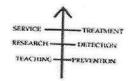


Box 3592, Dar es Salaam, Tanzania Tel. 2127597, Fax: 255-22-2118704



CONTRACTUAL AGREEMENT

Between

OCEAN ROAD CANCER INSTITUTE, TANZANIA

And

ANUDHA LIMITED, TANZANIA

TENDER NO. PA-010/2021-2022/G/38

FOR

SUPPLY OF AND INSTALLATION OF EMERGENCY MEDICAL EQUIPMENT FOR 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

Don

Form of Contract

THIS AGREEMENT made the F21khy 24 day of December 2021 between OCEAN ROAD CANCER INSTITUTE, junction of Luthuli road/ Barack Obama 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 AND INSTALLATION OF EMERGENCY MEDICAL EQUIPMENT FOR 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 Two Billion One Hundred Eighty Three Million and Eighty Two Thousand Only. TSH 2,183,082,000.00 (VAT inclusive) (hereinafter called "the Contract Price").

NOW THIS AGREEMENT WITNESSETH AS FOLLOWS:

- 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]



- 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 Two Billion One Hundred Eighty Three Million and Eighty Two Thousand Only. TSH 2,183,082,000.00 (VAT inclusive) in consideration of the provision of the goods and services and the remedying of defects therein, the Contract Price or such other 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

> Julius Mwarselage EXECUTIVE DIRECTOR Recuting Directo

IN PRESENCE OF:

NAME: ELIPENDOKAZIMOTO

DESIGNATION: HEAD OF LEGAL SERVICES

Signed by, for and on behalf of the Supplier: ANUDHA LIMITED P.O BOX 5962 DAR ES SALAAM
NAME: Anurag Hassija
DESIGNATION: Director
SIGNATURE: ANUDHA LIMITED P.O.Box 5982 DAR-ES-SALAAM TEL: 2122745, 212274 FAX. 2126490
IN PRESENCE OF: NAME: NIShita Patel
DESIGNATION: Managen
SIGNATURE: Notel

ace

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

THE SCHEDULE OF REQUIREMENTS

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TO



QUOTATION SUBMISSION FORM

DATE: 24th November, 2021

PURCHASER: OCEAN ROAD CARDIAC INSTITUTE TENDER NO. PA-010/2021-22/G/38

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 2,183,082,000/-, Two Billion, One Hundred Eighty Three Million and Eighty Two Thousand 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

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

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 Tender for and on beha

ANUDHA LTD. P. O. Box 5982

DAR-ES-SALAAM TEL: 2122745, 2122747

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ECG machine Model Manufacturer R12 Mindray - China	-	He Io weeks	-	6,460,000		4,000,000
Comprehensive maintenage for ECG Machine R12		to to weeks	=	2.500,000		6,400,000
Electrotonic Cardiac Maniter Model: ePM 15 with CO2_ECG Other	-	To to weeks	2	15,500,000		2,500,000
Comprehensive maintenace for Cardine Monitor			7 7	2,500,000		217,600,000
Fluid warmer Begler - Austra		I to J6 weeks	in	0.700.000		35,000,000
Mechanical Ventilator Medel, SV 300Pro Nitudity China	-	I to 16 weeks	CI	52,000,000		20,100,00
Comprehensive maintenance for Ventilator SV300 pro		to 16 weeks	es.	14,000,000		104,000,00
129 Opthalmascope Manufacturer, Koeler	-	I to 16 weeks	-	1,400,000		28,000,000



Duly authorized to sign the Tender for and on behalf of Dated on

Thy day of Stretchers and gurneys (wheeled 195 stretchers)
Model: YDC-3B Stethoscope - MDF 747Xp Vein finder Model: Vein Navi Manufacturer MDF - China Manufacturer Hebei - China 76 I to It weeks I to 16 weeks Direc Progres-SALAAM 6,500,000 TEL: 2122745, 2122747 FAX: 2126490 2,500,000 55,000 SBT1 + SBT2 + SBT3 TOTAL AMOUNT 2,183,082,000 15,000,000 6,500,000 110,000

Nate: In case of discrepancy between unit price and total, the unit price shall prevail

Do

TECHNICAL SPECIFICATIONS

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To



TECHNICAL SPECIFICATIONS SUPPLY OF EMD EQUIPMENTS AND CONSUMABLES FOR MTWARA ZONAL, LIGULA AND MOROGORO RHH

_ ITEM	DESCRIPTION	Yes/ No	If Yes, Provide Details or If No, Mention Why or Alternative if any
EMD MODEL: DA-2B	Fully Motorized ICU Bed) 5 Section PP Tops,	YES	5 section, with epoxy coated steel & X-ray board tops
'ith 10cm mattressH-13 +I.v	125mm Twin Wheel Castors. Central Locking Mech.,	YES	,
rod	Rubber Buffers at 4 corners Handset,	YES	
ad	ACP Box,	YES	Central control box under the bed only 1pcs battery
- 100 June	Battery Back Up,	NO	bed end can extension 20cm
7 - 7 - 7	SMPS Bed Extension,	YES	
-	Linen Holder,	YES	inside the side rail & head/foot
	Night Lamp,	YES	board
	Angle Indicator.	YES	
	Polymer Moulded Head and Foot Panels (Removable)	YES	
_	Polymer Moulded Split Safety Rails (2 sets)	YES	
	Urine Bag Holder (Single)	YES	H-13 10cm mattress, but with
	S.S Heavy Duty I.V. Pole (2 hooks)	YES	wave shape on one side
	Premium mattress, single section, 4", with wave shape on both sides, High quality, high density.	YES	
	Flame retardant foam and cover fabric having flame retardant, antibacterial, antiskid properties	YES	
	Mattress for Bed Extension	YES	
entilator	Should have facility for Invasive and Non-		COMPLIED
.1odel: SV 300Pro	Invasive ventilation		
	Five years comprehensive Maintenance inclusive		
	Microprocessor Control suitable for Neonatal.		
	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. Should have in-built facility to upgrade with EtcO2 Facility to Measure and Display: Status Indicator for ventilator mode Battery Indication Alarm 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 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





PATIENT	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 1 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		
MON ITOR	Modular & Suitable for Adult/Pediatric/ Patients monitoring	YES	
MODEL EPM 15 with CO2	Minimum 15 inches multicolour TET display screen	YES	with 15 inches capacitive touch screen
	Eight Channel digital and waveforms/traces display	YES	
	Capability of storage of patient data and printing of patient report	YES	
	Facility to monitor and display:-ECG, Respiration, NIBP, SpO2, EtCO2 and Temperature.	YES	with CO2 monitoring
	ECG	YES	with 12 leads ECG
	3 or 5 lead with cascade waveform facility	YES	
	Monitoring, Diagnostic & OT modes of monitoring ECG lead Simultaneous Multi - lead ECG monitoring of 7 ECG lead. HR range 20-350 bpm.	YES	Better, we have 3/5/12 leads monitoring
	HR/PR Source selection facility from Automatic, SpO2 IBP and NIBP	YES	





