BREAST PUMP

10.17		CONTRAIL
	Clinical purpose	A breast pump is a mechanical device that extracts milk from the breasts of a lactating individual. Breast pumps is an electrical devices powered by electricity or batteries.
2	Used by clinical department/ward	NICUandPICU
The second		ECHNICAL CHARACTERISTICS
1	Technical characteristics (specific to	
•	this type of device)	Cushioninsertedinsidethebreastcupsothatitdoesnothurtthem other.
		3) SuctionPressure100to250mmhb;useradjustable.
		4) Abletoexpressmilkfrombothbreastssimultaneously.
		5) Collectionbottlescanbeusedforstorageofmilkshouldbeautoclavablean dbiocompatible.
		6) Doublealternatingpumps/doublecyclingpumps.
		7) Shouldbemotorizedbreastpumpunits.
		8) Shouldbehospitalgradeandheavyduty.
		9) Breast pump should not come in contact with Milk
		10) shall be supplied with expression set (complete) reusable (washable and autoclavable)
		11) shall have dual mode (start with stimulation for 2 minutes followed by suction mode)
2.2	User'sinterface	Manual
2.3	Software and/or standard ofcommunication(where everrequired)	NA
		B.PHYSICALCHARACTERISTICS
3.	Dimensions(metric)	Portable
3.2	2 Weight(lbs,kg)	Compactunit(weightlessthan4kg)
3.	3 Configuration	LCD/LEDdisplaysuctiontiming
3.	4 Noise(indBA)	<60db
3.	5 Heat dissipation	NA
3.		Yes
		URCE(electricity,UPS,solar,gas,water,CO2)
4		220VAC+10%,50Hzpowersupply;5Aplug.
\vdash	.2 Batteryoperated	NA
4	.3 Tolerance (to variations, shutdowns)	±10%ofinputAC
4	.4 Protection	Electrical protection by resettable overcurrent breakers or replace able fuses.
4	Powerconsumption	Shouldbecompatible withother lifes a vingequipments running parallel.

पान के 9/3

	5.ACCESSO	RIES,SPAREPARTS,CONSUMABLES
	ccessories	1) Reusablecollectionbottlesalong-withbreastcups-50sets.
1	mandatory,standard, optional);Spareparts(mainones	2) Allkindsoftubes-50sets(Ifapplicable).
	optional);Spareparts(mamories ;Consumables /	3) Diaphragm-100Nos.
/	eagents(open,closedsystem)	4) Otheraccessoriesrequiredforoptimumfunctioningoftheequioment.
/ '	L.	5) All accessories shall be BPA free & CE marked (including
		disposable bottles, tubing, Breast shield- 24mm, Valve
		membrane, and extra as per need)
	BIDDIXG/BROGU	REMENTUERNS/DONATIONREQUIREMENTS
	STANCE OF THE PROPERTY OF THE	TALANDDEPARTMENTAL CONSIDERATIONS
	Atmosphere/Ambiance(airconditioning,humidity,dust)	1) Operating condition: Capable of operating continuously in ambienttemperature of 10to 40 deg Candrelative humidity of 15to 90% in ideal circumstances.
		2) Storagecondition:Capableofbeingstoredcontinuouslyinambie nttemperatureof0to50degCandrelativehumidityof15to90%.
		Disinfection: Partsofthe Device that are designed to come into contact with the patient or the operators hould either be capable of easy disinfection or be protected by resingle was discovered.
11,50	S	dbyasingleuse/disposablecover. STANDARDSANDSAFETY
	and the first of the state with the series had been also as the first of the series where the series are	
	Certificates(pre- market,sanitary,);Performanceands	
	afety standards (specific tothe	2) Manufacturer/suppliershouldhaveISO13485certificateforqualitys tandard.
	device type);Local	
	and/orinternational	3) ElectricalsafetyconformstostandardsforelectricalsafetyIEC-60601-1.
	8.TI	RAININGANDINSTALLATION
8.1	Pre-installation requirements:nature, values, quality,tolerance	Suppliertoperforminstallation, safety and operation checks before handover.
8.2	Requirementsforsign-off	Certificateofcalibrationandinspectionfromthefactory.
8.3	Training of staff (medical,paramedical,technicia	Trainingofusersinoperationandbasicmaintenanceshallbeprovided.
30	9.W.	ARRANTYANDMAINTENANCE
9.1	Warranty	5 years
9.2	Maintenancetasks	Maintainancemanualdetailingcompletemaintainingschedule.
9.3	Service contract clauses,includingprices	warrantyofthreeyearwithfreeservicing(min.3)duringwarranty.
		10.DOCUMENTATION
10.1	Operating manuals, servicemanuals, othermanuals	Userandmaintenancemanualstobesuppliedin Englishlanguage. Certific
	Servicemanuais, other manuais	ateofcalibrationandinspectiontobeprovided.
		Listtobeprovidedofequipmentandproceduresrequiredforlocalcalibration nandroutine maintenance.
		Listtobeprovidedofimportantsparesandaccessories, with their partnumbers and cost.
		Contactdetailsofmanufacturer, supplier and local service agent to be provided the contact of the
10.2	Recommendationsformaintenance	User/Technical/MaintenancemanualstobesuppliedinEnglish.

VIOT 3 2/3

1	11.NOTES		
A	Service Support Contactdetails (Hierchy Wise; including a toll free/landlinenumber)	Contactdetailsofmanufacturer, supplier and local service agent to be provided.	
1.2	Recommendationsorwarnings	Anywarningsignswouldbeadequatelydisplayed.	

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Specifications Colonoscope (Low End)

Technical Specifications:

Video Processor/Light source:

- Should be a Compact Processor, long life LED /Xenon light source having light weight.
- Should be able to provide 16:9 and 16:10 output for a HDTV monitor and should be compatible with Analog and digital (HD-SDI and DVI output) to reproduce high definition images/videos
- Should be equipped with special detection technology for detail observation by enhancing visibility of blood capillaries and mucosa
- Should contain portable memory(minimum 2 GB) & USB Slot for image recording and minimum 1 GB internal buffer memory of processor
- Should have automatic white balance
- Should be able to function Picture in Picture display & index function ability.
- Should be equipped with memory back up for settings & Lithium battery.
- Should have pre-freeze function to select the clearest still image automatically and noise reduction technology.
- Should have structure enhancement A for better observation of larger mucosal tissues in lower Gl and structure enhancement B for better observation of vascular tissues in upper
- Company Should have certificate of USFDA & CE notified body

