printer and Recorder, UPS
		- The entire networking and cabling with hardware Wall mounts
	Central Vernous catheters (CVC)	Power Supply: 220 - 240 V, 50Hz Set of Sizes 3, 4, 5, 6, 7, 8, 9, 10
	Orophary ngeal Air way (OPA	Set of Sizes 0,1, 2, 3, 4, 5 for Adult and Pediatrics
	Nasalphar yngeal Air	Set of Sizes 3, 4, 5, 6 for Adults and Pediatrics

S/N	Equipment Description	Technical specifications
	Way (NAP)	
	Endotrach	Set of Sizes 3, 3.5, 4, 5, 6, 6.5, 7, 7.5, 8
<u> </u>	ea	Should be a Low Energy Biphasic defibrillator monitor with Recorder, having capability to deliver shocks from 2 Joules to 200Joules. Five years
		comprehensive Maintenance inclusive. Should monitor ECG through paddles, pads and monitoring electrodes and defibrillate through pads and paddles
		for hady impedance for a range of 25 to 150 onms.
		time of loss than 5 seconds for maximum chorsy.
		Should have High resolution more than 8 inch colour display for viewing meaning parameters like ECG, SpO2, NIBP and EtCO2 with 4 waveform capability of 4
		seconds Both Adult and Pediatric paddles should be available Should have event summary facility for recording and printing at least 55 events
		the of usage for at least 5 hours of monitoring
	defilbrillat	Should be capable of printing Reports on event summary, configuration, sen test, easily a sent set up function.
		Should have facility to monitor parameters like SpO2, NIBP and etCO2 along with non invasive
	_	Should be able to upgrade the defibrillator for 12 lead ECG monitoring and ECG transmission
		System Configuration Accessories, spares and consumables:
		Defibrillator with AED and external Pacemaker - 1pc Adult with built in Pediatric External Pacemaker - 1pc Adult with built in Pediatric External Pacemaker - 1pc Adult NIBP Cuff and Hos 1pc Patient Cables - 1pc ECG Rolls 50 Adult SpO2 reusable Sensor - 1pc Adult NIBP Cuff and Hos
		- 1pc EtCO2 Tubing (box of 20) -1box AED Multifunction Pads for Adults - 10 pairs with Each unit.
		The unit shall be capable of operating continuously in ambient temperature of 3 to 430°C and relative
		humidity of up to 95% Shall meet General Requirements of safety for Electromagnetic Compatibility.
		December 240 VAC 50Hz
8		dividually wrapped device. Markings: at 20
	ea Tube Introduc rs (Styla	cm, 30 cm, and 40 cm intervals. It consists of a 50 to 60 cm stylet with the distance of the stylet with the s
	and	- size 6 FR, OD 2.0 mm, 530 mm
	Bougie)	Made from stainless steel and can be disinfected and sterilized Twin-bladed tong-
	Magills Forceps	like forceps Handles for gripping by the user Rounded ends for grasping conque

/N1	Equipment	Technical specifications
S/N	Description	Nasal cannula (nasal prongs), device designed for easy administration of oxygen into the patient nose through two small prongs placed in the nostrils. It consists of soft twin prongs nasal tips to ensure equal oxygen flow to both. Adjustable, smoothly finished nasal tips for maximum patient comfort Star lumen main tube to avoid accidental blockage. Made from soft and kink resistant polyvinyl chloride (PVC) material For Adult and Child use
	Wall mounted otoscope	(PVC) material For Adult and Child use Illumination: Pressurized halogen lamp Image Resolution: 1280 x 1024 pixels Connection/Interface: USB 2.0 with 3m cable System Requirements: Windows XP Service Pack Electrical Rating: Lamp 3.5V DC, 810mA USB: 5V DC, 150mA Focal length:114mm and Magnification:2.2X Constructed from ABS and Acrylic Plastic or Stainless Steel Automatic Image Brightness and white balancing for optimal image
	-	2 1: 1 walete with Charger
	wall mounted Opthalmo scope	Range of lenses not smaller than -20D to +29D with steps not greater than 1D. Anti- reflection lens Magnification: 12 - 15X Apertures: Small, Large and Semi Circle, reflection Star Colour Temperature: Cool White in the range 3100 - 5500K Light Fixation Star Colour Temperature: Cool White in the range 3100 - 5500K Light Intensity: 8,000 - 12,000 lux. Dust free sealed optics and aspherical optical system Red-free, Blue, Green and polarization filters Handle with On/Off switch. Scratchproof lens: Glass or Plastic Batteries: AA or rechargeable. Battery life at least 5 hours of use on full charge or fresh batteries Accessories: Battery Charger (a applicable to rechargeable type only) Sufficient light bulbs for 3 years use. Rechargeable Battery 1.5V.
	Instrumer t trolley	Stainless steel shelves and railings Available in full stainless steel powers frame. Stainless steel bottom and top shelves with railing on top shelf Frame Mounted on 100 mm diameter wheels Overall Size: L 710 mm x W460 mm x H830
	100	Made of stainless steel with removable top tray
	Adjustabl e Mayo table	Height adjustable by foot 75 mm swivel castors Dimensions: Width 60 cm, Depth 40 cm and Height 120 - 130 cm
	IV Pole	 Five-claw wheel Adjustable IV Pole • Material: Cast iron base 5 Wheels.• Down Tube: 25.• Top Tube: φ 16.• Adjustment mode: Rotary Knob / Jackscrew.• Height: 2m, 2 hooks

S/N E	quipment escription	
	90.	 Dimensions(mm): 850x500x39mm Film: 14 x 17" View area: 730x442mm Lamp: CCFL 12pcs Luminance: 12,000lux or more Power consumption: 60W Power: 220VAC 50/60HZ Life Time: over 80,000hr Film Clipping: Spring Roller or similar
	Digital Mobile X- ray	This Machine should be able to provide a minimum of 300 images per day. Should high frequency generator of 50-150KW, Automatic exposure device, Anatomical programming radiography, overloading protection feature, digital display of KV and mAs. X-Ray tube should be Floor stand mounted. Axis should rotate 360 degree Five years comprehensive Maintenance inclusive. High speed rotating anode and exposure should be 50 – 150KV and 0-600mA. Heat strength capacity of the anode at least 150,000HU. The digital detector (Two fixed Detectors) should be flat panel of latest technology. The digital workstation should have high speed processors, preview time of 5s or less. The workstation should provide basic functions for image processing and be compatible to other HIS Displayed Parameters: The console monitor should display patient ID, Exposure factors,



