

नेपाल सरकार

स्वास्थ्य तथा जनसख्या मन्त्रालय लुम्बिनी अञ्चल अस्पताल बुटवल, रुपन्देही

आई.सि.यू.को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्यको शिलबन्दी बोलपत्र फाराम

ठे.नं. १६ क/०७१/०७२

फर्मको नाम :-

प्रोपाइटरको नाम/थर :-

सम्पर्क फोन/मोवाइल नं. :-

मिति:-

बिक्रि अन्तिम मिति : सूचना प्रकाशित मितिको ३० औ दिनसम्म

दाखिला अन्तिम मिति : सूचना प्रकाशित मितिको ३१ औ दिनको १२:०० बजेसम्म

खोल्ने मिति : सूचना प्रकाशित मितिको ३१ औ दिनको २:०० बजे



स्वास्थ्य तथा जनसंख्या मन्त्रालय

ब्रिविति अञ्चल अस्पताल ब्रटवल

बोलपत्र आव्हानको सूचना

सूचना प्रकाशित मिति २०७१।११।२५

यस अस्पतालको लागि आ.व. २०७१।०७२ मा **ICU** संचालन गर्नको निम्ती आवश्यक पर्ने मेशिनरी उपकरणहरु बोलपत्रका माध्यमवाट खरिद गर्नुपर्ने भएकोले निम्न शर्तहरूको अधिनमा रहि शिलबन्दी बोलपत्र पेश गर्न सम्बन्धित सबैको जानकारीको लागि यो सूचना प्रकाशित गरिएको छ ।

- १) सूचना प्रकाशित भएको मितिले ३० दिन सम्म कार्यालय समयमा तपिसलमा उल्लेखित रकम (पिछ फिर्ता नहुने गरी) तिरी यस अस्पताल बाट खिरद गर्न सिकनेछ बिकि भएका बोलपत्रहरु ३१ औं दिनको १२.०० बजे भित्र यस अस्पतालमा दर्ता भईसक्नु पर्नेछ । म्यादिभित्र प्राप्त भएका बोलपत्रहरु सोहि दिनको २.०० बजे बोलपत्रदाता वा निजको प्रतिनिधिको रोहवरमा यस अस्पतालमा खोलिनेछ । बोलपत्रदाता /प्रतिनिधि उपस्थित नभएमा पिन बोलपत्र खोल्न क्नै बाधा पर्ने छैन ।
- २) बोलपत्र फारम खरिद गर्न लिखित निवेदन साथ आ.व. २०७१।०७२ सम्मको लागि निवकरण गरिएको सम्बन्धित व्यवसाय सञ्चालन गर्न पाउने इजाजत पत्र, फर्म दर्ता प्रमाण पत्र, आयकर दर्ता प्रमाण पत्र तथा मु.अ.कर दर्ता प्रमाणपत्र, आ.व. २०७०।०७५ सम्मको कर चुक्ता प्रमाणपत्र नोटरी पब्लीकवाट प्रमाणित गराई संलग्न राखी प्रोप्राइटर स्वयं उपस्थित भै वा Letter Head मा दस्तखत नमूना सहित आधिकारिक प्रतिनिधि पठाई बोलपत्र खरिद गर्न सिकनेछ ।
- बोलपत्र फारम खरिद तथा दर्ता गर्ने अन्तिम दिन विदा पर्न गएमा सो पछि कार्यालय खुलेको पिहलो दिनलाई अन्तिम दिन मानिनेछ । बोलपत्र
 पेश गर्दा खामको बाहिर यस अस्पतालको नाम सम्बोधन गरी ठेक्का नं. र सप्लाइको नाम खुलाई शिलवन्दी गरी पेश गर्नुपर्नेछ ।
- ४) बोलपत्र दाताले मान्यता प्राप्त वाणिज्य बैंक वाट यस अस्पतालको नाममा जारी भएको १२० दिन(बोलपत्र खोल्ने दिनबाट) म्याद कायम भएको वैंक जमानत (विडवण्ड) वा <u>नेपाल राष्ट्र बैंक भैरहवा स्थित खाता नं १९०१२००.००१.००२.४२४</u> <u>कार्यालय कोड नं. ३७१३७०।०४ मा</u> नगद जम्मा गरेको सक्कलै वैंक भौचर बोलपत्र साथ संलग्न राख्नु पर्नेछ ।
- प्र) बोलपत्र दाताले आफूले कबोल गरेको प्रत्येक आइटमहरुको मु.अ.कर बाहेकको दररेटलाई अंक र अक्षर दुवैमा उल्लेख गर्नु पर्नेछ । अंक र अक्षरमा लेखेको दररेट फरक परेमा अक्षरमा लेखिएकोलाई मान्यता दिइनेछ ।
- ६) बोलपत्रदाताले दररेट उल्लेख गर्दा केरमेट गरेको ठाउँमा र प्रत्येक पानामा सिहछाप नगरेको, टिपेक्स लगाएको, बोलपत्र फारमको फोटोकपी गरी पेश गरेको यस अस्पतालवाट विकि नभएको, एक व्यक्ति वा फर्मका नाउँमा विकि भएको बोलपत्र फारम अर्को व्यक्ति वा फर्मको तर्फवाट पेश गरेको वा आफ्नो तर्फवाट कुनै शर्त राखी पेश गरेको बोलपत्रलाई मान्यता दिईनेछैन ।
- ७. कारण जनाई वा नजनाइ बोलपत्र पूर्ण वा आशिक रुपमा स्विकृत गर्ने वा नगर्ने सम्पूर्ण अधिकार यस अस्पतालमा निहित रहनेछ
- ९. उपरकरणहरुको स्पेशिफिकेसन बोलपत्र फाराममा संलग्न गरिएको छ बोलपत्रदाताले उल्लेख भएको स्पेशिफिकेशन अनुशारको मेशिनरी उपकरणहरु उपलब्ध गराउन् पर्नेछ ।
- शिलबन्दी बोलपत्र फारामका साथमा उपकरणहरुको निम्नानुसारका कागजात पेश गर्नुपर्नेछ ।