High DefinitionLCD Monitor

- Full HD LCD monitor Medical Grade 24inch and above with high resolution 1920X1200 (WUXGA) & contrast ratio 2000:1.
- Lower Power consumption.
- Aspect ratio 16:9 & 16:10 with output of (1080/601:NTSC) (1080/501:PAL) with RGB
- DVI, SVideo, Input Compatible with video processor.
- Should have Picture-in-Picture and Picture-out-Picture for viewing side-by-side
- split screen images.
- Stand should supplied separately

Should have following specifications:

Four or more no. of remote control switches on control body. R

Compatible with leakage testing device with its air flow and pressure regulation through light sources air

Field of view 140 degree or more

Direction of view:

O degree, forward viewing

Depth of field:

3 to 100 mm or better

Distal end outer diameter: 12-13 mm or less

Insertion tube outer diameter: 12-13 mm or less Tip bending rage:

Up 180 deg, Dn 180 deg, Rt & Lt 160

Working length:

1600 mm or more

Channel inner diameter: 3.7mm or more

Minimum visible distance of instrument used through channel: 5mm or closer from the distal end Software -To store the data/Record patient information

Trolley-Local/indian make MS trolley with scope holder

Biopsy Forceps for Colonoscope -Reusable ellipsoid cup with needle biopsy forceps made from enhanced, easy to clean autoclavable materials to facilitate reprocessing.

Leakage testing Device to be provided with separate maintenance unit as well as the device should be compatible with video processor supplied

Laptop (4GB RAM+512GB SSD) Laser Printer to be provided

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Dr. Gurav P. D. Professor & Head of Dept.Surgery, Govt Medical College, Miraj & P.V.P.G.H., Sangli,

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DOUBLE SURFACE PHOTOTHERAPY

45	CIONIDIZATE DE LA CONTRACTOR DE LA CONTR		
1	Clinical purpose	1.USE	
4		Emits in the main radiation spectrum in the range between 400nm an 550nm for reducing the concentration of Bilirubin	
	Used by clinical department/ward	New born stablisation unit,SNCU	
`	Over view of functional		
	requirements	a) Provides filtered light using radiant electric lights,not fibreoptics.	
		b) Infantsupportedsecurelyinbassinettebelowbulbs.	
		c) Monitorshoursofradiantlightexposure.	
en a	2.T	IDECHNICAL CHARLESTON	
1	Technical characteristics	ECHNICALCHARACTERISTICS 1. Phototherapyshouldbebasedonl EDtechnology which afterfiltering	
	(specific to this type of device)	 PhototherapyshouldbebasedonLEDtechnology,whichafterfiltering shouldprovide,alightofwavelengthapproximately450to470nmwith peakwavelengthof450-460nmrange. 	
		2. Irradiancetobeminimum35µW/cm2/nmat40cmheightandUV shouldnotexceed10-4W/m2in180nmto400nm.	
		3. DigitalHourmetershowingtotalexposuretimeforcurrentpatienttobeclearlyvisiblebyoperator.	
		4. Effectivelightfield>700cm2.	
		5. Lamplifeshouldbeminimum20000hoursforLEDandshouldhav etimertoindicateitsusage.	
		6. Over temperaturesafetycutouttobeincluded.	
		7. Up,downandtiltingofheadshouldbepossible.	
		8. Theunitshouldbemountedwithcastorwheelswithbrakes.	
		9. Variationinintensityover5-6hours<10%.	
		10. The irradianceratio(min tomax)shall begreaterthan 40%onmattress.	
		 Greenindicatorlightshallbeprovidedtoindicatethatequipmentisrea dyfornormaluse. 	
		12. Interruptionandarestorationofthepowersupplydonotchangepreset alues.LEDheatcanbereducedbynaturalcooling.	
		13. LEDshouldbeprotectredfromfreefall.	
	역본 2	14. Itshouldnottoppleon10deginclinedangle.	
	,	15. Thetemperature of baby bedand metal surfaces should not exceed 40 deg Cand 43 deg C for other accesible surfaces.	
		16. Thereshouldbeintutivemethodtoindicatethelightsurfaceisattheappr opriatetreatmentdistance.	
		17. Mobilestandwithmovablecastorsandheightadjustmentfacilityalon gwitheasyswivellingofsourcebox. UnitcanbeusedalongwithInfantcar trolley,RadiantWarmerandIncubator.	
2	2.2 Settings	UP/DOWNadjustmentofOverHeadUnit;Thephototherapyunitshou ldbeabletoprovideeffectivetreatmentforbedsandincubatorsofvaryinghe ights (generally 1.0 to 1.6m). Adjustment of light intensity may be provided.	
12	2.3 User'sinterface	Manual	

Software and/or standard L ofcommunication(where everrequired)		EDDisplayandinbuiltsoftware	
0	thers		
	3.PH)	YSICALCHARACTERISTICS	
D: () ()		minimumspec:1650mmHeightX750mmWidthX500mmLength	
W		<20kg	
C	Configuration	Clearcabinetforobservationofinfant.	
		nfantbassinettetobeanintegralunitwhichshouldbedetachable.Unit	
	t	to provide shielding of infant in the event of bulb breakage.	
		Bulbmounttohaveangleadjustmentofatleast30degrees.Alls	
		urfacestobemadeofcorrosionresistantmaterials.	
1		Lightunittiltingfacilityandheightadjustmentfacility.	
4		<60dBA	
_	heat dissipation	The temperature of baby bedand metal surfaces should not exceed 40 deg Cand 43 deg C for other accesible surfaces.	
.6		Minimum3castorsandatleast2withbrakes	
	4.ENERGYSOURC	E(electricity,UPS,solar,gas,water,CO2)	
.1	PowerRequirements	220to240V,50Hz	
.2	Batteryoperated	NA	
1.3	Tolerance (to variations, shutdowns)	±10%ofinputAC	
4.4	Protection	Electrical protection by resettable overcurrent breakers or replaceable fuses, fitted in both live and neutral lines.	
4.5	Powerconsumption	Shouldnotbemorethan 160W	
4.6	Otherenergysupplies	Mainscabletobeatleast2.5mlength	
	5.ACCESS	ORIES,SPAREPARTS,CONSUMABLES	
5.1	Accessories (mandatory,standard,optiona	Completesetofreplacementtubestoallow3months' continuous operation Two replacements ets of fuses, if replace able type used.	
5.2	Spareparts(mainones)	Nosparesrequired	
5.3	Consumables / reagents(open,closedsystem)	Total 500nos. Infanteyemasks ofbothavailable sizes(termand pretermbabies).	
	6.ENVIRONMEN	TALANDDEPARTMENTALCONSIDERATONS	
6.1	Atmosphere/Ambiance(airconditioning,humidity,dust)	Capableofoperațingcontinuouslyinambienttemperatureof10to40degC ndrelativehumidityof15to90%inidealcircumstances.	
6.2	User's care, Cleaning,Disinfection&Sterilityissue	Completeunittobeeasilywashableandsterilizableusingbothalcoholandc lorineagents.	