ECG	PULSE OXYMETRY: Display of Plethsmograph with Pulse strength indicator &SpO2 values and perfusion index. SpO2 range I - 100% and PR Range 20 - 230 BPM	YES	
MODEL R12	Electrocardiogram (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 lading
	ECG diagnostic monitor and recorder with printer 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
	ECG Features: 12 standard derivations aVR, aVL, aVF, I, II, III, and V1 - V6	YES	analysis),6 channels recording
	Simultaneous 12-leads acquisition adjustable ECG acquisition 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
	Adjustable ECG sweep (trace speed in mm/s) HR range 30 - 300 bpm with rhythm analysis Common Mode Rejection Ratio (CMRR) >89 dB.	YES	30 to 300 BPM CMRR > 110 dB
	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.	YES	Frequency 0.05 to 150 Hz Input Impedance > 50MOhm.
	Internal noise level <15 mV (p-p). Leakage current to patient <10 micro A With integrated keyboard, built-in printer with thermal	YES	
	printing head. Built-in rechargeable battery, autonomy >2.5 hours or 800 examinations.	YES	More than 3.5 hours of continuous
	Automatic switch to battery in case of power failure and automatic recharge on connection to mains.	YES	operation







francisco de la companya del companya del companya de la companya			
	ITEMS SUPPLIED WITH: 1 x patient cable 6 x reusable chest electrodes, suction ball type.	1	
	4 x reusable clamp electrodes 1 x supply of 960 thermal Z-folded sheets 300ml of ECG conductive gel		Thermal Z-fold A4 paper (210 mm x 295 mm)
	1 x spare rechargeable battery pack Power Input 240 VAC, 50Hz	: YES	
Portable ultrasound			
Model: Z-60	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 Ambulance and in the Emergency Room.
	Five years comprehensive Maintenance inclusive.	YES	5 years warranty
	Capable of generating imaging procedures involving lungs, heart, abdomen, pelvis, blood vessels, musculoskeletal and soft tissue. Five years comprehensive Maintenance inclusive.	YES	Capable of generating imaging procedures involving lungs, heart, abdomen, pelvis, blood vessels,
	optional touch screen combined with conventional user control panel. Mounted on trolley of Four casters two with brakes	NO	musculoskeletal and soft tissue. Laptop design, No touch screen
	normal use conditions Supplied complete with clear protective control panel cover for infection control. Imaging Focusing:	YES	endurance >120 min
	Adjustable focal depth, synchronization of focal zone to the selected scanning depth.	YES	Adjustable focal depth, synchronization of focal zone to
	Zooming Capability with automated image optimization. Image Orientation:	YES	the selected scanning depth.
1	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





1	Doppler, Colour Doppler Imaging (CDI), Power		2D/Colour Imaging mode with Cine loop.
	wave Doppler, Triple mode (optional) Software Application that include at least:	YES	Doppler, Colour Doppler Imaging (CDI), Power Doppler Imaging (PDI), Duplex, Continuous wave Doppler, Triple mode (standardl)
	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	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
	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	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
	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. Broadband Curvilinear transducer of at least 5-2	YES	annotations capture patient data, date and time, scanning 15" LCD screen, 38cm diagonal(across)
	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
lood gas analyzer	Portable		200~240V, 50/60MHz
Iodel: ST-200CC	Sample volume 110µl Result in one min after sample aspiration Power supply -220v with rechargeable battery at least 42mAh		YES 140μl YES 220V without rechargeable battery





	At least 10 parameter pH, pO2, pCO2, Na, K, Ca, Cl, Glu, Lac and Hct		YES
Suction machine Model: Askir C30	pump must be totally oil-free displacement	YES	
	international design for continuous use Motor shall be of Class "F" insulation to open to	YES	
	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. The suction capacity must be 25 liters per minute and can be regulated It must have two bottles of 21 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 provide foot switch. Base, top and panel made of rust proof and corrosion resistant moulded ABS.	YES YES	40Ltrs
	Accessories: Spare bottles: 02nos, Lid: 02nos, Rubber Seals: 02nos, Blades: 02nos, Suction Tubing set at least 5metres: 02nos,	YES	
	Supplied complete with all standard appearance	YES	
	1 ower Suppry: 220 - 240 V, 50Hz	YES	
yngoscope set	For Adults and Children (Marc and Miller Type) Large hollow, cylindrical, slightly ribbed handle. Handle made of either Chromium plated or Stainless Steel		COMPLIED





~	attorna nimited	
	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 Curved Nr 4, 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	
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	COMPLIED
BLOOD AND FLUID WARMER	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	COMPLIED





INFUSION PUMP/	One Charles		
VOLUMETRIC PUMP MODEL: VP3	One Channel at least Capable of accepting any kind o fluids (solutions and medications) Pump Capabilities: Flow Range: 01 to > 999 ml/hr Increments: 01 - 100 ml/hr	YES	Better Flow Rate 0.1-2000ml/hr
	Keep Vein Open (KVO) Rate: 1 - 5 ml/hr	YES	
	Volume to be Infused Selector (VTBI): 1 - 9999 ml Flow Pote Accessed Selector (VTBI): 1	YES	Better.KVO 0.1-5ml/hr
	Flow Rate Accuracy of +/- 5% When multiple channel automatic piggybacking	YES YES	
	Ingress protection not less than IPX2 Front Panel lockout Self-check carried out on	YES	IP23
	powering on	YES	
	Events Stored System: Log book Pause Infusion Facility required Anti-bolus System to reduce	YES	
_	pressure on sudden release of occlusion	YES	
1	IV Set Free-Flow Protection and Air Trapping Capability Needleless IV connection	YES	
	Dose Error" reduction system preferable Drug library software available, including updates (free during warranty)	YES	
	Air Bubble Detector with single and cumulative functions preferable Clearly visible optical alarms	YES	
	Acoustic Alarm not less than 45Db Real Time Display Availability of a Nurse call	YES	
	system connectable to a staff alerting system, 24V/02A static or dynamic preferable	YES	
I	Continuous operation within specification in ambient temperature of at least 5 - 40oC, Relative humidity of at least 10 - 90%	YES	
l a	Monitored and Displayed Parameters (colour and graphic preferable)	YES	
, F	Flow Pressure Dose Availability of software to nonitor the delivery of drugs preferable	YES	
	and delivery of drugs preserable	YES	







	Alarms Audible alarm required with volume control Momentary silence less than 2 min	7	
	Occlusion upstream and downstream Air in line and System malfunction Set loaded improperly	YES	not supply administration set. Open system please use commo
	and Door Open Infusion Complete and Loss of		universal Sets
	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	YES	Open system please use commo
	macro-bore or long extension sets: 100 Accessories Clamp for mounting pump on IV		universal Sets
	stand Clamp for external transportation		
	preferable Portability Data port required, at least RS 232 and/or USB interface Long analysis	YES	
	software and updates provided Wireless connectivity	123	
	Event log required and recording Software to	YES	
	diagnose and calibrate the equipment withaccess to calibration settings Power Supply Operates		
	from AC mains power: 220 - 240V 50Hz In-	YES	5 hours working
	built rechargeable battery Battery with operating time at least 4 hours at 25		
	IIII/III Automatic switch from AC mains power	YES	
	mode to battery operating mode and vice versa Total re-charging time not greater than 6 hours		
SYRINGE PUMP			
MODEL:	Microprocessor controlled programmed syringe pump	YES	
BENEFUSION SP3	Programming: Up to 99 steps of speed and time' Time Resolution: 0 to 999 minutes in 1 minute	YES	
	steps	YES	
	Time Resolution can be selected individually for each program step Accuracy: +- 1%	YES	
	Reproducibility: +/-0.2%Syringes: Glass plactic	YES	
	Metal syringes from 5 micro-liter to over 150ml Flow Rate Range: Depends on the inner syringe	YES	
	diameter 0.4nl/min with a 5ml syringe to 110ml/min with a 150ml syringe	YES	