Orginal Cataloge & Details Literature

- क) Company को डिलरशिप प्रमाणको प्रमाणित प्रमाण पत्र
- ख) उक्त मेशिन बिकि पश्यात सेवा दिने सम्बन्धी प्रस्तावको विवरण (warranty period समेत) ।
- ग) सो कम्पनी र मोडलको त्यस्तै उपकरण नेपालका अन्य स्वास्थ्य संस्थामा आपूर्ति गरेको भए सो सम्बन्धी कागजात,
- घ) ISO तथा CE प्रमाणको प्रमाणित पत्र
- ११. प्रत्येक ठे.नं. को प्रत्येक आईटमहरु छुट्ट छुट्ट स्वीकृत गिरनेछ। बोलपत्र फारम साथ संलग्न स्पेसिफिकेशन अनुसारका मेसिनरी उपकरण उपलब्ध गराउनु पर्नेछ। साथै आफुले आपुर्ति गर्ने मेसिनरी उपकरण यस अस्पतालमा उपलब्ध गराई सम्बन्धीत चिकित्सक, नर्स तथा कर्मचारिहरुलाई संचालन सम्बन्धी आवश्यक तालिम दिनुपर्नेछ।
- 9२. स्विकृत बोलपत्रदाताले उपकरण तथा मेशिनहरु यस अस्पतालमा ल्याइ जडान गरिदिनुपर्ने छ । मेशिन तथा उपकरणहरु सञ्चालनका लागी आवश्यक प्राविधिक तालिम समेत गराई मेशिन सन्चालन भए पश्चात मात्र भुक्तानी गरीने छ । ढुवानी तथा जडान खर्च बोलपत्रदाता स्वयले नै व्यहोर्नु पर्नेछ ।
- १३. यस सम्बन्धमा क्नै क्रा ब्फन् परेमा कार्यालय समयमा यस अस्पतालमा सम्पर्क राख्न सिकनेछ
- १४. बोलपत्रसाथ उपकरणको क्याटलक र स्पेशिफिकेशन समेत पेश गर्नुपर्नेछ
- ৭২) बोलपत्रदाताले दर्ता भै सकेको बोलपत्रमा कुनै संशोधन गर्न वा बोलपत्र फिर्ता लिन चाहेमाबोलपत्र दर्ता गर्ने अन्तिम मिति र समय अगावै नै व्यहोरा खुलाई सिलवर्न्दी निवेदन दिनु पर्नेछ ।
- १६) यो सूचनामा उल्लेख हुने छुट भएका अन्य क्राहरु सार्वजिनक खिरद ऐन २०६३ र सार्वजिनक खिरद नियमावली २०६४ बमोजिम हुनेछ ।
- १७) बोलपत्र सम्बन्धी सूचना यस अस्पतालको वेभसाईट www.lzhospital@gov.np मा पनि हेर्न सिकनेछ।