	Certificates(pre- market,sanitary,);Performanceands afety standards (specific tothedevicetype);Localand/orinter national	ShouldbeFDA/CEapprovedproduct Shall meet IEC-60601-1-2:2007 Medical electrical equipment Part 1- 2:Generalrequirements forbasicsafetyandessentialperformance- Collateralstandard: Electromagnetic compatibility - Requirements and tests (OrEquivalentBIS) ShouldmeetIEC60601-1:2005standardrequirements ShallmeetIEC60601-2-50:2009MedicalElectricalEquipment-Part2- 50:ParticularRequirementforthebasicsafetyandessentialperformanceofin fantphototherapyequipment;
	87	ManufacturershouldbeISO13485certified RAININGANDINSTALLATION
8.1	Pre-mstallation requirements: nature, values, quality, tolerance	Suppliertoperforminstallation,safetyandoperationchecksbeforehandover.
8.2	Requirementsforsign-off	CertificateofCalibrationandinspectionfromthefactory.
8,3	Training of staff (medical,paramedical,technicia ns)	Trainingofusersinoperationandbasicmaintenanceshallbeprovided
8.4	Others	
	9. V	VARRANTYANDMAINTENANCE
9.1	Warranty	5 yearsforthemachineand20,000hoursforLEDs
9.2	Maintenancetasks	Maintenancemanualdetailingcompletemaintainingschedule
9.3	Service contract clauses,includingprices	Localclinicalstafftoaffirmcompletionofinstallation
9.4	Others	
		10.DOCUMENTATION
10.1	Operating manuals, servicemanuals, other manuals	Advancedmaintenancetasksrequiredshallbedocumented User,technicalandmaintenancemanualstobesuppliedinenglishlangua ge. Listtobeprovidedofequipmentandproceduresrequiredforlocalcalibrati onandroutinemaintenance
1.00	Otheraccompanyingdocuments	List to be provided of important spares and accessories, with their partnumbersandcost. Certificateofcalibration and inspection to be provided.
		11.NOTES
11.1	Service Support Contactdetails (Hierchy Wise;including a toll free/landlinenumber)	Contactdetailsofmanufacturer, supplier and local service agent to be provided
11.2	Recommendationsorwarnings	List to be provided of important spares and accessories, with their partnumbersandcost. Certificate of calibration and inspection to be provided.

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Specifications - Laparoscopy (Instrument, Trolly Kit & Accessories) (Code No 11472)

1. Camera System-01

Integrated video Laparoscope, 10 mm 30 degree direction of view.

- Two distal full HD sensors with depth of focus ?om no ?oo mm, optimized for stereoscopic endoscopy.
- Slender and light design weight should not be able to more than 840 gm for optimal ergonomics, integrated in a fine titanium case.
- Sterilisation options: Autoclavable, Stend R 900 S, NX, Steris R system V-pro and Et O
- Should have DVI-D output for transmitting the 3D sight ijn 9ο/ο p
 format at 4ο/ξο Hz to a 3D monitor
- Early switching between 2D and 3D modes.
- Integrated USB interface for saving captured videos or still images in 2D while working on 2 D mode
- Should have input keyboard for tile generator, 4-pn DIN socket.
- System should have facility of controlling additional equipment's though camera head buttons from sterile zone like light sources, insufflators and recording device etc.
- Camera processor should be upgradeable in 3D giving full HD motion form both sensors
- Focus range should be optimal
- should be autoclavable
- Easy switching from 3D to 2D from camera head can be done

should meet-CEA certification OR US FDA

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High Definition Medical Grade Monitor Capable of displaying 3D-01.

- LCD panel minimum २९ inches.
- High Definition 9820* 9200 pixel resolution.
- should supply clip on type glasses also for selectable wearing surgeons.
- Should be supplied with 20 pieces of light, passive circularly polarizes 3D . glasses
- Viewing angle-Horizontal: 900 degrees, vertical: 900 degrees
- Contrast 9800:9
- Should be supplied with 90 pieces of light, passive circularly polarized 3D glasses Clip on glasses:4

3. LED Light sources performance- Qty 1 this should be in high end

- Lamp type LED 904
- Color temperatus &oook
- Light Outlets- 9 Certified to CFA and USFDA.
- Light Intensity Adjustment: Contineuously adjustable either manually or automatically by camera head.
- Certified to CEA and USFDA

4. Fibre optic cable and Illuminator

- High light transmission for optimal imaging.
- Extreme heat resistance
- Should be supplied with diamter 80 mm, Longih 300 cm-Qy 2 nos.

५. Electronic CO२ Insufflator with Gas heating and smoke Education sytems, Qty-9

it should have the flowing features

heating facility must be inbuilt in thermoflator. Both Insufflator and smoke evacuation system should be separate units for convenience.

Heatable insufflation tube for sensitive the COR gas up to the

ate Profess patient body temperature rept of General Surgery

Medical College Mirro Vative Sensitive Mode for sensitive areas such as paediatric

application with safety limits in the pressure up to maximum 94 mm Hg and flow ranges up to 94 L/min.

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- High flow mode with flow performance upto 80-40 L/min.
- Individually programmable procedure list for starting device with personal pre ets.
- Easy and intuitive use with user friendly colour touch screen for easy and precise setting of set values for pressure and flow and of insufflations modes. As well as for clear display of corresponding set values and actual values.
- Optical and acoustic alarm signals in the event of patient over pressure.
- Fully automatic, electrically controlled gas refill.
- Safety systems: Constant monitoring of intra abdominal pressure; any overpressure is reduced immediately.
- Applicable for use in Laparoscopy and to assure an optimal air extrusion and minimizing risk of air micro emboli and for decreasing rate of contamination and minimizing risk of post operative wound infection.
 - control via Realistic User Interface

Technical Specification

- gas flow: 0-40 L/min
- Pressure :0-30 mm Hg(8000 Pa)
- High Flow mode (o to 40 L/min)
- Sensitive mode pressure 94 mm Hg and flow 94 L./min for sensitive application.
- Electronic control and colour touch screen
- Following data are displayed on touch screen

Associate Professions ufflations mode.

Dept of General Surger Walue pressure (0 to 30 mmHg)

Nedical Colleges of the profession of the surger of the surger

- Gas consumption (o-9991)

-Status of gas heating

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- Power supply: 900-280 VAC 40/ξο Hz
- Certified to: IEC ξοξο9-9, ce Lable acc to MDD, medical product class lib.

Type of protection against electrical shocks; Protection Class 9.