1	Maximum Force: 300N reducible by a switch to 80N	YES	
	Motor: Microprocessor controlled brushless long life BLDC motor with Neodymium magnets	YES	
	Transmission: Efficient force transmission by a ball screw with highest mechanical load capacity of 12,000N	YES	
	Pusher Travel: 120mm	YES	
	Pusher Travel Rate: Minimum: 008 mm/min	YES	
	Maximum: 80 mm/min	YES	
	Speed Control Range: 0 to 999	YES	
	Non-volatile Memory: Storage of all settings Power Supply: 220 - 240V, 50Hz	YES	
	Interface: RS-485 or RS-232 automatic valve	YES	
	control	YES	
	Remote Control: 0 - 10V	YES	
	Operation Temperature: 0 - 400c	YES	
	Operation Humidity: 0 - 90% RH not	YES	
Portable Pulse	condensing	110	1
eximeter	Patient Range: Adult, Pediatrics and Neonatal	YES	4.3" TOUCH LCD
Anneter	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		Display range 0% ~ 100% SpO2 display resolution 1% Spo2 AccurayAdult/Pediatric : 70 100% ±2% Neonate : 70 ~ 100% ±3% 0 ~ 69% : unspecified
	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	YES	Measuring range 25 ~ 250bpm Resolution ±1bpm Accuracy ±2% or ±2bpm, whichever is greater Type Color TFT touch screen LCD Size 4.3"
	Alarm Audible alarm, audible button tone Supports Pitch Tone and multi-level volume	YES	SPO2 alarms and various system





CENTRAL		
MONITORING	General Description Modular and Suitable for	
STATION 16	Adult/Paediatric/ Patient monitoring	
CHANNEL	Minimum 15" multi colour TET display screen	
7	Eight Channel digital and waveforms/traces display	
	Capability of storage of patient 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	
	Temperature	
	ECG Multichannel (uo to 12 lead)	1
	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	1
	Display of Plethysmograph with Pulse Strength	
	a sour sine graph with I take Strength	







indicator and SpO2 values and perfusion index	
Sp02 Range: 1 - 100% PR Range: 20 to 230bpn	
ETCO2 Should be Main stream capnography	
with display of CO2 and digital Values of	
EtCO2, FiCO2 and RR	
EtCO2 Range: 0 - 99 mmHg	
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 Neonate	
Use selectable alarm settings: Mode: Manual,	
STAT (continuous 5 minute operation) and	
automatic (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	
Should be provided Respiration: RR range 1 -	
150 bpm	
Apnea alarms should be provided	
rends 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	





Recorder In-built dual channel thermal array	The same of the sa	
recorder		
Include Laser Printer and dual channel strip		
chart recorder		
Others Defibrillator and Cautery protection		
should be provided		1
Should work on Mains as well as Battery		
(backup for 2 hours) Automatic zoom in facility		
in the monitor display	1	
Should have facility to download trend data on		
USB and SD Card	1	
Supplied complete with all standard accessories		
and consumables that will enable the machine to		better, if display resolution
operate	YES	1920×1080, support display 36
Power Supply: 220 - 240V,50 Hz	IES	Patient Sectors with max.3 waves
CENTRAL STATION Central Station should	1	per patient
have facility to display up to 20 real time waves		
at a time and upgradable		
	YES	
Central Station should have		
Central Station should have separate patient		
window for viewing detailed real-time or stored data for individual patient	YES	1
It should have 24 by standard		1
It should have 24 hr stored patient data		
monitoring trends and 24 hr event review facility		better,
The Central Nurse Station should also have the		Full disclosure: Most recent 240
following: - Multi-lead arrhythmia and ST		hours of full-disclosure waveforms
review facility - 50 alarms strips storage per had	YES	and compressed waveforms
oner wave review with 24 hr full disclosure		Alarms: Most recent 3000 events
Optional facility for dual dieplay for detailed		including the parameter name and
analysis of individual bed without compromising		16-second waveform before and after an alarm is triggered
on full ICU monitoring		an anathr 12 miggered
		We offer ECG 24h Summary report
Facility for intenfactor xx .	YES	which is reference to doctor; we
Facility for interfacing Holter data for analysis, in case of the Holter from the same brand is		don't have holter,







	available. Export the ICU Patient data to Holter for analysis	YES	
	Remote display (Slave) facility should be available	YES	see the document accompanying the CMS
_	Real time recording thru Dual Channel Recorder	YES	3
	patient information and trend formats	YES	
	Advanced arrthythmia analysis package of at least 20 arrthythmia analysis 12 lead ECG Monitoring and view possible at Central Nurse Station	YES	Display: 1 page maximum: 3, tur the page to view all waves the monitor have
	Continuous full disclosure of up to 4 configurable waves per patient		Yes. Support Windows® 7 Professional SP1 Support Windows® 10
	Operate on Microsoft Windows NT workstation operating system	YES	■ Support Windows® Server 2008 ■ Support Windows® Server 2012 R2 ■ Support Windows® Server 2016 better, our screen is 24 inches
	Supplied with 19" Flat Screen TFT display, Laser Printer and Recorder, UPS	YES	
	The entire networking and cabling with hardware Wall mounts Power Supply: 220 - 240 V, 50Hz		
CENTRAL VEIN	Size 2 4 5 5 = 2		
CATHETER (CVC)	Size 3,4,5,6,7,8,9,10		Central Venous Catheter Triple Lumen 5FR 200mm Central Venous Access Catheter - Triple Lumen 7FR Central Venous Access Catheter - Triple Lumen 4.5FR 80mm





Oropharyngeal Air way (OPA)	Sizes 0,1, 2, 3, 4, 5 for Adult and Pediatrics	
Nasalpharyngeal Air Way (NAP)	Sizes 3, 4, 5, 6 for Adults and Pediatrics	
Endotrachea	Sizes 3, 3.5, 4, 5, 6, 6.5, 7, 7.5, 8	
DEFIBRILLATOR	Should be a Low Energy Riphagia J. Cl. 11	
	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 Should compensate for body impedance for a range of 25 to 150 ohms. Should have charging time of less than 5 seconds for maximum energy. Should have High resolution more than 8 inch colour display for viewing monitoring 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 Should have a battery capable of usage for at least 5 hours of monitoring Should be capable of printing Reports on event summary, configuration, self test, battery capacity etc Should have facility for self rest/check before usage and set up function. Should have facility to monitor parameters like SpO2, NIBP and etCO2 along with non-invasive facing (Demand and Fixed mode) facility should be able to upgrade the defibrillator for 12 read ECG monitoring and ECG transmission system Configuration Accessories, spares and	COMPLIED
C	onsumables:	