S/N	Equipment	Technical specifications
	Description	
		Electrical Requirements: Should meet Tanzania Electrical Standards (Voltage of between 220-240V with the standard frequency of 50Hz) with type G adaptor System. Accessories: Multi-tray printer for different X-Ray film sizes (10 x 12, 14 x 14, 14 x 17, 10 x 8). Protective gear (lead apron minimum of 2 small, 2 medium and 2 large), Googles Protective gear (lead apron minimum of 1 small, 1 medium and 2 small, 2 medium 2 small, 2 medium, 2 (Minimum 1 small, 1 medium and 1 large). Gonad shield (minimum 2 small, 2 medium, 2 large), neck collar shield (minimum of 1 small, 1 medium, 1 large), Gloves (Minimum 2 small, 2 medium and 2 large). Radiation hazards warning signs to be supplied with the small, 2 medium and 2 large). Radiation hazards warning signs to be supplied with the small, 2 medium and 2 large). Radiation protective gears: X-ray should come with;(i) two (2) not less than 15 Minutes. Radiation protective gears: X-ray should come with;(i) two (2) pairs of lead aprons with back protection 0.35/0.25mmPb, (ii) two (2) pairs of Thyroid pairs of lead aprons with back protection 0.35/0.25mmPb, (ii) two (2) pairs of Thyroid Shield Model Classic 0.35mmPb (iii) two (2) pairs of Patient Apron with belt 0.50mmPb Shield W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 x 12, 14 Leadlite W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 x 12, 14 Leadlite W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 x 12, 14 Leadlite W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 v 12, 14 Leadlite W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 v 12, 14 Leadlite W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 v 12, 14 Leadlite W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 v 12, 14 Leadlite W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 v 12, 14 Leadlite W/G-30 L-30PrinterMulti-tray printer for different X-Ray film sizes (10 v 12, 14 Leadlite W/G-3
		day of the acceptance testing of the machine. Should be available throughout the lifetime period of the machine. Uptake time should be a minimum of 90%. Proof of locally available technical support personnel, including CVs and work permit for foreign personnel. Availability of technical personnel within the country should be stated; this should include CVs, work permits for foreign personnel. Software should be flexible and provide the room for upgrade to add new parameters to be measured by the Machine and report format. Documentation: Operating and service manuals (In English) including lists of important spares and accessories - with their part numbers and list of equipment and procedures required for calibration and routine maintenance should be provided. Documentation must also show recommended procedures for disposal and any probable hazards to the environment and/or community. Life span: Life span of the machine should be not less than 10 Years. Risk Classification: As per ISO 14971:2007- Application of risk management to medical devices. Regulatory Approval / Certification: TAEC, TBS and TFDA



51(512)	Equipment Description	Technical specifications
	Neonatal ventlilator	The Ventilator should be microprocessor controlled designed for neonatal use. There should be possibility to upgrade with additional features Continuous flow, pressure limited, time cycled ventilator design Ventilator modes: should have following modes available in the unit IMV/IPPV CPAP including non-invasive ventilation SIMV, SIPPV/Assist-control Volume targeted/guarantee mode of ventilation with ability to deliver and monitor tidal volume as low as1-2ml (Range 2ml to 50 ml) Pressure support mode of ventilation Apnea back-up ventilation Should have integrated high resolution TFT medical grade screen (size 4") for real time display Scalar (Pressure, Flow and Volume against time) and loop (Pressurevolume, volume flow and pressure-flow) pulmonary graphics Digital display of FiO2, Peak pressure, mean airway pressure, CPAP/PEEP, Expiratory tidal volume, expiratory minute volume, total frequency, spontaneous frequency, lung function monitoring Compliance, resistance, lung distention coefficient, (C20/C), Lung time constant, Rate volume ratio etc. Should have built-in logbook for recording events like various alarms. Integrated monitoring: Integrated volume and pressure monitoring i.e. monitoring of PEEP Pmax, Pmean and VT, VTspont, MV and MVleak. The volume monitoring should have NTPD to BTPS correction. Monitoring of ie, Frequency and Spont. Frequency Settings range: Trigger Flow/volume, leak adapted PIP 8 to 80cm of water PEEP/CPAP 0 to 25 mbar. Five years comprehensive Maintenance inclusive. I.E ratio I:0 to 1:10 Inspired Time 0.1 to 2 sec Expired Time 0.2 to 30 sec Frequency Up to 200 BPM Base Flow (VIVE) 1 to 30 LPM Patient synchronization Integrated bender for Oxygen 21 % to 100% Integrated nebulization facility Integrated screen for display of Pressure-Time, Flow-Time and Volume-Time curves Integrated monitoring of FiO2 Monitoring of flow: At the Y piece with facility to activate it Audiovisual alarms with advisory onscreen message: MV high/low, Apnea, tube obstruction, FiO2, high/low high PIP, low PEEP/CPA
0	Glucomet	·Hand held type Glucometer ·Battery operated ·Memory up to 10 measurements ·Sticks method measurements ·Code of sticks interring is available ·Indication of high and low measurements One



S/N	Equipment	Technical specifications
	Hemoglob inometer	· Parameter: Hemoglobin, HCT(Hematocrit)· Principle: Optical reflectance · Sample: Capillary or venous whole blood (13~15ul) · Strip: H12 Hemoglobin Test Strip · Speed: ≤ 10 seconds · Measuring Range: 4.0g/dL~25.6g/dL(g/L optional) · Memory: 1200 test results · Automatization: Self-checking, auto-judging and displaying malfunction · Power: DC 6V(Two CR2032 Batteries) · Precision: CV<5%,testing samples including high, medium & low concentration, the CV of 10
	Patient Beds with matress,	• Heavy duty bed Size (L×W×H): 2150×950×500mm• High quality section steel welding• The surface electrostatic sprayed• Corrosion resistant• Easy to clean• Side guard made of well plastic• I.V pole• Four casters, two caster with brakes
	Bedside lockers	 dimension 520*480*740 made of high density wood one drawer and one door with handles, top with rail two tower hanger, four wheels with cross breaks
	Cardiac table	 Size:900*450*900-1200mm Wooden dinning board(800*400mm) Steel frame, ABS gas spring bottom Height is adjustable, controlled by gas-spring Four silent castors,φ55mm

SPECIAL CONDITIONS OF CONTRACT

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Special Conditions of Contract (SCC)

The following Special Conditions of Contract (SCC) shall supplement the GCC. Whenever there is a conflict, the provisions herein shall prevail over those in the GCC. The corresponding clause number of the GCC is indicated in parentheses.

SCC Clause Number	GCC Clause Number	Amendments of, and Supplements to, Clauses in the GCC
	Definition	ons (GCC 1)
1.	1.1	The Purchaser is: Ocean Road Cancer Institute Luthuli/Samora avenue, P.o Box 3592 Tel. +255 22 2127597 Fax 255-22-2118704 Dar Es Salaam
2.	1.1(j)	The Supplier is: ANUDHA LIMITED of P. o. Box 5982. Plot 2169/82& 8170/82, Morogoro Road, Opp. DART Kisutu Bus stand, DAR ES SALAAM TANZANIA
3.	1.1(q)	The Project is: SUPPLY OF ICU EQUIPMENTS AND CONSUMABLES FOR ORCI, MTWARA ZONAL, LIGULA AND MOROGORO RHH
	Governin	g Language (GCC 4)
4.	4.1	The Governing Language shall be: ENGLISH
	Applicabl	e Law (GCC 5)
5.	5.1	The Applicable Law shall be: Laws of Tanzania.
	Country of	f Origin (GCC 6)
ó.	6.1	Country of Origin is all countries and territories as indicated in the section of the Tendering Documents, Eligibility for the Provision of Goods, Works and Services.