Degree of protection against shocks: Applied part type CF International protection code: IP2

Smoke evacuation System: Should supply smoke ecacuation system with endoflator.

Intelligent, fully automatic control of vacuum generated for smoke for smoke evacuation in combination with HF units integrated in SCB

Should include a footswitch that can be activated from sterile area to provide surgical smoke evacuation for standalone activation during Laparoscopy surgery

Electrosurgical Integration Compatibility: Atomic Activation via FSU footswitch of advance ESUs

Ouch key on the unit to enable smooth insertion of the tube Should be compatible with all standard insufflators with pressure control, ranging up to 50L flow capacity.

Should have parallel suction and smoke evacuation facilities.

Technical Specification

Line Voltage: 100.....240 V Power frequency: 52/60 Hz

Dimensions: (wxhxd): 30.5cm X 5 cm X 32 cm

- **6. IMAGE/VIDEO RECORDING AND DATA ARCHIVING SYSTEM-01** State of the art user friendly Medical grade system (certified to be used in OT) Should be offered with following features.
 - User should have full control of the system from the sterile field via camera head buttons, optional touch screen, optional foot switch
 - parallel (synchronic or independent) recording of two image sources.
 - Still images and videos in 2D FULL HD and 3D.
 - Intelligent, adaptive storage mamegment

Dept. of General Surgeriage location is freely definable and configurable. Storage in internal sovt. Medical College Singliv (? TB FIFO) USB storage media via ?.o.

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- Playback of 2D and 3D content on separate monitor (optional 3D system required)
- Controllable via 92" inbuilt touch screen
- Input voltage: 900-280VAC
- Input frequency: 40-ξο Hz
- Type approval: ICE ξοξο9-9-9, ENξοξο9-9, ENξο409-2

7. Endoscopic Trolley

- Equipment cart suitable for the equipment
- Monitor holding arm

8. Accessories

- Should be supplied with sterilization wire tray for all 3D Videos Scopes Qty - 9 no from same manufacturer.
- Should be EDA & CE approved product

All equipment's and accessories mention above should be from single manufacture better compatibility and service.

JUSTIFICATION

- > For diagnosis of various Gynaecological condition in diagnostic Laparoscopy.
- > For Gynaecological Endoscopies operative procedures like TLH, LAVH, Laparoscopies myomectomy, Lap tub plastic.

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P.M.P. Govt. Hospital, Sanc

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ultipara Monitor Specification (Lower End)

- Monitors ECG, NIBP, SpO2, RESP, TEMP, in standard configuration
- 2) Suitable for adult, paediatric and neonatal patient with specific parameter.
- 3) Arrhythmia analysis, S-T segment detection for 10 arrhythmias.
- 4) 10-12" High Resolution (touch screen), colour TFT display with 7 Waveforms.

Selectable waveform as well as background colour

- 5) Maximum 24 hours graphical and tabular trends of all S parameters
- 6) Built in rechargeable battery with 2 hrs. Of battery backup.

Supports Ethernet, wired LAN aud Central Monitoring system connectivity (if CMS and

- Multipara monitor should be of the same company would be preferred.)
 7) Built in protection against Electro Surgical Unit (Cautery)
- 8) Keys for Menu, alarm, Freeze, NIBP Start
- 9) Operation through single rotary switch

ECG

- 1) Lead 5-lead
- 2) Lead selection I, II, III, VR, aVI., aVF, V
- 3) Sweep speed 12/25/50mm/sec
- 4) Heart rate Range/ Accuracy Adult 15-300 bpm with accuracy of \pm 1 bpm or 1% whichever is greater
- 5) Alarm Range Adult 15-300 bmp
- 6) Paediatric/ Neonatal 15-350 bpm. Alarm events recallable
- 7) Arrhythmia and S-T analysis YES
- 8) Review Up to 300 sec real time waveform replay with se selectable sweep speed

NIBP

- 1) Method Oscillometric
- 2) Operation modes Manual/Automatic/STAT
- 3) Measurement unit mmHg/Kpa selectable.
- 4) Curf pressure display range: 0 to 300 mm Hg
- 5) Accuracy: +3 mmHg 6) Measurement types Systolic, diastolic, Mean.

SPO₂

- 1) Measurement Range 0-100%
- 2) Accuracy + 2 digits (80-100 %)
- +3 digits (70-80%)

Alarm Range (Visible audible) 0-100%

Pulse Rate Range 25-224 bpm, resolution lbpm

Accuracy + 2 bpm or 2% whichever is greater

Alarm range (Visible audible) 3-300 bpm

Wave from Plethysmograph

Saturation Technology Nelcore / Nonin / Equivalent.

Probe Neonate, paediatric, Adult or Universal (3 No.s)

410/31.9/2

surement range 25 degree-45 degree C solution 0.1 degree C (including probe)

Measurement unit Degree C/ Degree F selectable Channels 2 channels

IBP/ Dual IBP NA.

EtCo2

NA.

Warranty: - 2 Years.

Certification : - CE (NB)/USFDA Approval.

Various accessories and their number varies as per user requirements.

- 1. 5 lead reusable patient cable for ECG monitoring.
- 2. Adult Reusable SpO2 Sensor
- 3. Pediatric reusable SpO2 Sensor
- 4. Neonate reusable SPO2 Sensor
- 5. Adult & Pediatric Cuff for NIBP monitoring
- 6. Skin probe compatible for Temp. monitoring
- 7. Rectal probe compatible for Temp monitoring
- 8. Sure CO2 sampling line with Nafion Dryer for intubated patient 5 nos.
- 9. Adult Nasal sampling line 100 nos.
- 10. Water Trap-50 No.

370 SHING 6 & PUPGHSANAII

Multipara Monitor Specification (Lower End)

- (1) Monitors ECG, NIBP, SpO2, RESP, TEMP, in standard configuration
- 2) Suitable for adult, paediatric and neonatal patient with specific parameter.
- 3) Arrhythmia analysis, S-T segment detection for 10 arrhythmias.
- 4) 10-12" High Resolution (touch screen), colour TFT display with 7 Waveforms.

Selectable waveform as well as background colour

- 5) Maximum 24 hours graphical and tabular trends of all S parameters
- 6) Built in rechargeable battery with 2 hrs. Of battery backup.

Supports Ethernet, wired LAN aud Central Monitoring system connectivity (if CMS and Multipara monitor should be of the same company would be preferred.)

