

Technical Specifications of emergency equipment for pediatric and perinatal wards in CP Tiraspol, CP Bender and Rabnita, Slobozia, Grigoriopol hospitals

Item		Quantity
ITEM 1. PULSE OXIMETER		5
1.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipment
		ISO 9001: 2000 certified companies
1.2.	Basic Structure	Monitor for SpO2 and pulse rate
		Stationary and portable pulse oximeter
		Removable hand held display pulse oximeter
		Universal connection for sensors
1.3.	Technical Characteristics	a) Monitor:
		SpO2 and pulse rate ensures continuous monitoring during transportation
		Oximetry technology clinically proven to work under motion and poor perfusion conditions
		Patients perfusion level display
		Sensor signal quality indicator to ensure best sensor position placement
		Full patient alarms including high and low saturation and pulse rate
		Downloadable min. 24 hour memory at 2 seconds resolution with the ability to review trend data on screen
		Fast time to initial reading after sensor placement
		Probe off detection system to alert user of probe displacement
		High sensitivity mode for improved performance on poorly perfused patients
		LCD waveform display
		User selectable averaging time
		SpO2 range: 30-100%
		Pulse rate: approximately 0 ÷ 250 bpm
		Perfusion (optional): 0.02% ÷ 10%
		Saturation accuracy in motion for infant: ± 3 digits
		Minimum 6 hours battery life for the standalone device for bedside monitoring
		Configurable display, including big numbers option
		On screen alarm values
		Alimentation with power cord for 220 V
		b) Sensors:
		Availability of complete range of adhesive and reusable sensors for all patient weights
		Sensors with recessed photo detector to reduce ambient light and electromagnetic interference
Extended adhesive sensor life by use of rejuvenatable adhesive		

Item		Quantity
ITEM 1. PULSE OXIMETER		5
		Low weight adhesive sensors, <5gms Adhesive sensors manufactured from durable, moisture resistant material
1.4.	Accessories and Consumables	Interconnection cable 1 pcs 1 SpO2 sensor reusable for adults (finger) 1 SpO2 sensor reusable for children over 10 kg 100 single patient use SpO2 sensors for infant 25 single patient use SpO2 sensors for neonates lower than 2 Kg
1.5.	Warranty and Service Conditions	24 months from the moment of installation Maximum response time at beneficiary site: 72 hours
1.6.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language Training for at least 2 technical personnel on site at beneficiary location in Russian language Installation and commissioning User manual in English or translated into Russian language

Item		Quantity
ITEM 2. BASIC INCUBATOR		1
2.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark Compliant to IEC 60601 and amendments for medical electrical equipments ISO 9001: 2000 certified companies
2.2.	Basic Structure	Incubator hood Temperature controller Standard humidity module Mobile stand
2.3.	Characteristics	a) Incubator hood and mattress: Double wall and other parts of incubator easily removable for cleaning At least two opposite doors for quick access Small range temperature drop with door down Front door access One iris port on each lateral side, two irises on front and back or doors for access Hood can be raised separately from front door Mattress with minimum surface of 2,000 cm ² Tilting of the mattress Tubing access – minimum 4 b) Temperature controller: Microprocessor controlled Automatic test on start-up Heat up time to 39°C to be 30 minute or less Removable control unit for easy service Air flow pattern directed from front to rear of incubator across