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	Perform	mance Security (GCC 10)		
7.	10.1	The amount of performance security, as a percentage of the Contract Price, shall be: \\ \{__\)10 percent of the Contract Price		
8.	10.4	After delivery and acceptance of the Goods, the performance security shall be reduced to two (2) percent of the Contract Price to cover the Supplier's warranty obligations in accordance with GCC 18.2.		
	Inspect	tions and Tests (GCC 11)		
9.	11.1	Inspection and tests prior to shipment of Goods and at final acceptance are as follows:		
		Quality and quantity inspection shall be carried out prior to shipment of Goods by the manufacturer(s) at the supplier's own expense and responsibility in terms of the items specified in the specifications. The supplier shall submit the inspection certificate issued by himself which should be attached with the certificate(s) of the manufacturer(s) to the PE in order to ensure that the goods are manufactured in compliance with the contract.		
	Packing	g (GCC 12)		
10.	12.2	The Goods shall be packed properly in accordance in a second state of the control		
- 1 - 2 - 2 - 2 - 2 - 2 - 2 - 2 - 2 - 2		The Goods shall be packed properly in accordance with standard export packing specified by the PE in the Technical Specification.		
	Delivery and Documents (GCC 13)			
11.	13.1	For Goods supplied from abroad: Upon shipment, the Supplier shall notify the Purchaser and the Insurance Company by cable the full details of the shipment,		
		including Contract number, description of Goods, quantity, the vessel, the bill of lading number and date, port of loading, date of shipment, port of discharge, etc. The Supplier shall mail the following documents to the Purchaser, with a copy to the Insurance Company:		
		(i.)One original plus four copies of the Supplier's invoice showing Goods' description, quantity, unit price, and total amount;		



		 (ii.)original and four copies of the negotiable, clean, on-board bill of lading marked "freight prepaid" and four copies of nonnegotiable bill of lading; (iii.)One original plus four copies of the packing list identifying contents of each package; (iv.)insurance certificate; (v.)Manufacturer's or Supplier's warranty certificate; (vi.)inspection certificate, issued by the nominated inspection agency, and the Supplier's factory inspection report; and (vii.)certificate of country of origin issued by the chamber of commerce and industry or equivalent authority in the country of origin in duplicate The above documents shall be received by the PE at least one week before arrival of the Goods at the port or place of arrival and, if not
		before arrival of the Goods at the port of place of difficulty and received, the Supplier will be responsible for any consequent expenses. [Other similar documents should be listed, depending upon the Incoterm retained.]
12.	13.3	For Goods from within the United Republic of Tanzania:
		Upon delivery of the Goods to the transporter, the Supplier shall notify the PE and mail the following documents to the PE:
		(i.) one original plus four copies of the Supplier's invoice showing Goods' description, quantity, unit price, and total amount;
		(ii.) delivery note, railway receipt, or truck receipt;
		(iii.) Manufacturer's or Supplier's warranty certificate;
		(iv.) inspection certificate issued by the nominated inspection agency, and the Supplier's factory inspection report; an
		(v.) certificate of country of origin issued by the Tanzan Chamber of Commerce, Industry and Agriculture of equivalent authority in the country of origin in duplicat

		The above documents shall be received by the PE before arrival of the Goods and, if not received, the Supplier will be responsible for any consequent expenses.
	Insuranc	e (GCC 14)
13.	14.1	The Insurance shall be in an amount equal to 110 percent of the DDP value of the Goods from "warehouse" to "warehouse" on "All Risks" basis, including War Risks and Strikes.
	Inciden	tal Services (GCC 16)
14.	16.1	Incidental services to be provided are: [Selected services covered under GCC 16 and/or other should be specified with the desired features. The price quoted in the tender price or agreed with the selected Supplier shall be included in the Contract Price.]
	Spare 1	Parts (GCC 17)
15.	17.1	Additional spare parts requirements are: Supplier shall carry sufficient inventories to assure ex-stock supply of consumable spares for the Goods. Other spare parts and components shall be supplied as promptly as possible, but in any case within six (6) months of placing the order and opening the letter of credit.
	Warr	anty (GCC 18)
16.	18.2	GCC 17.2—In partial modification of the provisions, the warranty period shall be hours of operation or months from date of acceptance of the Goods or () months from the date of shipment, whichever occurs earlier. The Supplier shall, in addition, comply with the performance and/or consumption guarantees specified under the Contract. If, for reasons attributable to the Supplier, these guarantees are not attained in whole or in part, the Supplier shall, at its discretion either:
		(a) make such changes, modifications, and/or additions to the Goods or any part thereof as may be necessary in order to attain the contractual guarantees specified in the Contract at its own cost and expense and to carry out further performance tests in accordance with SCC 4,
		or



		 (b) pay liquidated damages to the PE with respect to the failure to meet the contractual guarantees. The rate of these liquidated damages shall be 0.20 per cent per day of undelivered materials/goods value up to the sum equivalent to the amount of ten percent of the contract value. Five (5) years comprehensive Maintenance
17.	18.4 & 18.5	The period for correction of defects in the warranty period is: Five (5) years comprehensive Maintenance
	Paymen	The method and conditions of payment to be made to the Supplier
18.		under this Contract shall be as follows: Payment for Goods supplied from abroad: Payment of foreign currency portion shall be made in
		(i) Advance Payment: 65 percent of the Contract Price shall be paid within thirty (30) days of signing of the Contract, and upon submission of claim and a bank guarantee or insurance bond for equivalent amount valid until the Goods are delivered and in the form provided in the Tendering Documents or another form acceptable to the PE.
		(ii) On Shipment: 30 percent of the Contract Price of the Goods shipped shall be paid through irrevocable confirmed letter of credit opened in favor of the Supplier in a bank in its country, upon submission of documents specified in GCC 10.
		(iii) On Acceptance: 5 percent of the Contract Price of Goods received shall be paid within thirty (30) days or receipt of the Goods upon submission of clair supported by the acceptance certificate issued by the PE.

		Documents or another form acceptable to the PE. (ii) On Delivery: 30 percent of the Contract Price shall be
		(ii) On Delivery: 30 percent of the Contract Price shall be paid on receipt of the Goods and upon submission of the documents specified in GCC 11.
		(iii) On Acceptance: The remaining 5 percent of the Contract Price shall be paid to the Supplier within thirty (30) days after the date of the acceptance certificate for the respective delivery issued by the PE.
19.	19.3	Rate to be used for paying the Supplier's interest on the late paymen made by PE shall be [insert: rate] N/A
	Prices (GCC 20)
20.	20.1	Prices shall be adjusted in accordance with provisions in the
		Attachment to SCC.
-9100-		The price shall be fixed
	Liquida	ated Damages (GCC 26)
	-	

		Maximum deduction: is equal to the performance security. Note: 0.1 to 0.2 per cent per day of undelivered materials/good's value.
	Proced	ure for Dispute Resolution (GCC 32)
23.	32.3	Arbitration institution shall be TANZANIA INSTITUTE OF ARBITRATION Place for carrying out Arbitration TANZANIA INSTITUTE OF ARBITRATION
24.	33.1	Appointing Authority for the Adjudicator TANZANIA INSTITUTE OF ARBITRATION
	Notices	s (GCC 35)
26.	35.1	-PE's address for notice purposes: Executive Director, Ocean Road Cancer Institute Luthuli/Samora avenue, P.o Box 3592 Tel. +255 22 2127597 Fax 255-22-2118704 Dar Es Salaam - through ONLINE- TANePS (Tanzanian National e-Procurement System)
		-Supplier's address for notice purposes:- ANUDHA LIMITED of P. o. Box 5982. Plot 2169/82& 8170/82, Morogoro Road, Opp. DART Kisutu Bus stand, DAR ES SALAAM TANZANIA

GENERAL CONDITIONS OF THE CONTRACT (GCC)