- 7) Built in protection against Electro Surgical Unit (Cautery)
- 8) Keys for Menu, alarm, Freeze, NIBP Start
- 9) Operation through single rotary switch

ECG

- 1) Lead 5-lead
- 2) Lead selection I, II, III, VR, aVI., aVF, V
- 3) Sweep speed 12/25/50mm/sec
- 4) Heart rate Range/ Accuracy Adult 15-300 bpm with accuracy of $\pm\,1$ bpm or 1% whichever is greater
- 5) Alarm Range Adult 15-300 bmp
- 6) Paediatric/ Neonatal 15-350 bpm. Alarm events recallable
- 7) Arrhythmia and S-T analysis YES
- 8) Review Up to 300 sec real time waveform replay with se selectable sweep speed

NIBP

- 1) Method Oscillometric
- 2) Operation modes Manual/Automatic/STAT
- 3) Measurement unit mmHg/Kpa selectable.
- 4) Curf pressure display range: 0 to 300 mm Hg
- 5) Accuracy: +3 mmHg 6) Measurement types Systolic, diastolic, Mean.

SPO₂

- 1) Measurement Range 0-100%
- 2) Accuracy + 2 digits (80-100 %)
- +3 digits (70-80%)

Alarm Range (Visible audible) 0-100%

Pulse Rate Range 25-224 bpm, resolution lbpm

Accuracy + 2 bpm or 2% whichever is greater

Alarm range (Visible audible) 3-300 bpm

Wave from Plethysmograph

Saturation Technology Nelcore / Nonin / Equivalent.

Probe Neonate, paediatric, Adult or Universal (3 No.s)

सहमागी प्राध्यापक, सिंहिंग्याख्यापी क्रियाखाँ स्वारोग व प्रस्तीवी क्रियाखाँ से स. सिरंज व पं. व. पा. क्रि. सांगली,

mot \$ 9/2

Temperature

Measurement range 25 degree-45 degree C
Resolution 0.1 degree C
Accuracy ± 0.2 degree C (including probe)
Measurement unit Degree C/ Degree F selectable
Channels 2 channels

IBP/ Dual IBP

NA.

EtCo2

NA.

Warranty: - 2 Years.

Certification: - CE (NB)/USFDA Approval.

Various accessories and their number varies as per user requirements.

- 9. 4 lead reusable patient cable for ECG monitoring.
- २. Adult Reusable SpO⊋ Sensor
- 3. Pediatric reusable SpOR Sensor
- 8. Neonate reusable SPOR Sensor
- 4. Adult & Pediatric Cuff for NIBP monitoring
- ξ. Skin probe compatible for Temp. monitoring
- 6. Rectal probe compatible for Temp monitoring

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- ८. Sure CO२ sampling line with Nafion Dryer for intubated patient 4 nos.
- Adult Nasal sampling line 900 nos.

90. Water Trap-40 No.

4107. B. 2/2

Specifications O.T. TABLE (Medium End)

Application

Surgical Table is a mobile, hydraulic surgical table designed to support virtually major surgical procedures including general surgical, orthopedic, neurology, ERCP, urology, gynecology etc with the addition of table accessories. Table features can be opting by its mechanism i.e. Lateral tilt, Trendelenburg/Reverse Trendelenburg, Chair Position, Flex – reflex and Adjustable height functions.

Should be constructed of stainless steel and other high quality materials. The table should be equipped with a large translucent top, for C-arm use.

Standard Accessory package -

- · Anesthesia Screen
- · Arm Board
- · Shoulder Support
- · Lateral Support
- · Hand Rest
- · Knee Crutches

Key Structure

- · Specialized for orthopedic, neurology, ERCP, laparoscopy and urology.
- · should provide excellent C-arm and surgical access.
- Slim line column offset relative to tabletop and base.
- · 150 kgs maximum patient weight.
- · Furnished with hard, robust stainless steel shell which provides the sense of dirt free &) sanitized operations. I
- -Smooth height adjustment of the table can be achieved by using the foot pedal.
- · Interchangeable general sections (head/leg).

Technical Constraints

Top dimension L 1980 x W 533 mm Height adjustment 750 mm - 1000 mm Trendelenburg / Reverse 30° / 25° Lateral tilt 20° / 20° Kidney elevator 150 mm Back Rest (up / down) 80° / 25° Leg Rest (up / down) 15° / 90° Head Rest (up / down) 20° / 60° Patient Weight Capacity 150 kgs (330 lbs)

STANDARD ACCESSORIES

Anesthesia Screen - The L-type anesthesia screen frame is the most commonly used and can be mounted on either side of the table. It can be adjusted in height and angle and conveniently fixed on the table Side-Rail Clamp to be ordered separately.

Shoulder Support - Made of stainless steel frame, foam pad and adjustable with require height and

स्वीरोग व प्रसुतीशास्त्र विभागः बा. बे. म. मिरज न प. वं. पा, या. रू, सांगली.

" स्त्रीरोग व प्रमुर्ताला ने विश्वामा ल्जाराम व अधुवास्तान्त्र । ता. क. संविकी।

प्राचिम वं प्रदुर्गातान्त्रं विश्विति. प्राचिम वं प्रदुर्गातान्त्रं विश्विति. प्रा. वै. म. मिर्श्व व प्रवा. कि. वि. वि

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PVPGHSangli

DEPARTMENT OF CONSERVATIVE DENTISTRY AND ENDODONTICS SPECIFICATIONS FOR SUCTION MACHINE (HIGH VOLUME)

- 1. Average value of various specification should be in the following range
- 2. Ex. Noiseless operation due to oil immersed vacuum pump
- 3. Mounted on castor for easy mobility
- 4. Hi-power suction due to ¾ hp piston type vacuum pump
- 5. Vacuum capacity:760 mm/hg
- 6. Power 230 V ac, 50 Hz
- 7. Automatic on and off with the help of suction tip holder
- 8. Operational: colour choice at extra cost
- 9. Size 38"x16x 11 (h x l x w)
- 10. Weight 20 kg
- 11. This machine should be used for 2 chairs one at a time
- 12. The machine can continuously use for 2-3 hours
- 13. It should have maintenance free mechanism for auto drain operation