	Defibrillator with AED and external Pacemaker - Ipc Adult with built in Pediatric External Paddles - Ipc Patient Cables - Ipc ECG Rolls 50 Adult SpO2 reusable Sensor - Ipc Adult NIBP Cuff and Hose - Ipc EtCO2 Tubing (box of 20) - Ibox AED Multifunction Pads for Adults - 10 pairs with Each unit. The unit shall be capable of operating continuously in ambient temperature of 5 to 45oC and relative humidity of up to 95% Shall meet General Requirements of safety for Electromagnetic Compatibility. Power Input: 240 VAC, 50Hz	
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 - size 6 FR, OD 2.0 mm, 530 mm	
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	





Nasal Cannula	Nasal cannula (nasal prongs), device designed	
(prongs)	for easy administration of oxygen into the	
	patient nose through two small prongs placed in	
	the nostrils. It consists a factor in	1
	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	Illumination: Pressurized halogen lamp Image	
otoscope	1000 x 1074 pivels	
	Connection/Interface: USB 2.0 with 2m and 1	
	System Requirements: Windows VD Comit	
	Tack Electrical Rating. Lamp 3 5V DC 910	
	OSB. 3 V DC, 150mA Focal length: 114mm	
	inaginitation: 2.2X Constructed from ADC 1	
	recryfic Flastic of Stainless Steel Automatic	
	image Brightness and white balancing for	
	optimal image	
vall mounted	Range of lenges not and II	
Opthalmoscope	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, Fixation Star Colour	
	1 competature: Cool White in the range 2100	
	5500K Light Intensity: 8 000 - 12 000 lim	
	nee sedied optics and aspherical optical and	
	Red-fiet, Dive. (Teen and polarization Ch	1
	Training with On/Off switch Scratcheroof I	1
	Olass of Flastic Batteries. A A or recharge all	
	Dattery life at least 5 hours of use on full -	
	of fiesh batteries Accessories. Rattery Change	
	(do applicable to rechargeable type and a	
	Sufficient light bulbs for 3 years use. Rechargeable Battery 1.5V.	
	1.5 V.	







	anudha limited		
INSTRUMENT TROLLEY	Stainless steel shelves and railings Available in full stainless steel/powder coated 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 mm	YES YES YES	Stainless steel surface, without powder coated size: L730*W470*H970mm
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 Made of stainless steel with removable top tray	YES	Size: L600*W400*H850-1000mm
Neonatal ventlilator Model: SV 300Pro	I: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 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 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		COMPLIED





	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 trolley with wheels and brake facility Integrated medical air compressor Humidifier with autoclavable chamber and complete with patient circuit Circuit support arm Nebulization 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)	
Glucometer Glucoplus	·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	COMPLIED
Hemoglobinometer Hemocontrol	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	COMPLIED







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anaethetic Machine	□ Prequency (Hz): 50 / 60 □ Power (W): 300 □ Energy consumption (kWh/24h): 1.60 / 1.70 □ Heat emission (Kcal/h): 41 □ Compressor running time (%): 41 □ Noise level (dB(A)) (at 1m height & 1m	*Comply with Glostevace Helix Anesthesia-Diamedica from U.K -UAM(comply) .oxg concentrator . cylinder .pipeline -Vaporiser capacity(comply) -maximum liters capacity 10Lpm -Back up for ventilator 6hrs(more than 12hrs) -full fuction with UPS 20min(comply)
	□ Hold over time (+5°C to +10°C): 2 h 42 □ Climate class (ambient temperature range): SN / T (+10°C to +43°C) □ Defrosting technique: Natural □ Refrigerant type: R600a □ External dimensions H x W x D (mm): 1988 x 1139 x 1039 □ Inner dimensions H x W x D (mm) □ 1167 x 887 x 713 □ Supply voltage (V): 220-240 □ Frequency (Hz): 50 / 60 □ Power (W): 300	
refrigerator	□□Gross / Net volume (I): 895 / 763, Storage capacity (blood bags gross volume): 540 (450ml) / 750 (350ml) □Set temperature (preset): +4°C □Temperature cold / warm alarm limit: +2°C / +6°C	

P. O. BOX 5982, DAR ES SALAAM, TANZANIA

Plot 2169/82 & 8170/82, Morogoro Road, Opp. DART-KISUTU Bus stand

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Email: sales@anudha.com, sales@anudha.com, anudha@ctvsatcom.net, service@anudha.com





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	stabilisation. Full functionality with UPS for minimum of 20 minutes. Built in scavenger for exhaled gas Suitable for adults, paediatrics and neonates Peep 0-20cm H2O Patient monitoring:- 12.1" COLOUR TFT-LCD 60 minutes battery life ECG 5 leads, 3 as standard Sweep speed 12.5mm/s, 25mm/s, 50mm/s Heart rate range 10 -300 bpm Heart rate accuracy ± 1 % RESP 0 - 150rpm Temperature measurement range 25 - 50°C SPO2 Measurement range 0 - 100% Pulse measurement range 25 250bpm NIBP: adult/paediatric/neonate	uMEC 12 (Mindray) COMPLY COMPLY COMPLY COMPLY Sweep speed from 6.25-50mm/s comply comply Adult -120,ped/neonatal-150 comply
Operatting Table	Equipped with imported electric actuators for smooth & effortless working of the table. Must have hydraulic operation options also. It ensures the actuators working in a constant speed without any noise Designed to run even at low voltage to carry on the surgery. Top Dimension: L 1930 XW 553 m Height Adiustment: 762mm -1012 mm Trandelenburg / Reverse: 30°/25° Lateral Tilt: 20°/20° Kidney Elevator: 150mm Back Rest (up / Down): 80°/25° Leg Rest (up / Down): 15°/90° Head Rest (up / Down) °: 20°/60 Power Supply: 24v Dc Battery Backup: 2-3 Hours Key Structure The table has been made from acid proof, stainless steel (the face of the table is easy to clean and immune to disinfection agents) Strong	COMPLIED

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	& rust free base of stainless steel is provided to make the working conditions pleasant for the doctors & nurses. Orthopedic attachment (hips and legs support) including the patient's pelvis support and legs support enables the limbs strain performing. Design of the Orthopedic attachment & eccentric column provides monitoring of the limbs with	
	The various positions such as Up-Down movements, Trendelenburg Reverse Trendelenburg, Lateral Tilts, Flex-Reflex, Chair Position as realized by remote control. Leg section & head section are interchangeable. All accessories must be included	
Operating Lamp	LED Ceiling mounted surgical light \cdot Diameter of Lights: $\ \ \ \ \ \ \ \ \ \ \ \ \ $	COMPLIED

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

Equipment Descript	Technical specifications
EMD Beds	Fully Motorized ICU Bed) 5 Section PP Tops, 125mm Twin Wheel Castors. Centra Locking Mech., Rubber Buffers at 4 corners Handset, ACP Box, Battery Back Up, SMPS Bed Extension, Linen Holder, Night Lamp, Angle Indicator.Polymer
	Mattress for Bed Extension Should have facility for Inc.
	Should have facility for Invasive and Non-Invasive ventilation Five years comprehensive Maintenance inclusive.
	Microprocessor Control switch L. C. D. H.
	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 Should have in-built facility to upgrade with EtcO2
	Facility to Messure and Disabase and Disabas
/entilator	Facility to Measure and Display: Status Indicator for ventilator mode Battery Indication Alarm 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 200
	Aprilea 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
	Ose 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 Land Control of Ventilation:
	The state of the s
	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 1 year Expiratory Unit-life