Item		Quantity
ITEM 2. BASIC INCUBATOR		1
	entire width	
	Centralized display for patient, air temperatures control with alarms	
	display for function settings, parameters and trending, easy to operate and clean	
	Accurate air temperature control using two air probes	
	Air temperature range between: 20 and 39°C	
	Baby temperature control using reusable skin probe	
	Baby skin temperature range between: 35 and 37°C	
	Keypad lock function for protection	
	Override function at air temperature over 38°C	
	c) Standard humidity module:	
	Internal reservoir included	
	Front mounted for easy accessibility	
	All humidifier parts can be completely removed for filling and cleaning	
	Humidity range up to 70% RH	
	d) Mobile stand:	
	Stand on four anti-static wheels, from which two locking	
	Height variable 90-110cm or more	
	Two front drawers	
	e) Alarms:	
	audio alarms	
	Baby set temperature: + / - 1 ° C	
	High/Low air temperature: +3 / -1 ° C or better	
	Baby skin temperature probe fail	
	Air temperature probe fail	
	Power fail	
	Air flow fail	
	System fail	
	f) Accessories included:	
	Front drawers – 2 pieces	
	Mattress – 1piece	
	Rails for accessories – 2 pieces	
	Patient skin probe reusable –2 pcs	
	g) Physical properties:	
	Interior Noise level <60dB	
	Interior Aer flow <0,35m/s	
	Power requirements: 220/240 V, 50-60 Hz	
	Ambient operating temperature: minimum 20- 30° C	
	Ambient humidity: 0 – 90 %RH	
2.4.	Accessories	Front storage drawers
		Mattress
		Rails for accessories
		Patient skin probe
2.5.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours

Item		Quantity
ITEM 2. BASIC INCUBATOR		1
2.6.	<i>Incidental Services</i>	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation and commissioning
		User manual in English or translated into Russian language

Item		Quantity
ITEM 3. PHOTOTHERAPY LAMP		3
3.1.	<i>Standard requirements</i>	Compliant to EU Medical devices Directives CE Mark
		Compliant to IEC 60601 and amendments for Medical electrical equipment
		ISO 9001 certified companies
3.2.	<i>Basic Structure</i>	Phototherapy Unit
		Mobile Stand
3.3.	<i>Technical Characteristics</i>	a) Phototherapy Unit:
		Light characteristics:
		The possibility of adjusting the angle
		Irradiance >20 $\mu\text{M}/\text{cm}^2$
		Lamp characteristics:
		-LED Bulbs / tubes / estimated life time >30000 h
		-average life time fluorescent tube: minimum 1.500 hours
		Specify the field size [cm] -Diameter
		Low energy consumption
		Fault protection (specify)
		Maintenance free operation
		Easy light source replacement
		Eye protection 3 pics.
		b) Mobile Stand:
– The possibility of adjusting the angle		
– on 3-4 castors with locks		
3.4.	<i>Consumables</i>	Lamp 1 set
3.5.	<i>Warranty and Service Conditions</i>	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
3.6.	<i>Incidental Services</i>	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 4. SYRINGE INFUSION PUMP		14
4.1.	Standard requirements	<p>Compliant to EU Medical device Directives CE Mark</p> <p>Compliant to IEC 60601 and amendments for medical electrical equipment</p> <p>ISO 9001: 2000 certified companies</p>
4.2.	Basic Structure	Equipment for IV administration of drugs at controlled quantity and time
4.3.	Technical Characteristics	<p>Syringe type infusion pump for constant drug administration</p> <p>Flow rate: min range 0.1 ml – 999 ml/h (max 0.1 ml increments)</p> <p>Possibility of changing the flow rate whilst infusing</p> <p>Syringe loading sensor. The equipment will include KVO function</p> <p>High accuracy over the entire delivery range min +/-5%</p> <p>Accept all standard type syringe from 50/60 ml, 30/35 or 20 ml, 10 ml or 5ml</p> <p>Set-up fast and simple</p> <p>Bolus mode programmable</p> <p>Antibolus system</p> <p>Warns of pressure variation when there is a risk of occlusion or a possible leak in the infusion line</p> <p>Occlusion in preselected mode: three pre-selectable pressure alarm limits (occlusion)</p> <p>Infusion continuity protection</p> <p>Automatic internal battery operation during patient transfer AC power failure</p> <p>Infusion data memorization (flow rate, bolus rate, volume, volume limit, KVO rate)</p> <p>Event logging</p> <p>Electronic pressure management</p> <p>Infusion alarm:</p> <p>Pre-alarm end of infusion</p> <p>Pre-alarm volume limit</p> <p>Occlusion alarm</p> <p>Alarm for the end of infusion</p> <p>Alarm for volume limit</p> <p>Technical alarms:</p> <p>Disengaged driving mechanism alarm</p> <p>Low battery pre-alarm</p> <p>Discharged battery alarm</p> <p>Battery capacity display</p> <p>Unconfirmed programming</p> <p>Main malfunction alarms</p> <p>Syringe position control</p> <p>Syringe barrel clasp check</p> <p>Plunger head detection</p> <p>Occlusion pressure</p> <p>Locking syringe</p> <p>Fixing accessory</p>