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PUPGHSangly,

SYRINGE PUMP

			GENERAL	
	1.USE			
1.1 C	linica		designed toprecisely drive the plunger of a syringe down its barrel to infusea solution when it must be administered with a high degree of volume accuracy and rate consistency.	
1.2 U	Jsed b	oy clinical	NICU/PICU ·	
		tment/ward		
			TECHNICAL	
		2.	TECHNICALCHARACTERISTICS	
2.1	Clini	cal performances	Shouldacceptallinternationallyproduced/marketedsyringesandshouldbea bletodetectitautomatically,ShouldsupporttheBolussupplyofdrugonpresso fsinglebutton,asperneedandshouldbeabletopresetdifferentrangeofBolus supply.	
			PreferablytheunitshouldbeofBottom/sideloadedtoavoidaccidentalspillingo fdrugsanddamagetothemachine. (should not be top loaded)	
2.2		chnical characteristics ecifictothistypeofdevice)	1.Flowrateprogrammablerangeatleastfrom0.1to200ml/hr,instepsof 0.1ml/hr;andatleastfrom100to1200ml/hrinstepsof1ml/hr.	
			2. SaveslastinfusionrateevenwhentheACpowerisswitchedoff.	
			3. Bolusrateshouldbeprogrammabletoapprox500ml,withinfused volumedisplay.	
			4. Selectableocclusionpressuretriggerlevelsselectablefrom300,500and9 00mmHg.	
			5. Mustworkoncommonlyavailable 20-, 30- and 50-mlsyringes	
			6. Accuracyof±2%orbetter.	
			7. Maximumpressuregenerated ≤ 20psi.	
			8. Automaticdetectionofsyringesizeandproperfixing.	
		•	9. Anti-bolussystemtoreducepressureonsuddenreleaseofocclusion.	
			10. Pauseinfusionfacilityrequired.	
			11. Self-checkcarriedoutonpoweringon.	
			12. Comprehensivealarmpackagerequiredincluding:occlusionalarm,neare ndofinfusionpre-alarmandalarm,volumelimitpre-alarmandalarm,low battery pre-alarm and alarm, AC power failure, drive disengaged,syringeloadingerror,maintenancerequired.	
			13. ShouldincludeKVO(Keepveinopen)enablingfeature.	
			14. Itshouldbeanopensystemcompliant.	
			15. Adjusted stand for stacking up to 5 syringe pump	
	2.3	Settings	Singleloadablewithonesyringeofminimum20ml.	
+	2.4	User's interface	Automatic	
	2.5	Software and/orstandard of communication	Inbuilt	
			3.PHYSICALCHARACTERISTICS	
_	3.1	Dimensions(metric)	NA NA	
	3.2	Weight(lbs,kg)	NA	

4 1 1 1	7 figures 4	
1	Configuration	ramper-resistantcasemadeofimpactresistantmaterial.
\perp		Securelymountableontabletop, IV standorbed fitting.
1	Noise(indBA)	Noisefree
ŀ	heat dissipation	
	Mobility,portability	Yes
	DEVELOPE CENTER AND	
	Voltage(value, ACorDC, monophas	RCE(electricity, UPS, solar, gas, water, CO2)
	eortriphase)	220t0240 v,30HZ
	Batteryoperated	Internalrechargeablebatteryhavingat4to6hoursbackupfor10ml/hrflowratewith50mlsyringe.
3	Tolerance (to	10%
	variations, shutdowns)	
4	Protection	Batterypoweredalarmforpowerfailureordisconnection.
5_	Powerconsumption	25W
.6	Otherenergysupplies	Na
	5.ACCES	SORIES,SPAREPARTS,CONSUMABLES
5.1	Accessories	ClampformountingpumponIVstand.
	(mandatory,standard,optio	
<u> </u>	nal)	
5.2 5.3		Battery, syringe holder, PMO lines
	, — , — , — , — , — , — , — , — , — , —	Dattery, syringe notices, 1 two mies
د.د	reagents(open.closedsystem	
J.3	reagents(open,closedsystem)	
5.4	Others	
	Others	CUREMENTTERMS/DONATIONREQUIREMENT
	Others BIDDING/PRO	S
5.	Others BIDDING/PRO 6.ENVIRONM	S ENTALANDDEPARTMENTALCONSIDERATIONS
	Others BIDDING/PRO 6.ENVIRONM 1 Atmosphere/Ambiance(aircond	S
5.	Others BIDDING/PRO 6.ENVIRONM	S ENTALANDDEPARTMENTALCONSIDERATIONS Operating condition: -
5.	Others BIDDING/PRO 6.ENVIRONM 1 Atmosphere/Ambiance(aircond	S ENTALANDDEPARTMENTALCONSIDERATIONS
6.	Others BIDDING/PRO 6.ENVIRONM Atmosphere/Ambiance(aircond itioning,humidity,dust)	S ENTALANDDEPARTMENTALCONSIDERATIONS Operating condition: Capable of operating continuous ly in ambient temperature of 0t
6.	Others BIDDING/PRO 6.ENVIRONM Atmosphere/Ambiance(aircond itioning,humidity,dust)	S ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes.
6.	Others BIDDING/PRO 6.ENVIRONM Atmosphere/Ambiance(aircond itioning,humidity,dust) 1.2 User's care,	ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes.
6.	Dibling/Pro BIDDING/Pro 6.ENVIRONM Atmosphere/Ambiance(aircond itioning, humidity, dust) User's care, Cleaning, Disinfection & Sterility is ues	S ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7:STANDARDSANDSAFETY
6.	BIDDING/PRO 6.ENVIRONM 1 Atmosphere/Ambiance(aircond itioning, humidity, dust) 5.2 User's care, Cleaning, Disinfection & Sterilityise ues	S ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7.STANDARDSANDSAFETY CEorFDAcertified.
6.	Others BIDDING/PRO 6.ENVIRONM 1 Atmosphere/Ambiance(aircond itioning,humidity,dust) 5.2 User's care, Cleaning,Disinfection&Sterilityissues 7.1 Certificates(premarket,sanitary,);Performanceanceanceanceanceanceanceanceanceance	ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7.STANDARDSANDSAFETY CEorFDAcertified. Manufacturer/suppliershouldhaveISO13485certificateforqualitystanda
6.	Dothers BIDDING/PRO 6.ENVIRONM 1. Atmosphere/Ambiance(aircond itioning, humidity, dust) 6.2 User's care, Cleaning, Disinfection & Sterility is ues 7.1 Certificates(premarket, sanitary,); Performancean safety standards (specific	ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7.STANDARDSANDSAFETY CEorFDAcertified. Manufacturer/suppliershouldhaveISO13485certificateforqualitystanda ElectricalsafetyconformstostandardsforelectricalsafetyIEC-60601-
6.	Others BIDDING/PRO 6.ENVIRONM 1 Atmosphere/Ambiance(aircond itioning,humidity,dust) 5.2 User's care, Cleaning,Disinfection&Sterilityissues 7.1 Certificates(premarket,sanitary,);Performanceanceanceanceanceanceanceanceanceance	ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7.STANDARDSANDSAFETY CEorFDAcertified. Manufacturer/suppliershouldhaveISO13485certificateforqualitystanda
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6.	Dothers BIDDING/PRO 6.ENVIRONM 1. Atmosphere/Ambiance(aircond itioning, humidity, dust) 6.2 User's care, Cleaning, Disinfection & Sterility is ues 7.1 Certificates(premarket, sanitary,); Performancean safety standards (specific	ENTALANDDEPARTMENTALCONSIDERATONS Operating condition: Capable of operating continuously in ambient temperature of 0 to 50 deg Candrelative humidity of 15 to 90% in ideal circumstances. Capable of cleaning with alcoholor chlorine wipes. Capable of cleaning with alcoholor chlorine wipes. Capable of cleaning with alcoholor chlorine wipes. CE or FDA certified. Manufacturer/suppliers hould have ISO 13485 certificate for quality standare lectricals a fety conforms to standards for electricals a fety IEC - 60601-1, class II. Shall meet IEC 60601-1-2 EMC standard requirements. Certified to IEC - 60601-2-24: Particular requirements for the safety of infusion
6.	BIDDING/PRO 6.ENVIRONM 1 Atmosphere/Ambiance(aircond itioning, humidity, dust) 5.2 User's care, Cleaning, Disinfection & Sterilityist ues 7.1 Certificates(pre-market, sanitary,); Performanceand safety standards (specific tothedevicetype)	ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7:STANDARDSANDSAFETY CEorFDAcertified. Manufacturer/suppliershouldhaveISO13485certificateforqualitystanda ElectricalsafetyconformstostandardsforelectricalsafetyIEC-60601-1,classII.ShallmeetIEC60601-1-2EMCstandardrequirements. Certified to IEC-60601-2-24: Particularrequirements for the safetyof infusion pumpsandcontrollers.
6.	BIDDING/PRO 6.ENVIRONM 1 Atmosphere/Ambiance(aircond itioning,humidity,dust) 5.2 User's care, Cleaning,Disinfection&Sterilityiss ues 7.1 Certificates(premarket,sanitary,);Performancean safety standards (specific tothedevicetype)	ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7:STANDARDSANDSAFETY CEorFDAcertified. Manufacturer/suppliershouldhaveISO13485certificateforqualitystanda ElectricalsafetyconformstostandardsforelectricalsafetyIEC-60601-1,classII.ShallmeetIEC60601-1-2EMCstandardrequirements. Certified to IEC-60601-2-24: Particularrequirements for the safetyof infusion pumpsandcontrollers. ETRAININGANDINSTALLATION
6.	BIDDING/PRO 6.ENVIRONM 1. Atmosphere/Ambiance(aircond itioning, humidity, dust) 5.2 User's care, Cleaning, Disinfection & Sterilityist ues 7.1 Certificates(pre-market, sanitary,); Performanceand safety standards (specific tothedevicetype)	ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7:STANDARDSANDSAFETY CEorFDAcertified. Manufacturer/suppliershouldhaveISO13485certificateforqualitystanda ElectricalsafetyconformstostandardsforelectricalsafetyIEC-60601-1,classII.ShallmeetIEC60601-1-2EMCstandardrequirements. Certified to IEC-60601-2-24: Particularrequirements for the safetyof infusion pumpsandcontrollers. ETRAININGANDINSTALLATION
6.	BIDDING/PRO 6.ENVIRONM 1 Atmosphere/Ambiance(aircond itioning,humidity,dust) 5.2 User's care, Cleaning,Disinfection&Sterilityiss ues 7.1 Certificates(premarket,sanitary,);Performancean safety standards (specific tothedevicetype)	ENTALANDDEPARTMENTALCONSIDERATONS Operatingcondition: Capableofoperatingcontinuouslyinambienttemperatureof0t o50degCandrelativehumidityof15to90%inidealcircumstances. Capableofcleaningwithalcoholorchlorinewipes. 7:STANDARDSANDSAFETY CEorFDAcertified. Manufacturer/suppliershouldhaveISO13485certificateforqualitystanda ElectricalsafetyconformstostandardsforelectricalsafetyIEC-60601-1,classII.ShallmeetIEC60601-1-2EMCstandardrequirements. Certified to IEC-60601-2-24: Particularrequirements for the safetyof infusion pumpsandcontrollers.