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GENERAL CONDITIONS OF THE CONTRACT (GCC)

1.	Definitions	1.1	Th	e following words and expressions shall have the canings hereby assigned to them:
			a)	The Adjudicator is the person appointed by the appointing authority specified in the Special Conditions of Contract (SCC), to resolve contractual disputes in the first instance, and as provided for in General Conditions of the Contact (GCC) 31 hereunder.
			b)	The Arbitrator is the person appointed by the appointing authority specified in the SCC, to resolve contractual disputes.
			c)	"The Contract" means the agreement entered into between the Procuring Entity (PE) and the Supplier, as recorded in the Contract Form signed by the parties, including all attachments and appendices thereto and all documents incorporated by reference therein.
			d)	The Commencement Date is the date when the Supplier shall commence execution of the contract as specified in the SCC
			e)	"Completion" means the fulfillment of the related services by the Supplier in accordance with the terms and conditions set forth in the contract
			f)	The Contract Price is the price stated in the Letter of Acceptance and thereafter as adjusted in accordance with the provisions of the Contract Days are calendar days
			g	A Defective Goods are those goods which are below standards, requirements or specifications stated by the Contract.
			h)	"Delivery" means the transfer of the goods from the supplier equipment, machinery, and /or other materials which the Supplier is required to supply to the PE under Contract.
			i)	"Effective Contract date" is the date shown in the Certificate of Contract Commencement issued by the Employer upon fulfillment of the conditions precedent stipulated in GCC 3.
			j)	"The Purchaser" means the person named as purchaser in the SCC and the legal successors in title to this person



k)	"The Related Services" means those services ancillary to the supply of the Goods, such as transportation and insurance, and any other incidental services, such as installation, commissioning, provision of technical assistance, training, initial maintenance and other such obligations of the Supplier covered under the Contract.
1)	"GCC" means the General Conditions of Contract contained in this section.
m)	The Intended Delivery Date is the date on which it is intended that the Supplier shall effect delivery as specified in the SCC
n)	"SCC" means the Special Conditions of Contract.
0)	"The PE" means the entity purchasing the Goods and related service, as named in SCC.
p)	"The Supplier" means the individual private or government entity or a combination of the above whose Tender to perform the contract has been accepted by the PE and is named as such in the Contract Agreement and includes the legal successors or permitted assigns of the supplier and shall be named in the SCC.
q)	"The Project Name" means the name of the project stated in SCC.
r)	"Day" means calendar day.
s)	'Eligible Country" means the countries and territories eligible for participation in procurements financed by the specified institution.
t)	"End User" means the organization(s) where the goods will be used, as named in the SCC.
u)	"Origin" means the place where the Goods were mined, grown, or produced or from which the Services are supplied. Goods are produced when, through manufacturing, processing, or substantial and major assembly of components, a commercially recognized new produce results that is substantially different in basic characteristics or in purpose or utility from its components.
v)	"Force Majeure" means an unforeseeable event which is beyond reasonable control of either Party and which makes a Party's performance of its obligations under the



2.	Application and interpretation	2.1	These General Conditions shall apply to the extent that they are not superseded by provisions of other parts of the Contract. In interpreting these Conditions of Contract headings and marginal notes are used for convenience only and shall no affect their interpretations unless specifically stated references to singular include the plural and vice versa; and masculine include the feminine. Words have their ordinary meaning under the language of the Contract unless specifically defined.
			Contract impossible or so impractical as to be considered impossible under the circumstances. For the purposes of this Contract, "Force Majeure" means an event which is beyond the reasonable control of a Party, is not foreseeable, is unavoidable, and its origin is not due to negligence or lack of care on the part of a Party, and which makes a Party's performance of its obligations hereunder impossible or so impractical as reasonably to be considered impossible in the circumstances. and includes, but is not limited to, war, riots, civil disorder, earthquake, fire, explosion, storm, flood, epidemics, or other adverse weather conditions, strikes, lockouts or other industrial action (except where such strikes, lockouts or other industrial action are within the power of the Party invoking Force Majeure to prevent), confiscation or any other action by Government agencies. Specification means the Specification of the Works included in the Contract and any modification or addition made or approved by the Project Manager. The Supplier is the person, whether natural or legal whose Tender to deliver goods or services has been accepted by the Employer The Supplier's Tender is the completed Tender document submitted by the Supplier to the Employer



2.3	The documents forming the Contract shall be interpreted in the following order of priority:
	(1) Form of Contract,
	(2) Special Conditions of Contract,
	(3) General Conditions of Contract,
	(4) Letter of Acceptance,
	(5) Certificate of Contract Commencement
	(6) Specifications
	(7) Contractor's Tender, and
	(8) Any other document listed in the Special Conditions of Contract as forming part of the Contract.
	2.3



3.	Conditions Precedent	3.1	 Having signed the Contract, it shall come into effect on the date on which the following conditions have been satisfied: a) Submission of performance Security in the form specified in the SCC; b) Furnishing of Advance Payment Unconditional Guarantee.
		3.2	If the Condition precedent stipulated on GCC 3.1 is not met by the date specified in the SCC this contract shall not come into effect;
		3.2	If the Employer is satisfied that each of the conditions precedent in this contract has been satisfied (except to the extent waved by him, but subject to such conditions as he shall impose in respect of such waiver)he shall promptly issue to the supplier a certificate of Contract commencement, which shall confirm the start date.
4.	Governing Language	4.1	The Contract as all correspondence and documents relating to the contract exchanged by the Supplier and the PE shall be written in the language specified in SCC. Subject to GCC 3.1, the version of the Contract written in the specified language shall govern its interpretation.
5.	Applicable Law	5.1	The contract shall be governed and interpreted in accordance with the laws of the United Republic of Tanzania, unless otherwise specified in SCC.
6.	Country of Origin	6.1	The origin of Goods and Services may be distinct from the nationality of the Supplier.



7.	Standards	7.1	The Goods supplied under this Contract shall conform to the standards mentioned in the Technical Specifications, and, when no applicable standard is mentioned, to the authoritative standards appropriate to the Goods' country of origin. Such standards shall be the latest issued by the concerned institution.
8.	Use of Contract Documents and Information; Inspection and Audit by the Government of Tanzania	8.1	The Supplier shall not, without the PE's prior written consent, disclose the Contract, or any provision thereof, or any specification, plan, drawing, pattern, sample, or information furnished by or on behalf of the PE in connection therewith, to any person other than a person employed by the Supplier in the performance of the Contract. Disclosure to any such employed person shall be made in confidence and shall extend only as far as may be necessary for purposes of such performance.
		8.2	The Supplier shall not, without the PE's prior written consent, make use of any document or information enumerated in GCC 7.1 except for purposes of performing the Contract.
		8.3	Any document, other than the Contract itself, enumerated in GCC 7.1 shall remain the property of the PE and shall be returned (all copies) to the PE on completion of the Supplier's performance under the Contract if so required by the PE.
		8.4	The Supplier shall permit the Government of the United Republic of Tanzania or / and donor agencies involved in financing the project to inspect the Supplier's accounts and records relating to the performance of the Supplier and to have them audited by auditors appointed by the Government of the United Republic of Tanzania or / and the appropriate donor agencies, if so required by the Government of the United Republic of Tanzania or / and the appropriate donor agencies.
9.	Patent and Copy Rights	9.1	The Supplier shall indemnify the PE against all third-party claims of infringement of patent, trademark, or industrial design rights arising from use of the Goods or any part thereof in the United Republic of Tanzania.
		9.2	The patent right in all drawings, documents, and other materials containing data and information furnished to the PE by the Supplier herein shall remain vested in the supplier, or, if they are furnished to the PE directly, or through the Supplier by any third party, including suppliers of materials, the patent right in such materials shall remain vested in such third party.