Equipment Description	recrinical specifications
	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.
	1 5 WOL SUPPLY, ZZU - 74HV SHIP2
	Modular & Suitable for Adult/Pediatria/ Potient
	inches multi colour TET display screen.
	Eight Channel digital and waveforms/traces display
	Capability of storage of patient data and printing of patient
	Facility to monitor and display:-ECG, Respiration, NIBP, SpO2, EtCO2 and Temperature.
Monitor	Multichannel (w. 12) and Temperature.
	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 of 7 ECG lead. HR range 20-350 bpm.
	HR/PR Source selection facility from A
	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. SpO2 range 1 - 100% and PR Range 20, 220 PPL
	SpO2 range 1 - 100% and PR Range 20 - 230 BPM
	Electrocardiogram (ECG) digital monitor and recorder, 12-leads detection, multi-channel recording portable AC and but
	multi-channel recording, portable, AC and battery powered, with printer and accessories. Five years comprehensive M.
	accessories. Five years comprehensive Maintenance inclusive.
	ECG diagnostic monitor - 1
	ECG diagnostic monitor and recorder with printer ECG analysis and full interpretation (rhythm and events)
	interpretation (rhythm and events), real-time and manual.6-channels
	ECG Features: 12 standard derivations aVR, aVL, aVF, I, II, III, and V1 - V6
	Simultaneous 12-leads acquisition adjustable ECG acquisition and visualization modes ECG gain 2.5.5.10.20
	visualization modes ECG gain 2.5, 5, 10, 20 mm/Mv and Auto, accuracy +-
	Adjustable ECG sweep (trace speed in mm/s) HR range 30 - 300 bpm with
.cc	rhythm analysis Common Mode Rejection Ratio (CMRR) >89 dB.
CG	Frequency response (minimum quaranteed) 0.05 150 75
	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 > 50 MOhm
	2.5 MOhm or > 50MOhm.
	Internal noise level <15 mV (p-p)
	Leakage current to patient <10 micro A With interest 11
	. Finding head.
	Built-in rechargeable battery, autonomy >2.5 hours or 800 eventions.
	Automatic switch to battery in case of power failure and automatic recharge on connection to mains.
	mains.
	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
	1 x spare rechargeable battery pack Power Input: 240 VAC, 50Hz



Equipment Description	Technical specifications
	Complete with vascular, cardiac and curvelinear probes and Jelly for use in Ambulance and in the Emergency Room. Five years comprehensive Maintenance inclusive.
	Capable of generating imaging procedures involving lungs, heart, abdomen, pelvis, blood vessels, musculoskeletal and soft tissue. Five years comprehensive Maintenance inclusive
	with conventional user control panel. Mounted on trolley of Four casters two with brakes.
	Battery Duration: minimum 2 hours under normal use conditions Supplied complete with clear protective control panel cover for infection control. Imaging Focusing:
	Adjustable focal depth, synchronization of focal zone to the selected scannin depth.
Portable ultrasound	Zooming Capability with automated image optimization.
	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 (PDI), Duplex, Continuous wave 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 diameter and abdominal circumference, 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, scanning
	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.
	Broadband Curvilinear transducer of at least 5-2 MHz and 12-5 MHz transducer Power Supply: 220 - 240V, 50 Hz
	 Portable Sample volume 110μ1
Blood gas analyzer	Result in one min after sample aspiration
7-91	Fower supply -220v with rechargeable battery at least 42 Al-
	• At least 10 parameter pH, pO2, pCO2, Na, K, Ca, Cl, Glu, Lac and Hct
	The machine shall be portable on four wheels and with the life
uction machine	transportation. The suction pump must be totally oil-free diaphragm type.
uction machine	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.





Equipment Description	Technical specifications
	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. The suction bottles shall be autoclavable On/Off Switch and power indicator must be available. Shall provide foot switch.
	available. Shall provide foot switch.
	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 5metres: 02nos,
	Spare fuse: 01 set and Bacterial filter: 05 nos. Supplied complete with all standard accessories Power Supply: 220 - 240 V, 50Hz
	For Adults and Children (Marc and Miller Type) Large hollow, cylindrical, slightly ribbed handle. Handle made of either Chromium plated or Stainless Steel
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 Curved Nr 4, length approx. 15 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
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 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	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
nfusion Pump	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 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)





Equipment Descriptio	lechnical specifications
	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
	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
	Availability of software to monitor the delivery of drugs preferable) Flow Pressure Dose required with volume control Momentum.
	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
To the second se	Total re-charging time not greater than 6 hours
Syringe pump	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 with a 5ml syringe to 110ml/min with a 150ml syringe
syringe pump	Maximum Force: 300N roducible la
	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 Paris 100 - 90% RH not condensing
Portable 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





Equipment Description	Technical specifications
	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 multilevel volume General Description Modular and Spitch Left and Indiana.
	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.
	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 values and perfusion index SpO2 Range: 1 - 100% PR Range: 20 to 230bpm ETCO2 Should be Main stream capnography with display of CO2 and digital Values of EtCO2, FiCO2 and RR EtCO2 Range: 0 - 99 mmHg 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 Neonate
Central Monitoring Station 16 Channel	Use selectable alarm settings: Mode: Manual, STAT (continuous 5 minute operation) and automatic (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 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 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 2.1)
	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 standard accessories.
	CENTRAL STATION Central Station should have 6. 111
1	Central Station should have senarate patient wind.
	data for individual patient It should have 24 hr stored patient data monitoring trends and 24 hr event review facility
	stored patient data monitoring trends and 24 hr event review facility

Equipment Description	Technical specifications
	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 ICLI 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 the control of the
	arrany annual analysis package of at least 20 arrany three arrangement
	12 lead ECG Monitoring and view possible at Central Nurse Station
	Continuous full disclosure of up to 4 configurable waves per potient
	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
	Power Supply: 220 - 240 V, 50Hz
Central Vernous catheters (CVC)	Sizes 3, 4, 5, 6, 7, 8, 9, 10
Oropharyngeal Air way (OPA)	Sizes 0,1, 2, 3, 4, 5 for Adult and Pediatrics
Nasalpharyngeal Air Way (NAP)	Sizes 3, 4, 5, 6 for Adults and Pediatrics
Endotrachea	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 2007.
	comprehensive Maintenance inclusive
	Should monitor ECG through paddles pads and manitoring 1
	and paddies
	Should compensate for body impedance for a range of 25 to 150 ohms.
	Should have charging time of less than 5 seconds for maximum anarous
lefilbrillator	Should have High resolution more than 8 inch colour display for viewing monitoring 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
	Should have a battery capable of usage for at least 5 hours of
	capacity etc Should have facility for self test/check before usage and set up function
	invasive pacing (Demand and Fixed mode) facility
	Should be able to upgrade the defibrillator for 12 lead ECG monitoring and ECG transmission System
	Configuration Accessories, spares and consumables:



Equipment Description	lechnical specifications
	Defibrillator with AED and external Pacemaker - 1pc Adult with built in Pediatric External Paddles - 1pc Patient Cables - 1pc ECG Rolls 50 Adult SpO2 reusable Sensor - 1pc Adult NIBP Cuff and Hose - 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 5 to 45oC and relative humidity of up to 95%
	Shall meet General Requirements of safety for Electromagnetic Compatibility. Power Input: 240 VAC, 50Hz
Endotrachea Tube Introducers (Stylate and Bougie)	A disposable, latex free, sterile and individually wrapped device. Markings: a 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
	- size 6 FR, OD 2.0 mm, 530 mm
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
Nasal Cannula (prongs)	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 Cliff.
Wall mounted otoscope	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.
wall mounted Opthalmoscope	Range of lenses not smaller than -20D to +29D with steps not greater than1D. Anti-reflection lens Magnification: 12 - 15X Apertures: Small, Large and Semi Circle, 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 (as applicable to rechargeable type only) Sufficient light bulbs for 3 years use. Rechargeable Battery 1.5V. Stainless steel shelves and railings Available in full stainless steel/powder coated frame. Stainless steel bottom and to
nstrument trolley djustable Mayo table	Frame Mounted on 100 mm diameter wheels Overall Size: L 710 mm x W460 mm x H830 mm
wastable Mayo table	Made of stainless steel with removable top tray





Equipment Description	Technical specifications
	Height adjustable by foot 75 mm swivel castors Dimensions: Width 60 cm, Depth 40 cm and Height 120 - 130 cm
Digital Mobile X-ray	This Machine should be able to provide a minimum of 300 images per day. Should have a high frequency generator of 50-150KW, Automatic exposure device, Anatomical programming radiography, overloading protection feature, digital displat 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, warning sign and other important parameters. Components: Patient table: Mounted on heavy duty four castors with brakes to allow Longitudinal and lateral movement and better patient position. Floating table with up and down movement Table Size from 1800x800mm to 2200x800mm Patient weight: at least 150Kg Bucky wall stand: Height: 1900mm Center Height Stroke Range from 400mm to 1600mm Source to image detector should include the range of 90cm to 125cm Detectors: Fixed Flat Panel Detectors Active Image Size at least 17inch x 17inch 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.
	Protection against insect and rodent ingress to be incorporated





- Tompuon	Technical specifications
in p	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 x14, 14 x 17, 10 x 8). Protective gear (lead apron minimum of 2 small, 2 medium and 2 large) Googles (Minimum 1 small, 1 medium and 1 large). Gonad shield (minimum 2 smal 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 machine. It should come with the Power Backup System with capacity of power storage for 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 Shield Model Classic 0.35mmPb (iii) two (2) pairs of Patient Apron with belt 0.50mmPb Leadlite W/G-30 L-30 Printer Multi-tray printer for different X-Ray film sizes (10 x 12, 14 x14, 14 x 17, 10 x 8). Training, Installation and Utilization Requirements for commissioning: Manufacturer/supplier should perform installation, Requirements for commissioning: Manufacturer/supplier should perform installation, Requirements for commissioning: Manufacturer/supplier should perform installation, Training of user/s: Application specialist shall provide training of users in operation and basic maintenance: Five years comprehensive 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 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 Availability of technical personnel within the country should be stated; this should nelude CVs, work permits for foreign personnel. Software should be flexible and provide the room for upgrade to add ne
i i S S to E E E E E E E E E E E E E E E E E E	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 parameter to be measured by the Machine and report format. Documentation: Operating and service manuals (In English) including lists of inportant spares and accessories - with their part numbers and list of equipment and rocedures required for calibration and routine maintenance should be provided.





Equipment Description	Technical specifications
i i i i i i i i i i i i i i i i i i i	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 as 1-2ml (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. IE ratio 1:0 to 1:10 Inspired Time 0.1 to 2 see 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 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 on-screen message: MV high/low, Apnea, tube obstruction, FiO2, high/low high
Glucometer m	Hand held type Glucometer · Battery operated · Memory up to 10 measurements · Sticks at the sticks interring is available · Indication of high and low easurements · One box of sticks and punctures is included · Operating instructions is included

Equipment Description	Technical specific u
Hemoglobinometer	· Parameter: Hemoglobin, HCT(Hematocrit)· Principle: Optical reflectance · Sample: Capillary or venous whole blood (13–15ul) · Strip: H12 Hemoglobin Test Strip · Speed: ≤ 16 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 & Concentration, the CV of 10
plood bank refrigerator	 □Gross / Net volume (I): 895 / 763, Storage capacity (blood bags gross volume): 540 (450ml) / 750 (350ml) Set temperature (preset): +4°C •Temperature cold / warm alarm limit: +2°C / +6°C Hold over time (+5°C to +10°C): 2 h 42 Climate class (ambient temperature range): SN / T (+10°C to +43°C) Defrosting technique: Natural Refrigerant type: R600a External dimensions H x W x D (mm): 1988 x 1139 x 1039 Inner dimensions H x W x D (mm) 1167 x 887 x 713 Supply voltage (V): 220-240 Frequency (Hz): 50 / 60 Power (W): 300 Energy consumption (kWh/24h): 1.60 / 1.70 Heat emission (Kcal/h): 41 Compressor running time (%): 41 Noise level (dB(A)) (at 1m height & 1m distance): 47



Equipment Descript	Technical specificant
	Process with gravity air removal Process with
	•Microprocessor control system
	◆Steam generated by Heater in the chamber ◆Door lock is Mechanical cam system ◆Reservoir Capacity is 5L
	Reservoir Capacity is 5L
	DRY system: Fully automatic steam ejection
	3126: 380(W)×430(H)×640(L)
	*Chamber capacity: 310(W)×310(H)×450(L)mm
	30 L
	◆Chamber material : Acid resistant stainless ◆steel
	+steet
	Automatic microprocessor control system including the following the fo
	melauling the following Afactory process
	a) Steam sterilization 20 min at 132°C
	b) Drying time 30 mi
	c) Fast exhaust
utoclave machine	* Process 2 wrapped Instrument, Linen
	a) Steam sterilization 15 min at 132°C
	b) Drying time 20 mi c) Fast exhaust
	* Process 3 Glove Disert
	* Process 3 Glove, Plastic material
	a) Steam sterilization 20 min at 121°C b) Drying time 20 min
	c) Fast exhaust
	* Process 4 User select process
	a) Steam sterilization of the
	a) Steam sterilization 0 ~59 min at 110°C~135°C b) Drying time 0~59 min
	5. DRY system
	Fully automatic steam ejection
	The DRY system use the hot wind from air pump.
	6. The water Reservoir is in the machine.
	a stainless steel feed water tank's capacity is 5L
	7. Power: AC 220V, 50Hz, 2200W
	8. Tray: 300(W)×50(H)×410(L)mm 2EA
	7 1 2 3 5 (N) A 3 5 (N) A 4 TO (L) mm ZEA