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- T - 1	Training of staff (medical,paramedical,technic ians)	Trainingofusersinoperationandbasicmaintenanceshallbeprovided.
8.4	Others	
	9. V	VARRANTYANDMAINTENANCE
9.1	Warranty	5 year
9.2	Maintenancetasks	Advancedmaintenanceandcalibrationtasksrequiredshallbedocumented.
9.3	Service contract clauses,includingprices	Localclinicalstafftoaffirmcompletionofinstallation.
9.4	Others	
		10. DOCUMENTATION
10.1	Operating manuals, servicemanuals	User, technical and maintenance manual stobes upplied in english language.
		Listtobeprovidedofequipmentandproceduresrequiredforlocalcalibratio nandroutinemaintenance.
10.	2 Otheraccompanyingdocuments	List to be provided of important spares and accessories, with their partnumbersandcost. Certificate of calibration and inspection to be provided.
	The property of the property o	11.NOTES
11	.1 Otherinformation	Contactdetailsofmanufacturer, supplier and local service agent to be provided.
1	1.2 Recommendationsorwarnings	

E-UPROM code No- 11480. 31994105 31-90

PVPGHBangli

Specification

Vaccum Delivery System

- > Automatic Hands free Vacuum delivery system.
- > Ideal for Second stage prolonged labour with Oxipeto anterior position with outlet presentation.
- > Comes with complete autoclavable (formalin chamber or regular autoclave with 120de) silicone cups both for assisted vaginal delivery as well as for 'c' section.
- > Imported Dry motor with unbreakable Jar, SS surgical grade Trolley.
- > Heavy footswitch with Guarantee of 6000 pedal operation.

Justification:

> गंभीर प्रकारच्या प्रसुतिच्या वेळेला Vaccum Delivery System ही मशीन अति उपयोगी पडतेण्

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PVPGH5angli

Vessel Sealer with Electrosurgical Unit

(Code No.11626)

Monopolar, low-carbonization contact coagulation for microsurgery.

(soft coagulation current with low HF output voltage for minimal tissue impact)

- Adjustable power limit in watts.
- Nominal power up to 70 watts.
- Peak HF output voltage max. 240 Vp.
- Auto-stop function.
- Activation optionally via footswitch or handle pushbutton(s).
- Crest factor CF 1.5 CF

Bipolar coagulation with forceps and MIS instruments.

- Adjustable power limit in watts.
- Nominal power up to 120 watts.
- Minimum power selectable: 1 W.
- Auto-stop function selectable.
- Auto-start function selectable.
- Auto-start/auto-stop functions selectable.
- Coagulation degree adjustable.
- Min. number of available coagulation degrees 9 degrees
- · Impedance-modulated activation tone selectable.
- Activation/start delay.
- Activation optionally with double-pedal or single-pedal footswitch
- Peak HF output voltage max. 180 Vp
- Crest facto: CF 1.8 CF.