0.	Performance Security	10.1	no later than the date specified in the Ected of an and shall be issued in an amount and form and by a bank or surety acceptable to the Employer, and denominated in the types and proportions of the currencies in which the Contract Price is payable as specified in the SCC.		
	*	10.2	The proceeds of the performance security shall be payable to the PE as compensation for any loss resulting from the Supplier's failure to complete its obligations under the Contract.		
		10.3	The performance security shall be in one of the following forms:		
			a) A bank guarantee, an irrevocable letter of credit issued by a reputable bank, or an insurance bond issued by a reputable insurance firm located in the United Republic of Tanzania or abroad, acceptable to the PE, in the form provided in the Tendering Documents or another form acceptable to the PE; or		
-			b) A cashier's or certified check.		
		10	The performance security will be discharged by the PE and returned to the Supplier not later than thirty (30) days following the date of completion of the Supplier's performance obligations under the Contract, including any warranty obligations, unless otherwise specified in SCC.		
-		10	0.5 Where circumstances necessitate the amendment of the contract after signature, and such amendment is effected, the PE share require the Supplier to provide additional Performance Securit to cover any cumulative increase of more than ten percent of the initial Contract Price.		



11.	Inspections and Test	11.1	The PE or its representative shall have the right to inspect and /or to test the Goods to confirm their conformity to the Contract specifications at no extra cost to the PE. SCC and the Technical Specifications shall specify what inspections and tests the PE shall notify the Supplier in writing or in electronic forms that provide record of the content of communication, in a timely manner, of the identity of any representatives retained for these purposes.
		11.2	The inspections and tests may be conducted on the premises of the Supplier or its subcontractor(s), at point of delivery, and/or at the Goods' final destination. If conducted on the premises of the Supplier or its subcontractor(s), all reasonable facilities and assistance, including access to drawings and production data, shall be furnished to the inspectors at no charge to the PE.
		11.3	Should any inspected or tested Goods fail to conform to the Specifications, the PE may reject the Goods, and the Supplier shall replace the rejected Goods to meet specification requirements free of cost to the PE.
		11.4	The PE's right to inspect, test and, where necessary, reject Goods after the Goods' arrival in the PE's country shall in no way be limited or eared by reason of the Goods having previously been inspected, tested, and passed by the PE or its representative prior to the Goods' shipment from the country of origin.
		11.5	Nothing in GCC 10 shall in any way release the supplier from any warranty or other obligations under this Contract.
12.	Packing	12.1	The supplier shall provide such packing of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in the Contract. The packing shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt and precipitation during transit, and open storage. Packing case size and weights shall take into consideration, where appropriate, the remoteness of the Goods final destination and the absence of heavy handling facilities at all points in transit.
		12.2	The packing, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the Contract, including additional requirements, if any, specified in SCC, and in any subsequent instructions ordered by the PE.



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3.	Delivery and Documents	13.1	Delivery of the Goods shall be made by the Supplier in accordance with the terms specified in the Schedule of Requirements. The details of shipping and or other documents to be furnished by the Supplier as specified in SCC .
		13.2	For purposes of the Contract, "EXW" "FOB" "FCA", "CIF", "CIP," and other trade terms used to describe the obligations of the parties shall have the meanings assigned to them by the current edition of INCOTERMS published by the International Chamber of Commerce, Paris.
		13.3	Documents to be submitted by the Supplier are specified in SCC.
14.	Insurance	14.1	The Goods supplied under the Contract shall be fully insured in a freely convertible currency against loss or damage incidental to manufacture or acquisition, transportation, storage, and delivery in the manner specified in the SCC.
15.	Transportation	15.1	Where the Supplier is required under Contract to deliver the Goods FOB, transport of the Goods, up to and including the point of putting the Goods on board the vessel at the specified port of loading, shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price. Where the Supplier is required under the Contract to deliver the Goods FCA transport of the Goods and delivery into the custody of the carrier at the place named by the PE or other agreed point shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price.
		15.2	Where the Supplier is required under Contract to deliver the Goods CIF or CIP, transport of the Goods to the port of destination or such other named place of destination in the United Republic of Tanzania, as shall be specified in the Contract, shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price.
		15.3	Where the Supplier is required under the Contract to transport the Goods to a specified place of destination within the Unite Republic of Tanzania, defined as the Project Site, transport to such place of destination in the United Republic of Tanzania, including insurance and storage, as shall be specified in the Contract, shall be arranged by the Supplier, and related costs shall be included in the Contract Price.
10	6. Incidental Services	16.	The Supplier may be required to provide any or all of the following services, including additional services, if any, specific in SCC:
			a) Performance or supervision of on-site assembly and/ start-up of the supplied Goods;



			b)	Furnishing of tools required for assembly and/or maintenance of the supplied Goods;			
			c)	Furnishing of a detailed operations and maintenance manual for each appropriate unit of the supplied Goods;			
			d)	Performance or supervision or maintenance and/or repair of the supplied Goods, for a period of time agreed by the parties, provided that this service shall not relieve the Supplier of any warranty obligations under this Contract; and			
			e)	Training of the PE's personnel, at the Supplier's plant and/or on-site, in assembly, start-up, operation, maintenance, and/or repair of the supplied Goods.			
		16.2	inclu in ac	es charged by the Supplier for incidental services, if not uded in the Contract Price for the Goods, shall be agreed upon dvance by the parties and shall not exceed the prevailing rates ged to other parties by the Supplier for similar services.			
17.	Spare Parts	17.1	or a	As specified in SCC, the Supplier may be required to provide any or all of the following materials, notifications, and information pertaining to spare parts manufactured or distributed by the Supplier:			
			a)	Such spare parts as the PE may elect to purchase from the Supplier, provided that this election shall not relieve the Supplier of any warranty obligations under the Contract and			
			b)	 i) advance notification to the PE of the pending termination, in sufficient time to permit the PE to procure needed requirements; and ii) following such termination, furnishing at no cost to the PE, the blueprints, drawings, and specification of the spare parts, if requested. 			