Item		Quantity
ITEM 4. SYRINGE INFUSION PUMP		14
		Power 220 V AC / 50 Hz
		Battery life: min 3h/charge
4.4.	Consumables	200 connectors from the syringe to the patient for each syringe infusion pump
		200 syringe 20 ml for each syringe infusion pump
		200 syringe 50 ml for each syringe infusion pump
4.5.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
4.6.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Equipment assembly and installation on site
		User manual in English or translated into Russian language

Item		Quantity
ITEM 5. SUCTION UNIT		5
5.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipments
		ISO 9001: 2000 certified companies
5.2.	Technical Characteristics	Portable suction unit
		Suitable for adult and pediatric use
		Vacuum range: between around -20 and -300 mmHg, adjustable from a button
		Vacuum indicator
		Fast aspiration: at around 10-30 lpm flow
		Sealed water resistant unit
		Control panel tilted for easy viewing of the controls
		Main body made of durable and resistant material
		LED indication for bad battery
		Indication for battery charge status
		Remote internal AC charger for battery
		Charging circuit capable of running on low battery
		Power supply: 220VAC, 50Hz
		Rechargeable battery:
		Rechargeable battery
Long operating time at full vacuum (uninterrupted min 40-60 min)		
Low charge time: max 6-8 hours		
Fast charging: runs at least 1 hour after it is charged for 2 hours only		
5.3.	Accessories and consumables:	Reusable collecting container: min 1500 ml.
		Disposable hydrophobic bacteria filters – 3pcs.
		2 auto-clavable silicon suction tubes

Item		Quantity
ITEM 5. SUCTION UNIT		5
5.4.	Warranty and Service Conditions	Period: 24 months from the moment of installation of the equipment
		Battery warranty : at least 6 months
		Maximum response time : 72 hours
5.5.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation and commissioning
		User manual in English or translated into Russian language

Item		Quantity
ITEM 6. ELECTROCARDIOGRAPH (EKG)		4
6.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipment
		ISO 9001: 2000 certified companies
6.2.	Characteristics	12 channel ECG with 12 leads
		Minimum 6 Number of traces to be displayed simultaneously
		Screen LCD
		Preferable touch screen
		Resolution at least 640/480 pixels
		Possibility of introducing for each patient the following data: name, date of birth, date of recording, weight, age, diagnosis and comments
		The central Control panel will display all instructions in Russian or English language
		12 leads ECG: limb derivations: I, II, III, aVL, aVF, aVR; precordial derivations: V1, V2, V3, V4, V5, V6
		Manual selection for any 12-lead
		Graphic LCD viewing area for life 3-channel display with patient information
		Indicator for faulty leads
		Digital filters for all possible interferences. Band Filtres: muscle, low frequency, high frequency, 50 Hz
		Working frequency 0,67-150 Hz
		Calibration 1mV test
		Sensitivity and Frequency response test
		Sensitivity adjustment - 5, 10 and 20 mm/mV
		Gain adjustment - x ½, x 1 and x 2
		Full screen preview to determine quality of ECG
ECG interpretation with interpretation statements		
Low battery check and alarm		
Optional: indication of bradycardia and tachycardia		
Printer		