Bipolar, low-carbonization coagulation for microsurgery.

- Adjustable power limit in watts
- Nominal power up to 50 watts.
- Minimum power selectable: 0.1 W.
- Auto-stop function selectable.
- Auto-start function selectable
- Auto-start/auto-stop functions selectable.
- Coagulation degree adjustable.
- Min. number of available coagulation degrees 9
- Impedance-modulated activation tone selectable
- Activation/start delay.
- Peak HF output voltage max. 160 Vp
- Crest factor CF 1.7 CF.

Bipolar sealing mode for open surgery

- Process-controlled current with continuous impedance monitoring.
- Adjustable power limit in watts.
- Nominal power up to 320 watts.
- Reusable instruments with ratchet available.

Auto-stop function. ciate Professor Dept. of General Surgery Govt Medical College Miraj & P.v.P. Govt. Hospital, Sangli.

Associate Professor Dept. of General Surgery - Govt. Medical College Miraj & P.V.P. Govt. Hospital, Sangli.

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Dr. Gurav P. D. Professor & Head of Dept.Surgery. Govt Medical College, Miraj & P.V.P.G.H., Sangli.

- Sealing degree adjustable.
- Number of sealing degrees available 9 degrees
- Impedance-modulated activation tone selectable.
- Peak HF output voltage max. 180 Vp.
- Crest factor CF 1.9 CF.

Bipolar sealing mode for endoscopy

- Process-controlled current with continuous impedance monitoring
- Adjustable power limit in watts.
- Nominal power up to 100 watts
- Reusable instrument available; can be disassembled for cleaning
- Auto-stop function.
- Sealing degree adjustable.
- Number of sealing degrees available 9 degrees
- Impedance-modulated activation tone selectable.
- Peak HF output voltage max. 200 Vp.
- Crest factor CF 1.9 CF.

Neutral Electrode (NE)

- Automatic NE connection monitoring (optically and acoustically).
- Automatic monitoring of split NEs for proper contact with the patient's skin.
- For split neutral electrodes, continuous qualitative display of NE/tissue contact resistance during application (In addition to treatment parameters).
- Automatic recognition of single-surfaced and split neutral electrodes.
- Baby NE function with automatic power limitation or lock-out of certain currents.
- Baby NE available.
- Optional switchover for exclusive use of split NEs (prevents use of single-surfaced neutral electrodes).

Technical Requirement:

- Power requirements: 100-240 VAC
- Nominal frequency:50/60 Hz
- Maximum current input (line current): 6.3 A
- Maximum power input :600 VA
- Power input in standby mode: 40 VA
- Average heat dissipation: 40 W
- Protection class acc. to DIN EN 60-601-1:1
- Type (of applied part) :CF, defibrillation-proof
- Classification acc. to Directive 93/42/EEC Anh. IX I1 b
- Leakage currents (LF/HF): acc. To IEC 60601-1; IC 60601-2-2
- Duty type: int 10s/30s (=duty factor of 25%).
- Device fuses: T6.3 A (slow-blow)
- Sound level: HF indication: 55 dB (A), Alarm: 65 dB (A).
- Electromagnetic interference (EMI):acc. to EN 55011, DIN EN 60601-2-2.

Weight: Not more than 9 Kg.

Transport: maximum ambient temperature: from -25°C to +70°C, (-13°F to 158°F)

Transport: maximum humidity:10-100%

 Transportomaximum atmospheric pressure 700-00 Associate Professional Control of Con General Surgery ovi. Medical College Miraj & Dept. of General Surgery . Govt. Hospital, Sangli.

Boyt, Medical College Mirai & - 7 Govt Hospital, Sangli,

Dr. Gurav P. D. Professor & Head of Dept.Surgery, Govt Medical College, ""irai & P.V.P.G.H., Sangli,

- Operation: maximum ambient temperature: from +10°C to +40°C (50°Fto 104°F) Operation: maximum humidity:30-75%
- Operation: maximum atmospheric pressure: 700-1060 hPa
- Notified body: European CE or USFDA Certified.

Accessories

All accessories should be reusable, Autoclavable, made from the same manufacturer and should be

- 1. monopolar hand switch cable -
- 2. bipolar forceps cable -
- 3. Special bipolar non-stick forceps with 2 side irrigation facility. -
- 4. The unit should have special bipolar mode -fluide assisted coagulation for using the above
- 5. Special vessel sealing reusable instrument for open 200mm and laparoscopy 370mm, with integrated cutting facility. Cutting blade should be easily replaceable. Instrument should be easily removable for cleaning and sterilization. It should be Autoclavable. Plug and play with automatic recognition. Hand activation. 5mm and 10mm
- 6. Above same instrument 10mm, with double action jaws.
- 7. Vessel sealing claps for open surgery 18cm, 21cm, 23cm, 28cm.plug and play with automatic recognition.
- 8. Special vessel sealing reusable biclamp instrument with integrated cutting facility for pen surgery. Blades should be easily replaceable, it should have hand activation facility .it should plug and play with automatic instrument recognition.
- 9. Bipolar output socket should have facility for both single connector and 2 pin connector type
- 10. Vessel sealing option and all other bipolar modes should be accessible through both the 2 bipolar socket on the unit.
- 11. All monopolar current modes should be accessible through both the 2 monopolar socket on
- 12. There should option for swapping 2 programs through the normal hand pencil cable with 2 switches. No special cables are accepted for this function.
- 13. There should be option to integrate SMOKE EVACUATION SYSTEM with the unit
- 14. Disposable patients pads dual type with reusable cable
- 15. For bipolar saline tur unit should be compatible all reputed brand instruments

Demonstration:

Bidder to demonstrate the Equipment for evaluation of the techno commercial bid at his own cost and shall not charge on demand any extra cost from the Institution

Associate Professor Dept. of General Surgery ovt. Medical College Miraj & V.P. Govt. Hospital, Sangli.

Associate Professor Dept. of General Surgery **Sovt. Medical College Miraj &** P.V.P. Govt. Hospital, Sangli Professor & Head of Dept.Surgery, Govt Medical College, Miraj & P.V.P.G.H., Sangli.

419 3-3/3

319211105 of 92 PVPGH3amli Otoscope Mish End 10423

German Mini type with handle and disposable tips

- 1. Should be a convenient pocket type otoscope.
- 2. Should be provided with a halogen light source.
- 3. Should be able to detach the otoscope head.
- 4. Should provide no reflections and obstructions.
- 5. Should provide detachable accessories of various sizes.
- 6. Should have in built rechargeable battery. Recharge should be possible with direct mains supply

Ps = 14500