क्र. सं.	<i>ठे. न</i> ं.	विवरण	ल.ई रकम (भ्याट बाहेक)	धरौटी	बोलपत्र दस्तुर
٩	१६ "क"/०७१/०७२	आई.सि.यू.को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्य	<u>५३,९८,२३०।०८</u>	१,६८,०००।००	୩,୦୦୦।୦୦
२	१६ "ख" /०७१/०७२	आई.सि.यू को लागि मेशिनरी उपकरणहरू आर्पुर्त गर्ने कार्य	५१,३२,७४३।३६	१,४९,४००।००	૧,૦૦૦ ૦૦
३	१६ "ग" /०७१/०७२	आई.सि.यू को लागि मेशिनरी उपकरणहरू आपुर्ति गर्ने कार्य	४८,६७,२५६।६४	१,४१,४००।००	૧,૦૦૦ ૦૦
Х	१६ "घ" /୦७१/୦७२	आई.सि.यू.को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्य	३२,४४,२४७९२	9,09,000100	૧,૦૦૦ા૦૦
x	१६ "ड" /०७१/०७२	आई.सि.यू.को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्य	२७,८७,६१०।६६	56,000l00	૧,૦૦૦ા૦૦
(se	१६ "च" /०७१/०७२	आई.सि.यू.को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्य	२७,८३,१८५।८४	द६,५००।००	૧,૦૦૦,૦૦

मेडिकल सुपरिन्टेण्डेण्ट

Specification & Statement of Compliance, Fully Automated Bio-Chemistry Analyser

Bidders are to offer a standard production model most closely matching the specification below and provide details of the offer. The offer must be for brand new equipment.

These specifications are for the **minimum requirement.** Bidders may offer higher specifications but they are to highlight these in the Statement of Compliance column. Units are to be stated using the SI system. For example length in metres (m). Multiples and sub-multiples to be used are: 10^6 mega (M); 10^3 kilo (k); 10^{-3} milli (m) and 10^{-6} micro (μ).

Bidders must enter their offered specifications against each parameter of this Technical Specifications Form (TSF), comment as necessary, and sign and stamp each page. Failure to complete this statement of compliance may result in the offer being rejected. A Bidder who enters texts such as "Yes", "Complied", "Better", "Refer to catalogue", and directly copying the Purchaser's descriptions, leaving any parameter line blank and/or submit any text or content of this nature may result in the offer being rejected.