Item		Quantity
ITEM 6. ELECTROCARDIOGRAPH (EKG)		4
		Printing speed 25 mm/s or 50 mm/s
		Recording paper - thermal - A4 size
		Printout of patient ECG with patient information
		Storing at least 100 records (at least 10 seconds long strips)
		Defibrillation protection
		Pacemaker detection
		Built in accumulator. At least 2 hours autonomy
		The accumulator is re-charging automatically
6.3.	Consumables	Thermal paper rolls – 50 pcs
6.4.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
6.5.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 7. RESUSCITATION BALOON		6
7.1	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipment
		ISO 9001: 2000 certified companies
7.2.	Technical Characteristics	Resuscitation balloon for providing fast, simple and effective ventilation to non-breathing patients; self-inflating bags in three sizes: adult/pediatric, neonatal
		System assists the rescuer through :
		Pop-off pressure release valve which opens at pressures more than 20-40 cm H ₂ O
		Pressure manometer
		Valve system respond to the rescuer and the patient
		Visual alert of the pressure applied or for incorrect operation
		Audible overpressure alert
		Auto-clavable
		Oxygen reservoir system for each of the resuscitators
		Extension tube for patient connection
		Inspiratory resistance: 3-3.5 cm H ₂ O
		Expiratory resistance: 2-2.5 cm H ₂ O
		Dead space: max 8 ml
		Pressure relief valve
Low weight (for easy operation)		
Operating temperature range: min. -10°C to 50°C		
Storage temperature range: min. -40°C to +60°C		
7.3.	Accessories	Neonatal use masks – 6 pcs (2 of each size)
		Children use masks –6 pcs (2 of each size)

Item		Quantity
ITEM 7. RESUSCITATION BALOON		6
		Adult use masks – 6 pcs (2 of each size)
7.4.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
7.5.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 8. PORTABLE OXYGEN SYSTEM		5
8.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipment
		ISO 9001: 2000 certified companies
8.2.	Characteristics	Device for the oxygen administration of patients in emergency situations
		Robust and lightweight carrying case or carrying
		Oxygen cylinder, min. 5L approximate 150-200bar, filled (adapter to refill matching Moldavian connectors)
		Fixation for Oxygen cylinder
		Oxygen Pressure Reducer, infinitely adjustable, 3-15 L/min
		Pressure meter
8.3	Consumables	Flowmeter
		Tubing carries the oxygen from regulator to the delivery device
		Reusable ventilation masks for infants 3 sizes: 4 pieces each size
		Reusable ventilation masks for children 3 sizes: 4 pieces each size
		Reusable ventilation masks for adults 3 sizes: 4 pieces each size
		Set of reusable oropharyngeal tubes type Guedel (No.00, No.0, No.1, No.2, No.3, No.4, No.5): 10 pieces each size
8.4.	Warranty and Service Conditions	Set of reusable nasopharyngeal tubes type Wendel (Ch14, Ch16, Ch18, Ch19, Ch20, Ch22, Ch24): 10 pieces each size
		24 months from the moment of installation
8.5.	Incidental Services	Maximum response time at beneficiary site: 72 hours
		Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 9. GLUCOMETER		4
9.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipments
		ISO 9001: 2000 certified companies
9.2.	Technical Characteristics	Whole blood measurements
		Measurement range: min 10 – 700 mg/dl
		No light influence
		Compensated with temperature influence
		Serial imprecision: max 4 %
9.3.	Consumables	Strips for 500 of tests open market not limited to one company
9.4.	Warranty and Service Conditions	Period: 24 months from the moment of installation of the equipment
		Maximum response time : 72 hours
9.5.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

ITEM		Quantity
ITEM 10. SPHYNGOMANOMETER		6
10.1.	Standard requirements	Compliant to EU Medical devices Directives CE Mark
		Compliant to IEC 60601 and amendments for Medical electrical equipment
		ISO 9001 certified companies
10.2.	Characteristics	Digital Blood Pressure Monitor--Sphygmomanometer Upper Arm, for neonates, infant, children and adults
		Four mode for adult, pediatric ,infant, and neonatal selectable
		Measures systolic, diastolic pressure, and pulse
		Fully automatic one-touch operation
		Automatic inflation and deflation
		High resolution, large LCD display panel shows readings of blood pressure, pulse rate
		Optional 50 memories
		Cuff size
		Neonate cuff size
		Infant cuff size
		Pediatric cuff size
		Adult cuff size
		Battery and AC/DC Adapter
10.3	Accessories	All cuff size neonates, infant, children and adult - 1 pcs for each size
		Other accessories and consumables if needed
10.4.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 48 hours