18. Warranty	and to mate furth have (excessive) con	Supplier warrants that the Goods supplied under the fact are new, unused, of the most recent or current models that they incorporate all recent improvements in design and that they incorporate all recent improvements in design and warrants that all Goods supplied under this Contract shall are warrants that all Goods supplied under this Contract shall are no defect, arising from design, materials, or workmanship ept when the design and/or material is required by the PE, ept when the design and/or or material is required by the PE, develop under normal use of the supplied Goods in the ditions prevailing in the United Republic of Tanzania.
	after beed in of coo ot	en delivered to and accepted at the final destination interacted and elivered to and accepted at the final destination interacted the Contract, or for a period specified in the SCC after the date shipment from the port or place of loading in the source shipment from the port or place of loading in the source shipment from the port or place of loading in the source shipment from the period concludes earlier, unless specified therwise in SCC. The PE shall promptly notify the Supplier in writing or in the lectronic forms that provide record of the content of the communication of any claims arising under this warranty. Then receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall, within the period lipon receipt of such notice, the Supplier shall repeat lipon receipt of such notice, the Supplier shall repeat lipon receipt of such notice, the Supplier shall repeat lipon receipt of such notice shall
	1	the defective Goods of parts dielect, where applicable, the cost of inland delivery of the repaired or replaced Goods or parts from EXW or the port or place of entry to entry to the final destination.
	18.5	If the Supplier, having been notified, fails to remedy the defect(s) within the period specified in SCC, the PE may proceed to take such remedial action as may be necessary, at the Supplier's risk and expense and without prejudice to any other rights which the PE may have against the Supplier under the Contract.
19. Payment	19.1	The method and conditions of payment to be made to the Supplier under this Contract shall be specified in SCC.
	19.2	The Supplier's request(s) for payment shall be made to the PE in writing or in electronic forms that provide record of the content of communication, accompanied by an invoice describing, as appropriate, the Goods delivered and Services performed, and by documents submitted pursuant to GCC 13, and upon fulfillment of other obligations stipulated in the Contract.



		af m la th	ayments shall be made promptly by the PE, within sixty (60) days fer submission of an invoice or claim by the Supplier. If the PE nakes a late payment, the Supplier shall be paid interest on the late payment. Interest shall be calculated from the date by which he payment should have been made up to the date when the late by when the payment is made at the rate as specified in the SCC.
		1	The currency or currencies in which payment is made to the Supplier under this Contract shall be specified in SCC subject to the following general principle: payment will be made in the currency or currencies in which the payment has been requested in the Supplier's Tender.
		19.5	All payments shall be made in the currency or currencies specified in the SCC pursuant to GCC 19.4
20.	Prices	20.1	The contract price shall be as specified in the Contract Agreement Subject to any additions and adjustments thereto or deductions there from, as may be made pursuant to the Contract.
		20.2	Prices charged by the Supplier for Goods delivered and Services performed under the Contract shall not vary from the prices quoted by the Supplier in its Tender, with the exception of any price adjustments authorized in SCC or in the PE's request for Tender validity extension, as the case may be. The PE may at any time, by a written order given to the Supplier
21.	Change Orders	21.1	pursuant to GCC 22, make changes the following:
			a) Drawings, designs, or specifications, where Goods to be furnished under the Contract are to be specifically manufactured for the PE;
			b) The method of shipment or packing;
			c) The place of delivery; and/or
			d) The Services to be provided by the Supplier.
		21	1.2 If any such change causes an increase or decrease in the cost of, of the time required for, the Supplier's performance of an provisions under the Contract an equitable adjustment shall be made in the Contract Price or delivery schedule, or both, and the Contract shall accordingly be amended. Any claims by the Supplier for adjustment under this clause must be asserted within thirty (30) days from the date of the Supplier's receipt of the PE change order.



			any related services that		
$\overline{}$		21.3 Pr	rices to be charged by the supplier for any related services that hight be needed but which were not included in the Contract hall be agreed upon in advance by the Parties and shall not hall be agreed upon in advance by the parties by the		
	ı	Supplier for similar server			
22.	Contract	22.1	Subject to GCC 20, no variation in or modification of the the Contract shall be made except by written amendment signed		
	Amendments	1	by the parties.		
22	Assignment	23.1	Neither the PE nor the Supplier shall assign, in whole or in part, obligations under this Contract, except with the prior written		
23.	Assignment		obligations under this consent of the other party.		
		211	Entity in the event of		
24.	Subcontracts	24.1	The Supplier shall consult the Procuring Entity in the subcontracting under this contract if not already specified in the Tender. Subcontracting shall not alter the Supplier's		
			obligations.		
25	5. Delays in the	24.2	Delivery of the Goods and perform the time schedule prescribed		
25	Supplier's Performance		by the Supplier In accordance by the Procuring Entity in the Schedule of Requirements.		
	Periormance	25.2	If at any time during performance of the Contract, the Supplier or its subcontractor(s) should encounter conditions impeding timely delivery of the Goods and performance of Services, the Supplier shall promptly notify the Procuring Entity in writing or in electronic forms that provide record of the content of communication of the fact of the delay, its likely duration and its cause(s). As soon as practicable after receipt of the Supplier's notice, the Procuring Entity shall evaluate the situation and may at its discretion extend the Supplier's time for performance, with or without liquidated damages, in which case the extension shall be ratified by the parties by amendment of Contract.		
		25.3	Except as provided under GCC 28, a delay by the supplied performance of its delivery obligations shall render the Supplied liable to the imposition of liquidated damages pursuant to GCC 26 unless an extension of time is agreed upon pursuant to GCC 25 without the application of liquidated damages.		



5. I	iquidated Damages	5	of the special prejudent of the special perconduction of the special perco	fied dice tract enta ds con uction to the contract of the cont	GCC Clause 28, if the Supplier fails to deliver any or all Goods or to perform the Services within the period(s) in the Contract, the Procuring Entity shall, without to its other remedies under the Contract, deduct from the Price, as liquidated damages, a sum equivalent to the age specified in SCC of the delivered price of the delayed or unperformed Services for each week or part thereof of antil actual delivery or performance, up to a maximum on of the performance security specified in SCC. Once the aximum is reached, the Procuring Entity may consider ation of the Contract pursuant to GCC Clause 26.
27.	Termination for Default	27.1	ocuring Entity or the Supplier, without prejudice to any other y for breach of Contract, by written notice of default sent to neerned party may terminate the Contract if the other party a fundamental breach of the Contract.		
		27.2	Fi	ında	amental breaches of Contract shall include, but shall not be
			(a)))	the Supplier fails to deliver any or all of the Goods within the period(s) specified in the Contract, or within any extension thereof granted by the Procuring Entity pursuant to GCC Clause 24; or the Supplier fails to perform any other obligation(s) under the Contract;
				c)	Supplier's failure to submit performance security within the time stipulated in the SCC;
-		_		d)	the supplier has abandoned or repudiated the contract.
				e)	The Procuring Entity or the Supplier is declared bankrupt of goes into liquidation other than for a reconstruction of amalgamation;
				f)	a payment is not paid by the Procuring Entity to the Suppli- after 84 days from the due date for payment;
				g)	the Procuring Entity gives Notice that goods delivered with defect is a fundamental breach of Contract and the Suppli fails to correct it within a reasonable period of time determine by the Procuring Entity; and
				h)	if the Procuring Entity determines, based on the reasonable evidence, that the Supplier has engaged corrupt, coercive, collusive, obstructive or fraudule practices, in competing for or in executing the Contractions.