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogu e/datash eet/man ual
	Fully Automated Bio-Chemistry			1.11
	Analyser			
	Manufacturer			
	Brand			
	Type / Model			
	Country of Origin			
1	Description of Function			
1.1	For analysis of serum, plasma, urine, cerebrospinal fluid (CSF), hemolysate,			
	whole blood and therapeutic drugs			
	(TDM), drugs of			
	abuseimmunoturbidimetricassays.			
2	Operational Requirements			
2.1	Must be open system and fully			
	computerized with random access,			
	selective multi-batch type, providing			
	maximum flexibility in programming			
2.2	Must be capable of undertaking 200			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogu e/datash
				eet/man ual
	tests/hr. involving fixed time, end			uai
	point and kinetic chemistry.			
3	System Configuration			
3.1	Fully Automated Bio-Chemistry Analyser with integrated printer and computer and with complete accessories.			
4	Technical Specifications			
4.1	Optical Requirement: Wavelength Range: 340 to 700nm Absorbance: 0.000 to 3.000A Resolution: 0.0001A or better Measurement: Monochromatic & Bio chromatic options. Flow cell volume: approx. 50µl Source of light: Halogen lamp with very long life.			
4.2	Reagent Handling System: Pre and Post dilution: Automatic Aspiration volume: 5-50µl in 0-0.5µl increments Wash Cycles: Programmable for aspiration and sampling probes			
4.3	 Analytical Requirements: Sample Tray/reaction plate: >50 positions for samples/standards/ controls Sample cups: 0.5-1ml Reaction types: End point, kinetic- differential and initial rate bichromatic, with & without blank correction Test Parameters: 50 or more, all programmable as per user requirement. Incubation Temp: 37°C preferably with variable temperature options Cuvette Temp: 37°C +0.1°C Quality control: Daily and monthly with good real time QC programme with L-J graphsprintout of QC charts & reports. Calculated and precision check 			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogu e/datash eet/man ual
	facility			
4.4	Must have self-diagnostic tests with			
	error message & online display.			
4.5	Must be programmable for all test			
	menus & state of the art workstation.			
4.6	Must have built in cooled reagent			
	compartment tomaximise			
	reagentstability			
4.7	Must have continuous samples loading facility.			
4.8	Data Management Software: The equipment shall be supplied with compatible, programme windows basecomprehensive data processing & management system. Graphicaluser interface software, user friendly, test oriented, LIMS capability. Complete back up of the date base for calibration control and patients samples result.			
4.9	PC:Latest configuration, Core i5 with			
4.10	high speed processor and with instrument operating and data management software, 19" or more LCD flat monitor, wireless keyboard and mouse, 2 GB RAM, 180 GB HDD, 128 MB Graphic card, CD/DVD read/write drive and with latest version of Operating system MS Windows, OEM software. Facility to store at least 10,000patient'sdata storage and multitasking facility oncomputer. Inbuilt printer thermal type with 40			
4.10	characters/line or better.			
4.11	Shall come with compatible laser printer.			
4.12	Shall supply reagent pack for 1000 tests with cleaning solution and quality control solution and 3 set of printer paper roll.			
5	Accessories, spares and consumables			
5.1	All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above).			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogu e/datash eet/man ual
6	Operating Environment			
6.1	The system offered shall be designed to operate normally under the conditions of the purchaser's country. The conditions include Power Supply, Climate, Temperature, Humidity, etc.			
6.2	Power supply: 220 – 240 VAC, 50Hz fitted with appropriate plug. The power cable must be at least 3 metre in length.			
6.3	Shall provide compatible servo voltage stabilizer and UPS of suitable rating with voltage regulation and spike protection for 30 minutes back-up.			
7	Standards and Safety Requirements			
7.1	Must submit ISO13485:2003/AC:2007 for Medical Devices AND			
7.2	CE (93/42 EEC Directives) or USFDA approved product certificate.			
7.3	Shall meet IEC 61010-2-081safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-081: Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes.			
8	User Training			
8.1	Must provide user training (including how to use and maintain the equipment).			
9	Warranty			
9.1	Comprehensive warranty for 2 years afteracceptance.			
10	Maintenance Service During Warranty Period			
10.1	During the warranty period supplier must ensure planned preventive maintenance (PPM) and corrective/breakdown maintenance whenever required.			
11	Installation and Commissioning			
11.1	The bidder must arrange for the equipment to be installed by certified or qualified personnel; any prerequisites for installation to be communicated to the purchaser in advance, in detail.			
12	Documentation			
12.1	User (Operating) manual in English			
12.2	Service (Technical / Maintenance)			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogu e/datash eet/man ual
	manual in English			
12.3	List of important spare parts and accessories with their part number and costing.			
12.4	Certificate of calibration and inspection from factory.			