ITEM		Quantity
ITEM 10. SPHYNGOMANOMETER		6
10.5.	<i>Incidental Services</i>	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 11. NEBULIZER FOR DRUGS		9
11.1.	<i>Standard requirements</i>	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipments
		ISO 9001: 2000 certified companies
11.2.	<i>Technical Characteristics</i>	Equipment for broncho-pulmonary pathologies: asthma, pneumopathies, cystic fibrosis
		Dedicated for all medicines: bronchodilators, antibiotics, corticoids, and mucolytic
		Running mode: the aerosol is produced continuous
		Auto-clavable up to 134°C
		Preferable reusable accessories (tubes, masks)
		Nebulizer:
		Container capacity: approximately 5 ml
		Equipped with tube with length approximately 1m
		Multi position mask for adult and pediatric
		Equipped with mouth end-piece and flask for dosage
		Compressor:
		Oil free
		Maintenance free
		Small dimensions (portable device)
		Low weight (portable device)
		Low noise device: approximate 30 dB
		Pressure adjusted to 0.5 bar
Possibility to be fed by an oxygen central source: compressor flow in charge with minimum 5l/min		
Medical device from electrical class I		
Level of protection: IP 30		
11.3.	<i>Accessories and Consumables</i>	Mouth end-piece – 5pcs
		Nose plug – 2pcs
		Nose piece – 2pcs
		Masks adult – 2pcs
		Masks pediatric – 10 pcs
11.4.	<i>Warranty and Service Conditions</i>	Period: 24 months from the moment of installation of the equipment
		Maximum response time : 72 hours
		Training for at least 2 medical personnel on site at beneficiary

Item		Quantity
ITEM 11. NEBULIZER FOR DRUGS		9
11.5.	Incidental Services	location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 12. TABLE FOR NEWBORN RESUSCITATION		1
12.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipment
		ISO 9001: 2000 certified companies
12.2.	Basic Structure	Height adjustable table
		Resuscitation Radiant Warmer consisting of:
		Heat source
		Bassinet and mattress
		Mounting column with fixed height
		Suction devices for cleaning of the neonatal airways
		System for resuscitation
		Storage compartments
12.3.	Accessories	Front drawers for storage
		X-Ray tray
		Rails for accessories
		Patient skin probe
		a) Heat source controller and functions:
		Mean irradiance at mattress level: minimum 10 mW/cm ²
		The central Control panel will display all instructions in English or Russian language
		Control panel located centrally at eye level
		Microprocessor controller with control modes:
		– Manual mode in minimum 5% increments from 0 to 100%
		– Automatic (pre-warm) mode
		– Servo (baby) mode
		Servo control temperature range between approximate: 35 – 37 °C
		Display temperature range between 30 and 42 °C
		Examination lamp of minimum 50 W and minimum 1,000 Lux
		Apgar timer integrated
		Patient skin probe, reusable – 2 pieces
		b) Bassinet and mattress:
		Mattress area minimum: 3,000 cm ²
		Hydraulic tilt mechanism for the bassinet at least: +/- 10° minimum
Mattress thickness:		
– minimum 2 cm		
– Ray cassette tray to fit under mattress-1 piece		
Four foldable bassinet walls		