For the purpose of this clause:	
- I	
For the purpose of this	
"corrupt practice means the offering, gi of anything of value to influence the act procurement process or contract execut	pairing or harming, or or indirectly, any party or the of influencing improperly the of influencing improperly the of influencing improperly the of influencing improperly the or harming, or threatening to only part or the property of the improperly the action or a part or the property of the improperly the action or a part or the procurement or government or execution of a contract to the individues collusive practices of includes collusive practices of includes of free and oper of the benefits of free and oper intended to materially impedituent exercising a duty under the exercising a duty under the exercising a duty under the in such manner as it deem



28. Force Majeure	Notwithstanding the provisions of GCC Clauses 25, 26, and 27, neither Party shall have any liability or be deemed to be in breach of the Contract for any delay nor is other failure in performance of its obligations under the Contract, if such delay or failure is a result of an event of Force Majeure. For purpose of this clause, "Force Majeure" means an event which is beyond the reasonable control of a Party, is not foreseeable, is unavoidable, and its origin is not due to negligence or lack of care on the part of a Party, and which makes a Party's performance of its obligations hereunder
	impossible or so impraction impossible in the circumstances, and includes, but is not impossible in the circumstances, and includes, but is not impossible in the circumstances, and includes, but is not impossible or so impraction. In the circumstances, and includes, but is not impossible or so impossible or so includes, but is not impossible or so includes, and includes, but is not impossible or so includes, but is not impossible or so includes, and includes, but is not impossible or so includes, and includes, but is not impossible or so includes, but is not
	28.2 If a Party (hereinafter referred to as "the Affected Party") is or will be prevented from performing its substantial obligation under the contract by Force Majeure, it shall give a Notice to the other Party giving full particulars of the event and circumstance of Force Majeure in writing or in electronic forms that provide record of the content of communication of such condition and the cause thereof. Unless otherwise directed by the Procuring Entity in writing or in electronic forms that provide record of the content of communication, the Supplier shall continuate to perform its obligations under the Contract as far as is reasonable practical, and shall seek all reasonable alternative means for performance not prevented by the Force Majeure event.



9.	for	mination olvency	29.1	giv bar wi ter wl	ing V nkrup thout mina hich h	curing Entity may at any time terminate the Contract by written notice to the Supplier if the Supplier becomes of or otherwise insolvent. In this event, termination will be compensation to the Supplier, provided that such ation will not prejudice or affect any right of action or remedy has accrued or will accrue thereafter to the Procuring Entity.
30.	fo	ermination r onvenience	30.1	te co te is	ermina onver ermin s term effecti	1 i mont within thirty
			30.2		(30) d	Goods that are complete and ready for shipment within thirty lays after the Supplier's receipt of notice of termination shall cepted by the Procuring Entity at the Contract terms and price. The remaining Goods, the Procuring Entity may elect: To have any portion completed and delivered at the Contract terms and prices; and / or
					b)	To cancel the remainder and pay to the Supplier an agreed amount for partially completed Goods and Services and for materials and parts previously procured by the Supplier.
	31.	Disputes Resolution	44	1.1	par dar dis ne to	the event of any dispute arising out of this contract, either rty shall issue a notice of dispute to settle the dispute nicably. The parties hereto shall, within twenty-eight (28) ys from the notice date, use their best efforts to settle the spute amicably through mutual consultations and egotiation. Any unsolved dispute may be referred by party an adjudicator nominated by the appointing Authority becified in SCC.
				31.3	2 A 30 th re-	Ifter the dispute has been referred to the adjudicator, within a days, or within such other period as may be proposed by the Parties, the Adjudicator shall give its decision. The endered decision shall be binding to the Parties. If either Party is dissatisfied with the Adjudicator's decision and, within days specified in the SCC refer the dispute for arbitration. If either party within the period mentioned in the SCC has not referred the matter for arbitration the decision shall become final and binding to the Parties.



2.	Procedure For Disputes		arbit	ratio	tration shall be conducted in accordance with the on procedure published by the Institution named and accesshown in the SCC.
		32.2	The adjuand App	rate dica cos poin lity	of the Adjudicator's fee and administrative costs of ation shall be borne equally by the Parties. The rates sets shall be in accordance with the rules of the ting Authority. In conducting adjudication to its each party shall bear its incurred costs and expenses
		02.0	arb	itrat ce st	nown in the SCC.
33.	Replacemen of Adjudicator		Sh an in	ould d th	If the Adjudicator resign or die, or should the Employer are Supplier agree that the Adjudicator is not functioning cordance with the provisions of the contract, a new dicator will be appointed by the Appointing Authority.
34	Limitation Liability	of 34.	1 E	se O	t in cases of criminal negligence or willful conduct, and in the finfringement pursuant to GCC 8, The supplier shall not be liable to the Procuring Entity,
			a		whether in contract, tort, of old the consequential loss or damage, loss of use, loss of production, or loss of profits or interest costs, provided that this exclusion shall not apply to any obligation of the Supplier to pay liquidated damages to the Procuring Entity; and
				b)	The aggregate liability of the Supplier to the Procuring Entity, whether under the Contract, in tort or otherwise, shall not exceed the total Contract Price, provided that this limitation shall not apply to the cost of repairing or replacing defective equipment or to any obligation of the Supplier to indemnify the Procuring Entity with respect to patent infringement
	35. Notices		35.1	sha pro wr	y notice given by one party to the other pursuant to this Contract y notice given by one party in writing or in electronic forms that by it is sent to the other party in writing or in electronic forms that provide record of the content of the other party's address specified in SCC.
			35.2	A	notice shall be effective when delivered or on the notice's effective, whichever is later.



36. Taxes and Duties	36.1 A foreign Supplier shall be entirely responsible for all taxes, stamp duties, license fees, and other such levies imposed outside the United Republic of Tanzania. 36.2 If any tax exemptions, reductions, allowances or privileges may be a supplier in the United Republic of Tanzania the Supplier in the United Republic tax enable the Supplier to
	36.2 If any tax exemptions, reductions, allowances or privileges by available to the Supplier in the United Republic of Tanzania the Procuring Entity shall use its best efforts to enable the Supplier to benefit from any such tax savings to the maximum allowable extension between the Procuring Entity in the Procuring Entity. 36.3 A local Supplier shall be entirely responsible for all taxes, duties license fees, etc., incurred until delivery of the contracted Goods the Procuring Entity.



NEGOTIATION MEETING

SUPPLY OF ICU EQUIPMENT

Venue: Executive Director Board Room

Date: 3/12/2021

Quotation no: PA-010/2021-22/G/33

Attendance:

1.	Gabriel Sungi	Chairman	ORCI
2.	Chausiku Chapuchapu	Member	ORCI
3.	Jovitha Jovin	Secretary	ORCI
4.	Nafasa R. Marombwa	Member	Morogoro
5.	Dr. Lobikieki Kissambu	Member	Mtwara Zonal
6.	David Obed	Member	Anudha
7.	Nisha Patel	Member	Anudha

Agenda

- 1. Opening of the meeting
- 2. Adoption of the agenda
- 3. Discussion on areas for technical and financial negotiation
- 4. Closing of the meeting

1. Opening of the meeting

The meeting was opened by the chairman of negotiation team at 11:03AM, by welcoming all the members including the representatives from Anudha Limited to introduce themselves, which they all complied. The chairman went on explaining the purpose of the meeting and proceeded with agenda number 2.

2. Adoption of the Agenda

All Agenda were read out and confirmed.



3. Discussion on areas for technical and financial negotiation

The quoted price for supply of ICU equipment for ORCI, MTWARA ZONAL, LIGULA RRH and MOROGORO RRH were read out at the meeting to be TZS 6,876,916,400 while the budget was TZS 3,767,200,000.

The need for negotiation between these health facilities and Anudha was because some of the equipment were omitted according to the need/requirement from each facility, which needed to be resolved at the meeting and also to ask for the cost reduction.

The Anudha Limited agreed to drop the amount quoted for the accessories for each required machine. With regard to this discount the amount was still high as these consumable accessories has got very low price compared to the required machines and available funds. Hence, Anudha agreed to reduce the price for each specific item. General agreement was reached that, with the total budget of TZS 3,767,200,000, Anudha limited will supply the required items at the total cost of TZS 3,767,198,500.