Specification & Statement of Compliance, Fully Automated Haematology Analyser (Blood Cell Counter)

Bidders are to offer **a standard production model** most closely matching the specification below and provide details of the offer. **The offer must be for brand new equipment.**

These specifications are for the **minimum requirement.** Bidders may offer higher specifications but they are to highlight these in the Statement of Compliance column. Units are to be stated using the SI system. For example length in metres (m). Multiples and sub-multiples to be used are: 10^6 mega (M); 10^3 kilo (k); 10^{-3} milli (m) and 10^{-6} micro (μ).

Bidders must enter their offered specifications against each parameter of this Technical Specifications Form (TSF), comment as necessary, and sign and stamp each page. Failure to complete this statement of compliance may result in the offer being rejected. A Bidder who enters texts such as "Yes", "Complied", "Better", "Refer to catalogue", and directly copying the Purchaser's descriptions, leaving any parameter line blank and/or submit any text or content of this nature may result in the offer being rejected.

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/d atasheet/ma nual
	Fully Automated Haematology Analyser			
	(Blood Cell Counter)			
	Manufacturer			
	Brand			
	Type / Model			
	Country of Origin			
1	Description of Function			
1.1	Automated haematology analyser or			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/d atasheet/ma nual
	complete blood cell counter is used to count various types of blood cells in the blood.			
2	Operational Requirements			
2.1	Fully automated haematology analyser that measures 18 parameters including WBC 5-part differential is required.			
2.1	It shall be open system.			
3	System Configuration			
3.1	Fully Automated Haematology Analyser, complete unit with all standard reagents, consumables and accessories.			
4	Technical Specifications			
4.1	Parameters to be measured are: WBC, LYM%, LYM, MON%, MON, GRA%, GRA, RBC, HGB, HCT, MCV, MCH, MCHC, RDW, PLT, MPV, PCT, PDW.			
4.2	Large LCD display for all parameters, histograms, scattergrams.			
4.3	Shall have: • Histogram: WBC 5-part differential distribution, RBC distribution, PLT distribution. • Two Scattergrams.			
4.4	The instrument must have throughput of more than 60 samples per hour in all the discreteanalysis modes.			
4.5	The sample aspiration volume for the complete differential blood count must not be more than 150 µl.			
4.6	Principle of working: RBC/ Platelet count: Impedance Method/hydrodynamic focussing. Differential Parameters & Reticulocyte: Semiconductor Laser basedFlow Cytometry, fully automated. Haemoglobin Measurement: Colorimetric/ Cyanide free method.			
4.7	It shall have whole blood open vial and closed vial mode, capillary mode/ predilute mode.			
4.8	It shall have automatic start up, shut down and sample analysis.			
4.9	Multichannel analysis for better resolution.			
4.10	It shall have autoloader, 50 samples or more at a time.			
4.11	Automatic probe wipe.			
4.12	Automatic clot detection technology.			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/d atasheet/ma nual
4.13	It shall have user definable flagging system for various results.			
4.14	Various sensors must check the condition of the instrument, if any abnormality is detected, an error message be displayed so that occurrence of trouble is prevented.			
4.15	Shall have integrated thermal printer.			
4.16	Quality assurance system with calibrators & controls.			
4.17	It shall have Bar Code Generation and reader facility (External as well internal).			
4.18	On board memory for about 200-250 tests records.			
4.19	Shall have RS 232C port.			
4.20	PC: Computer with latest configuration, 19" or more flat monitor, optical mouse keyboard, CD, DVD read/write device and with latest software for storage of at least 5000 samplesdata with histogram & scattergram.			
4.21	Shall quote rates for reagents & consumables, calibrators & controls, printer paper, separately and it must be valid for at least 3 years.			
5	Accessories, spares and consumables			
5.1	Reagents & consumables, calibrators & controls, printer paper to be supplied for 1000 test.			
5.2	Shall provide compatible laser printer, 1 no.			
5.3	All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above).			
6	Operating Environment			
6.1	The product offered shall be designed to be stored and to operate normally under the conditions of the purchaser's country. The conditions include Power Supply, Climate, Temperature, Humidity, etc.			
6.2	Power supply: 220-240VAC, 50Hz fitted with appropriate plug type D (3 pins). The power cable must be minimum 3 meters long.			
6.3	Shall provide suitable servo voltage stabilizer and UPS with maintenance free batteries, voltage regulation and spike protection for minimum 30 min. back-up for the entire system including computer			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/d atasheet/ma nual
	and printer shall be supplied with the system.			
7	Standards and Safety Requirements			
7.1	Must submit ISO13485:2003/AC:2007 for Medical Devices AND			
7.2	CE (93/42 EEC Directives) or USFDA approved product certificate.			
7.3	Shall meet IEC 61010-2-081safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-081: Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes.			
8	User Training			
8.1	Must provide user training (including how to use and maintain the equipment).			
9	Warranty			
9.1	Comprehensive warranty for 2 years after acceptance.			
10	Maintenance Service During Warranty Period			
10.1	During the warranty period supplier must ensure preventive maintenance and corrective/breakdown maintenance whenever required.			
11	Installation and Commissioning			
11.1	The bidder must arrange for the equipment to be installed and commissioned by certified or qualified personnel; any prerequisites for installation to be communicated to the purchaser in advance, in detail.			
12	Documentation			
12.1	User (Operating) manual in English.			
12.2	Service (Technical / Maintenance) manual in English.			
12.3	List of important spare parts and accessories with their part numbers and costing.			
12.4	Certificate of calibration and inspection from factory.			