Item		Quantity
ITEM 12. TABLE FOR NEWBORN RESUSCITATION		1
		Ventilator tube supports
		c) Mounting column:
		Fixed height column on mobile castors
		Built in rails for accessories – 2 pieces
		d) Resuscitation module
		Wall supply pressure 3-6 bar
		Cylindre for oxygen and cylindre for compressed air pressure
		Pressure (PIP) min 50 cm H2O for flow 15L/min
		PEEP 0-15 cmH2O
		Adjustable positive end expiratory
		Gas bleed 0-15 LP/M
		Precision blender (optional) 21-100% O2 +/-3% O2
		Mask resuscitation size 0,1 and 2 two pieces for each size. Circuits T pieces
		d) Storage compartments:
		Front storage drawers – 2 pieces
		e) Alarms:
		Audio alarms
		Controller heating alarms
		Check patient 15 minutes in Manual Mode
		Baby temperature +/- 1° C from Set Point
		High temperature Skin Temp. 39.0° C +/- 0.2° C
		Probe Short or open circuit / No probe
		System fail Indicates System Fail
		Power fail AC Power Interruption
		Alarm silence/reset intervals
		Check patient Resets clock for 15 minutes manual mode
		Baby temperature 10 minutes
		High temperature 2 minutes
		Procedural silence Presilences baby temp alarm for 5 minutes
		Alerts
		Manual mode System alerts every 30 seconds > 10 minutes, for 15 minutes
		Apgar timer Alerts at 1, 5 and 10 minutes
		f) Physical properties:
		Total height: maximum 200 cm.
		Mattress height at approximate 100 cm.
		Weight: maximum 100 Kg. (excluding accessories)
		Power requirements: 220/240 V, 50-60 Hz
12.4.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
12.5.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 13. COMPRESSOR FOR MEDICAL AIR		2
13.1.	Standard requirements	Compliant to EU Medical devices Directives CE Mark
		Compliant to IEC 60601 and amendments for Medical electrical equipment
		ISO 9001 certified companies
13.2.	Characteristics	Compatible with the resuscitation module from resuscitation table ventilator and CPAP
		Stand-alone medical air compressor
		Air Flow >40 L/min
		Noise <60 dBA
		adjustable pressure 2-4 bar
		Snap fit with the resuscitation module from resuscitation table to provide medical air
		Air quality should comply with European Pharmacopoeia compressed air purity class or ISO 8573
		Replacement of internal filters should be performed without removing the compressor
13.3.	Consumables	Provide at least 5 sets of air filters
13.4.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
13.5	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 14. FETAL HEART DETECTORS		4
14.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipment
		ISO 9001: 2000 certified companies
14.2.	Characteristics	Compact, lightweight and portable device complete with carrying case, rechargeable batteries, gel and optional head-set if any
		Specify measurement method
		Audible and visible heart beat indication
		Fetal Heart rate calculation and display
14.3.	Technical Requirements	Specify FHR range [bpm]
		Display size
		Indicate displayed parameters
		Speaker yes/no
		Power Requirements: VAC 220, 50Hz / Plug Type F

Item		Quantity
ITEM 14. FETAL HEART DETECTORS		4
		CONTROLS:
		ON/OFF
		Volume
		Audio mute
		Calibration
		Easy to clean
		Waterproof / fluid resistant
		Battery operation
		Rechargeable standard batteries
		With integrated battery charger or ext. charging station
		Consumables if any covering 1200 patients
14.4.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
14.5.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 15 CARDIOTOCOGRAPH		2
15.1.	Standard requirements	Compliant to EU Medical devices Directives CE Mark
		Compliant to IEC 60601 and amendments for Medical electrical equipment
		ISO 9001 certified companies
15.2.	Characteristics	Standard requirements
		Fetal monitor for prenatal, labor and delivery monitoring
		LCD or LED display
		Built-in thermal recorder and printer
		Dual Ultrasound Twins available for reliable non-stress ² testing and routine labor monitoring
		Color display and keyboard or similar for entry of patient name and ID with numerical presentation of FHR and UA
		Tachycardia and bradycardia alarm management
		Adjustable alarms
		Water-proof ultrasonic transducers
		Backup memory
		Fetal Heart Rate Monitoring
		Ultrasonic measurement for fetal hear rate (FHR) with pulse Doppler technology
		Audible alarm and visual messages based on physiological alarms
		Transmitter frequency minimum 1 MHz
		Maximum constant intensity: < 15mW/cm ²
		Heart rate fetal counting range between approximately 60-200