4. Closing of the Meeting

The meeting was concluded at 4:00 PM.

Prepared by

Jovitha Jovin Secretary / ORCI

Approved by:

......

ANUDHA LIMITED P.O.Box 5982 DAR-ES-SALAMI TEL: 2122745, 2122,47 FAX: 2126490

Gabriel Sungi Chairman Ocean Road Cancer Institute

Nisha Patel CEO Anudha Limited

Ton

MEGOTIATION MEETING CCU & FMD EQUIPHENT

Attendance:

1. Grahmed Song:

2. Cha usiku Chapuchapu

3 Jovitha Jovin

4 NAFSA RIMARONBUM

5-DR LOBHERER KISSAMBU

5. Eng. David Obec!

1. Nicha Patel

TITLE Chrisman SIGH.

Chrisman Strip

Minber

Socretary

Member

Member

Amudha representative

Amudha seprentative

M.

3/12/2021

Mo

THE UNITED REPUBLIC OF TANZANIA

Ocean Road Cancer Institute

Telephone: +255 22 220002

Fax:

E-mail: info@orci.or.tz



Barrack Obama Road, Lithuli Road Dar es Salaam Ilala 3592, Dar es Salaam Tanzania, United Republic Of

Date 08/12/2021

In reply please quote
PA-010/2021-22/G/33
Name of awarded PE
OCEAN ROAD CANCER INSTITUTE

RE: Supply of ICU equipments and consumables for ORCI, Mtwara zonal, Ligula and Morogoro RHH SUB: NOTIFICATION OF CONTRACT AWARD

- 1. Reference is being made to the bid documents submitted by 26/11/2021, for the above captioned matter.
- 2. Kindly be informed that the Ocean Road Cancer Institute Tender Board during its ordinary Meeting held on 01/12/2021, approved award of the contract to ANUDHA LTD. For Supply of ICU equipments and consumables for ORCI, Mtwara zonal, Ligula and Morogoro RHH at the contract price of TZS 3767153500.00 VAT inclusive. We hope you will provide us with best services

EXECUTIVE DIRECTOR





anudha limited

Tender Securing Declaration

Date: 24/11/2021

PA-010/2021-2022/G/33

SUPPLY OF ICU EQUIPMENTS AND CONSUMABLES FOR ORCI, MTWARA ZONAL, LIGULA AND MOROGORO RHH - Tanzania

Executive Director, Ocean Road Cancer Institute Luthuli/Samora/Baraka Obama Road, P.o Box 3592 Dar - es- salaam

We, the undersigned, declare that:

We understand that, according to your conditions, tenders must be supported by a Tender Securing Declaration.

We accept that we will automatically be suspended from being eligible for tendering in any contract with the Procuring Entity for the period of time to be determined by the Authority, if we are in breach of our obligation(s) under the tender conditions, because we:

- (a) have withdrawn or modified our Tender during the period of tender validity specified in the Form of Tender;
- (b) Disagreement to arithmetical correction made to the tender price; or
- (c) having been notified of the acceptance of our Tender by the Procuring Entity during the period of tender validity, (i) failure to sign the contract if required by Procuring Entity to do so or (ii) fail or refuse to furnish the Performance Security or to comply with any other condition precedent to signing the contract specified in the tendering documents.

We understand this Tender Securing Declaration shall expire if we are not the successful Tenderer, upon the earlier of (i) our receipt of your notification to us of the name of the successful Tenderer; or (ii) twenty-eight days after the expiration of our Tender.

Signed: Julian In the capacity of Director

Name: Anurag Hassija

Name: Anurag Plassija

Duly authorized to sign the tender for and on behalf of: Anudha Limited 121188, 2125748

ANUDHA LIMITED P.O.Box 5982



STANDARD POWER OF ATTORNEY

BY

ANUDHA LIMITED

("Donor")

AND

ANURAG HASSIJA

("Donee")

IN RESPECT OF TENDER NO. PA-010/2021-2022/G/33 FOR SUPPLY OF ICU EQUIPMENTS AND CONSUMABLES FOR ORCI, MTWARA ZONAL, LIGULA AND MOROGORO RHH.

DRAWN BY:

Pride Attorneys

Mshihiri Street

Morogoro Road

P.O. Box 315

DAR ES SALAAM

Email: <u>info@prideattorneys.co.tz</u>
Website: www.prideattorneys.co.tz



STANDARD POWER OF ATTORNEY

TO ALL IT MAY CONCERN

THAT BY THIS POWER OF ATTORNEY given on the 24th day of November, 2021.

We the undersigned **ANUDHA LIMITED** of P.O. Box 5982 Morogoro Road, Dar es salaam by virtue of authority conferred to us by the Board Resolution No. 89 of 24th day of November, 2021, **DO HEREBY** ordain and nominate **ANURAG HASSIJA** of P.O. Box 2019 Dar es salaam, to be our true and lawful ATTORNEY and Agent, with full power and authority, for us and in our names, and for our accounts and benefits to do any, or all the following acts in in the execution of Tender No. **PA/140/2021-2022/G/33** that is to say;

To act for the company and do any other thing or things incidental for SUPPLY OF ICU EQUIPMENTS AND CONSUMABLES FOR ORCI, MTWARA ZONAL, LIGULA AND MOROGORO RHH.

AND provided always that this Power of Attorney shall not revoke or in any manner affect any future Power of Attorney given to any other person or persons for such other Power of Attorney or powers shall remain and be of the same force and effect as if this deed has not been executed.

AND We hereby undertake to ratify everything, which our attorney or any substitute or substitutes or agent or agents appointed by him under this power on his behalf hereinbefore contained shall do or purport to do in virtue of this Power of Attorney.

SEALED with the common seal of the said **ANUDHA LIMITED** and delivered in the presence of us this 24th day of November, 2021.

IN WITNESS WHEREOF we have signed this deed on the 24th day of November, 2021 at Dar es Salaam for and on behalf of ANUDHA LIMITED

BEFORE ME

COMMISIONER FOR OATHS

ADVOCATE NOTARY PUBLIC & COMMISSIONER FOR OATHS

ANUDHA LIMITED
P.O.Box 5982
DAR-ES-SALAAM
TEL: 2121188, 2125746
FAX: 2126490

ACKNOWLEDGMENT

I, ANURAG HASSIJA, doth hereby acknowledge and accept to be Attorney of the said ANUDHA LIMITED under the terms and conditions contained in the POWER OF ATTORNEY and I promise to perform and discharge my duties as the lawfully appointed Attorney. I hereby accept the Special Power of Attorney conferred to me and I will act in good faith in honoring the powers so conferred to me.

introduced/Identif	ied to me by	the latter bein	g l
this 24th day	onally in my presence of November	2021	DONEE
BEFORE ME:	1/14	.1 4	TOR MHORO
Name:	Victor N	There /s	I AND ATE I LI
Signature:		· · · · · · · · · · · · · · · · · · ·	NOTARY PUBLIC \\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\
Postal Address:	315 I	Sm (0)	COMMISSIONER TO FOR OATHS
Qualification:	COMMISSIONER	FOR OATHS	+315, DAR-85

SIGNED and DELIVERED at Dar es Salaam by

the said ANURAG HASSIJA who is