Equipment Description	Technical specifications
Anaethetic Machine	A robust Universal Anaethesia Machine (UAM) general anesthesia workstation that delivers anesthesia effectively with integrated oxygen concentrator to generate its own oxygen and provides standard connectors for cylinder, pipeline, and portable oxygen. Complete anaesthesia machine with integral pneumatic ventilator and patient monitor. Vaporiser: Low resistance, Suitable for draw over and continuous flow, Single vaporiser, dual agent use for isoflurane/halothane, Capacity 150ml. Integral pneumatic Ventilator: Back up for assisted ventilation minimum 6 hours (more hours is prefered) Manual ventilation, Integral oxygen concentrator provides oxygen and medical air 8 – 10lpm at >95%. Alternative oxygen supply options: cylinder, pipeline oxygen. UPS for battery backup and voltage stabilisation. Full functionality with UPS for minimum of 20 minutes. Built in scavenger for exhaled gas Suitable for adults, paediatrics and neonates Peep 0-20cm H2O Patient monitoring: 12.1" COLOUR TFT-LCD 60 minutes battery life ECG 5 leads, 3 as standard Sweep speed 12.5mm/s, 25mm/s, 50mm/s Heart rate range 10 -300 bpm Heart rate range 10 -300 bpm Heart rate accuracy ± 1 % RESP 0 - 150rpm Temperature measurement range 25 - 50°C SPO2 Measurement range 0 - 100% Pulse measurement range 25 250bpm NIBP: adult/paediatric/neonate

Equipment Description	recnnical specifications
Operatting Table	Equipped with imported electric actuators for smooth & effortless working of the table. Must have hydraulic operation options also. It ensures the actuators working in a constant speed without any noise Designed to run even at low voltage to carry on the surgery. Top Dimension: L 1930 XW 553 m Height Adiustment: 762mm -1012 mm Trandelenburg / Reverse: 30°/25° Lateral Tilt: 20°/20° Kidney Elevator: 150mm Back Rest (up / Down): 80°/25° Leg Rest (up / Down): 15°/90° Head Rest (up / Down) °: 20°/60 Power Supply: 24v Dc Battery Backup: 2-3 Hours Key Structure The table has been made from acid proof, stainless steel (the face of the table is easy to clean and immune to disinfection agents) Strong & rust free base of stainless steel is provided to make the working conditions pleasant for the doctors & nurses. Orthopedic attachment (hips and legs support) including the patient's pelvis support and legs support enables the limbs strain performing. Design of the Orthopedic attachment & eccentric column provides monitoring of the limbs with the C-ARM during the operation. The various positions such as Up-Down movements, Trendelenburg Reverse Trendelenburg, Lateral Tilts, Flex- Reflex, Chair Position as realized by remote control. Leg section & head section are interchangeable. All accessories must be included
Operating Lamp	LED Ceiling mounted surgical light \cdot Diameter of Lights: $\ \ \ \ \ \ \ \ \ \ \ \ \ $



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	
	Definitions (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 EMD EQUIPMENTS AND CONSUMABLES FOR MTWARA ZONAL, LIGULA AND MOROGORO RHH	
	Governing Language (GCC 4)		
4.	4.1	The Governing Language shall be: ENGLISH	
	Applicable Law (GCC 5)		
5.	5.1	The Applicable Law shall be: Laws of Tanzania.	
	Country of Origin (GCC 6)		
5.	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.	
	Performance 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	

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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.
	Inspec	ctions and Tests (GCC 11)
9.	11.1	Inspection and tests prior to shipment of Goods and at final acceptance are as follows:
		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 (GCC 12)		
10.	12.2	The following SCC shall supplement GCC 12.2:
		The Goods shall be packed properly in accordance with standard export packing specified by the PE in the Technical Specification.
	Deliver	y 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



		(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
		(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 weel before arrival of the Goods at the port or place of arrival and, if no 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;
	1	(iii.) Manufacturer's or Supplier's warranty certificate;
		(iv.) inspection certificate issued by the nominated inspection agency, and the Supplier's factory inspection report; and
		(v.) certificate of country of origin issued by the Tanzania Chamber of Commerce, Industry and Agriculture or equivalent authority in the country of origin in duplicate.
		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.

13.	14.1	The Insurance shall be in an amount equal to 110 percent of the DD value of the Goods from "warehouse" to "warehouse" on "All Risks basis, including War Risks and Strikes.
	Incide	ntal 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 I	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.
	Warran	ty (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 (spears comprehensive Maintenance
	Payment	(GCC 19)
18.	19.1	The method and conditions of payment to be made to the Supplie under this Contract shall be as follows: Payment for Goods supplied from abroad:
		Payment of foreign currency portion shall be made in <i>Tanzania</i> shillings in the following manner:
		(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 of receipt of the Goods upon submission of claim supported by the acceptance certificate issued by the PE.
	F	Payment of local currency portion shall be made in: TZS within thirty 30) days of presentation of claim supported by a certificate from the PE declaring that the Goods have been delivered and that all other ontracted Services have been performed.
	P	ayment for Goods and Services supplied from within the United Republic of Tanzania:
	P R	ayment for Goods and Services supplied from within the United epublic of Tanzania shall be made in Tanzanian Shillings, as follows:



		(i) Advance Payment: 65 percent of the Contract Price shall be paid within thirty (30) days of signing of the Contract against a simple receipt and a bank Guarantee or insurance bond for the equivalent amount and in the form provided in the Tendering Documents or another form acceptable to the PE.
		(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 - N/A
	Prices	(GCC 20)
20.	20.1	Prices shall be adjusted in accordance with provisions in the Attachment to SCC.
		The price shall be fixed
	Liquid	ated Damages (GCC 26)
21.	25.1	Applicable rate: 0.1%
		Maximum deduction: is equal to the performance security.
		Note: 0.1 to 0.2 per cent per day of undelivered materials/good's value.
	Procedure 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

		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	The following words and expressions shall have the meanings hereby assigned to them:		
		b	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. 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) m	"GCC" means the General Conditions of Contract contained in this section.
	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



	For the purpose of this clause:
	"corrupt practice means the offering, giving receiving or solicitir of anything of value to influence the action of a public officer in the procurement process or contract execution; "coercive practice" means impairing or harming, threatening to impair or harm directly or indirectly, any party or the property of the party for the purpose of influencing improperly the action or that party in connection with public procurement or if furtherance of corrupt practice or fraudulent practice;
	"collusive practices" means impairing or harming, or threatening to impair or harm directly or indirectly, any part or the property of the Party for the purpose of influencing improperly the action or a part or in connection with public procurement or government contracting or in furtherance of a corrupt practice or a Fraudulent Practice
	"fraudulent practice" means a misrepresentation of facts in order to influence a procurement process or the execution of a contract to the detriment of the Procuring Entity and includes collusive practices among Suppliers, prior to or after submission designed to establish tender prices at artificial non-competitive levels and to deprive the Procuring Entity of the benefits of free and open competition;
	"obstructive practice" means acts intended to materially impede access to required information in exercising a duty under this Contract;
27.4	In the event the Procuring Entity terminates the Contract in whole or in part, pursuant to GCC Clause 26.1, the Procuring Entity may procure, upon such terms and in such manner as it deems appropriate, Goods or Services similar to those undelivered, and the Supplier shall be liable to the Procuring Entity for any excess costs for such similar Goods or Services. However, the Supplier shall continue performance of the Contract to the extent not terminated.



28.	Force Majeure	28.1	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 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
		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 continue to perform its obligations under the Contract as far as is reasonably practical, and shall seek all reasonable alternative means for performance not prevented by the Force Majeure event.