Item		Quantity
ITEM 15 CARDIOTOCOGRAPH		2
		BPM
		Automatic detection of fetal movement
15.3.	Accessories	Uterine Activity
		Uterine activity (UA) measurement with toco-transducer
		Measurement range between: 0-100 relative units
		Manual or auto zero adjust
		FHR transducers- 2 pieces, with connection devices
		TOCO transducer- 1 piece, with connection devices
		Fetal monitor paper- minimum 6,000 sheets
		Table top or roll stand support for monitor and transducers- 1 piece
15.4.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
15.5.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 16. NEONATAL MATTRESS HEATER		9
16.1.	Standard requirements	Compliant to EU Medical devices Directives CE Mark
		Compliant to IEC 60601 and amendments for Medical electrical equipment
		ISO 9001 certified companies
16.2.	Characteristics	Equipment used for compensation of heat losses in neonates
		Complete unit, microprocessor controlled functions for temperature, set values, etc.
		Temperature setting range
		Main mode from 35,0 to 37,0
		Additional mode from 37,1 to 39,0
		Accuracy of temperature control $\pm 1,0$
		Maximum deviation of mattress surface temperature after switching to maximum temperature mode $\pm 1,0$
		Switch off temperature of the heater when there is inadmissible overheating of the mattress surface,
		Heating time, not more than 20 min
		Alarms
		Alarm signaling OVERHEATING
		Main mode at temperature within, °C 38,0 \pm 0,4
		Additional mode at temperature within, °C 40,0 \pm 0,4
		Other malfunctions are indicated by alarms
Mattress		
Overall dimensions of the mattress, mm, not more than		

Item		Quantity
ITEM 16. NEONATAL MATTRESS HEATER		9
		– Height 20
		– Width 600
		– Length 750
		Mattress cover antimicrobial, fire retardant, fluid-proof.
		Control unit
		Display with LED
		Built in handle
		ON/OFF
		Display the temperature
		Alarm ON/OFF
16.3.	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
16.4.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language

Item		Quantity
ITEM 17. MOBILE EXAMINATION LIGHT		4
17.1.	Standard requirements	Compliant to EU Medical devices Directives CE Mark
		Compliant to IEC 60601 and amendments for Medical electrical equipment
		ISO 9001 certified companies
17.2.	Technical Characteristics	Optical lighting system reflector for optimum illumination of the operation light field, cool, white light, free from deep shadows. Adjustable height and articulated arm.
		Classical or LED lamp technology
		Light-field size >120mm
		Low power consumption: max 150VA
		On-off switch
		Lamp life please specify
		Light characteristics: Lux: approx. 15,000 at 1m, Color temperature: approx. 4,300°K
		Mobile stand with 5 castors at least 2 with breaks
		Lamp must be water proof and fluid resistance for cleaning purposes.
17.3	Accessories	Spare lamp – 1 pcs (if applicable)
17.4	Warranty and Service Conditions	24 months from the moment of installation
		Maximum response time at beneficiary site: 72 hours
17.5.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation

Item		Quantity
ITEM 17. MOBILE EXAMINATION LIGHT		4
	User manual in English or translated into Russian language	

Item		Quantity
ITEM 18. BILIRUBINOMETE CUTANEOUS		1
18.1.	Standard requirements	Compliant to EU Medical device Directives CE Mark
		Compliant to IEC 60601 and amendments for medical electrical equipment
		ISO 9001: 2000 certified companies
18.2.	Technical Characteristics	Non-invasive fast measurement
		Measurement position forehead or sternum
		Easy to read display - Specify type and range
		Printer (preferably integrated) and paper rolls
		Alarms
		Measurements error
		Low battery
		Others (specify)
		Technical Requirements
		Power Requirements: VAC 220, 50Hz / Plug Type F
		Battery (rechargeable) and mains operation
		Charging device (220V)
		Carrying case
Consumables for 200 patients		
18.3.	Warranty and Service Conditions	Period: 24 months from the moment of installation of the equipment
		Maximum response time : 72 hours
18.4.	Incidental Services	Training for at least 2 medical personnel on site at beneficiary location in Russian language
		Training for at least 2 technical personnel on site at beneficiary location in Russian language
		Installation
		User manual in English or translated into Russian language