Do

29.	Termination for Insolvency	29.1	giv ba wi ter	ne Procuring Entity may at any time terminate the Contract by ving written notice to the Supplier if the Supplier becomes nkrupt or otherwise insolvent. In this event, termination will be thout compensation to the Supplier, provided that such mination will not prejudice or affect any right of action or remedy nich has accrued or will accrue thereafter to the Procuring Entity		
30.	Termination for Convenience					
		30.2	be a	Goods that are complete and ready for shipment within thirty days after the Supplier's receipt of notice of termination shall accepted by the Procuring Entity at the Contract terms and price. the remaining Goods, the Procuring Entity may elect:		
			a)	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.		
	Disputes Resolution	31.1	amid days disp nego to a	ne event of any dispute arising out of this contract, either y shall issue a notice of dispute to settle the dispute cably. The parties hereto shall, within twenty-eight (28) is from the notice date, use their best efforts to settle the ute amicably through mutual consultations and obtation. Any unsolved dispute may be referred by party in adjudicator nominated by the appointing Authority ified in SCC.		
			the rend	r the dispute has been referred to the adjudicator, within ays, or within such other period as may be proposed by Parties, the Adjudicator shall give its decision. The ered decision shall be binding to the Parties.		
		51.5	n eit may, arbiti SCC	her Party is dissatisfied with the Adjudicator's decision within days specified in the SCC refer the dispute for ration. If either party within the period mentioned in the has not referred the matter for arbitration the decision become final and binding to the Parties.		



32	. Procedure for Disputes	32.1	The arbitration shall be conducted in accordance with the arbitration procedure published by the Institution named and in the place shown in the SCC.	
		32.2	The rate of the Adjudicator's fee and administrative costs of adjudication shall be borne equally by the Parties. The rates and costs shall be in accordance with the rules of the Appointing Authority. In conducting adjudication to its finality each party shall bear its incurred costs and expenses	
		32.3		
33.	Replacement of Adjudicator	33.1	Should the Adjudicator resign or die, or should the Employer and the Supplier agree that the Adjudicator is not functioning in accordance with the provisions of the contract, a new Adjudicator will be appointed by the Appointing Authority.	
34.	Limitation of Liability	34.1	Except in cases of criminal negligence or willful conduct, and in the case of infringement pursuant to GCC 8,	
			a) The supplier shall not be liable to the Procuring Entity, whether in contract, tort, or otherwise, for any indirect or 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	
5.	Notices		Any notice given by one party to the other pursuant to this Contract shall be sent to the other party in writing or in electronic forms that provide record of the content of communication and confirmed in writing or in electronic forms that provide record of the content of communication to the other party's address specified in SCC.	
		35.2	A notice shall be effective when delivered or on the notice's effective date, 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 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 extent.
		36.3	A local Supplier shall be entirely responsible for all taxes, duties, license fees, etc., incurred until delivery of the contracted Goods to the Procuring Entity.



NEGOTIATION MEETING

SUPPLY OF EMD EQUIPMENT

Venue: Executive Director Board Room

Date: 3/12/2021

Quotation no: PA-010/2021-22/G/38

Attendance:

1.	Gabriel Sungi	Chairman	Ongr	
2.	Chausiku Chapuchapu		ORCI	
3.	Jovitha Jovin	Member	ORCI	
1	The state of the s	Secretary	ORCI	
4.	Nafasa R. Marombwa	Member	Morogoro	
5.	Dr. Lobikieki Kissambu	Member	Mtwara Zonal	
6.	David Obed	Member		
7.	Nisha Patel		Anudha	
		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 EMD equipment for MTWARA ZONAL, LIGULA RRH and MOROGORO RRH were read out at the meeting to be TZS 1,551,977,400 while the budget was TZS 2,180,102,000.

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 reduce 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. Hence, Anudha agreed to drop the price for each specific item.

General agreement was reached that, with the total budget of TZS 2,183,108,000 Anudha limited will supply the required items at the total cost of TZS 2,183,082,000 after reviewing the list

4. Closing of the Meeting

The meeting was concluded at 4:00 PM.

Prepared by

Jovitha Jovin Secretary / ORCI

Approved by:

Gabriel Sungi Chairman

Ocean Road Cancer Institute

ANUDHA LIMITEL

P.O.Box 5982

DAR-ES-SALAAM TEL: 2122745, FAX: 2126490

> Nisha Patel CEO

Anudha Limited

MEGOTIATION MEETING

four & FMD EQUIPMENT

Galinsel Sing, 2. Cha usiku Chapuchapu 3 Jovitha Jovin 4 NAFSA RIMARONIBION 5-DR LOBINGERCE KISSAMIBY 5. Eng. Dowid Obecil 1. Nisha Patel

717 LE Cherman Mamber Socretary

MEMBER Anudha representative

MEMBER

Anudha seprentative

116/2021

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/38
Name of awarded PE
OCEAN ROAD CANCER INSTITUTE

RE: Supply of EMD equipments for 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 EMD equipments for Mtwara Zonal, Ligula and Morogoro RHH at the contract price of TZS 2183082000.00 VAT inclusive.

 We hope you will provide us with best services

EXECUTIVE DIRECTOR



STANDARD POWER OF ATTORNEY

BY

ANUDHA LIMITED

("Donor")

AND

ANURAG HASSIJA

("Donee")

IN RESPECT OF TENDER NO. PA-010/2021-2022/G/38 FOR SUPPLY OF EMD EQUIPMENTS AND CONSUMABLES FOR MTWARA ZONAL, LIGULA AND MOROGORO RHH.

DRAWN BY:

Pride Attorneys

Mshihiri Street

Morogoro Road

P.O. Box 315

DAR ES SALAAM

Email: info@prideattorneys.co.tz
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/18 that is to say;

To act for the company and do any other thing or things incidental for SUPPLY OF EMD EQUIPMENTS AND CONSUMABLES FOR 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

ADVOCATE NOTARY PUBLIC

COMMISSIONER FOR OATHS

SEALED with the Common Seal of ANUDHA LIMITED

and DELIVERED at Dar es salaam in the presence of us this 24th day of November 2021

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

BEFORE ME

COMMISIONER FOR OATHS

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.

SIGNED and DEL the said ANURAG	IVERED at Dar es : HASSIJA who is	Salaam by]	1	i college
introduced/Identifie	d to me by	the latter be	eino L	Janes	
known to me person this 24 Th day of	nally in my presence Novembe	~ 2021		DONEE	_
BEFORE ME:	11:1- A	41	CTOR MH	*	
Name:	Victor 1	VIhoro /	ADVOCATE	011	
Signature:	- J	- 10	NOTARY PUBL	10 *	

Dsm

COMMISSIONER FOR OATHS

COMMISSIONER FOR OATHS



Postal Address: Qualification:



TENDER SECURING DECLARATION

Date: 24th November, 2021 TENDER NO: PA-010/2021-22/G/38

To, Executive Director Ocean Road Cancer Institute 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.

In the capacity of DIRECTOR
Name: ANURAG HASSIJA
Duly authorized to sign the tender for and on behalf of: ANUDHA LIMITED
Dated on 24th day of November, 2021

Corporate Seal

Corporate Seal

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