Specification & Statement of Compliance, Coagulometer

Bidders are to offer a standard production model most closely matching the specification below and provide details of the offer. The offer must be for brand new equipment.

These specifications are for the **minimum requirement.** Bidders may offer higher specifications but they are to highlight these in the Statement of Compliance column. Units are to be stated using the SI system. For example length in metres (m). Multiples and sub-multiples to be used are: 10^6 mega (M); 10^3 kilo (k); 10^{-3} milli (m) and 10^{-6} micro (μ).

Bidders must enter their offered specifications against each parameter of this Technical Specifications Form (TSF), comment as necessary, and sign and stamp each page. Failure to complete this statement of compliance may result in the offer being rejected. A Bidder who enters texts such as "Yes", "Complied", "Better", "Refer to catalogue", and directly copying the Purchaser's descriptions, leaving any parameter line blank and/or submit any text or content of this nature may result in the offer being rejected.

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/datas heet/manual
	Coagulometer			
	Manufacturer			
	Brand			
	Type / Model			
	Country of Origin			
1	Description of Function			
1.1	Coagulometer measures the blood clotting			
	parameters.			
2	Operational Requirements			
2.1	Microprocessor controlled system.			
3	System Configuration			
3.1	Coagulometer complete unit with printer and			
	with complete accessories.			
4	Technical Specifications			
4.1	16 incubation positions for samples (4 cells x 4 columns).			
4.2	Shall have 2 measurement channels.			
4.3	2-4 positions for reagents (one with magnetic			
	stirrer) and 2 pipette wells.			
4.4	Four independent built in timers for incubation.			
4.5	Measurement possible in plasma.			
4.6	Automatic pipette (electronically connected or			
	manual start up).			
4.7	Backlight LCD display, 4 lines of 40 characters			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/datas heet/manual
	with built in printer.			
4.8	Results in seconds and in various units (% INR,			
	Ratio, Gm. / L mg/ds, IC/ml).			
4.9	RS 232 interface.			
4.10	Incubation and measurement wells at 37°C +/- 0.5°C.			
4.11	Tests: PT, PTT, TT, FIB (Clauses and PT derived), Factor II, V, VII, VIII, IX, X, XI, XII, Fletcher, VT (Venom time), APCR, AT-III (clot), Protein C (clot), Protein S (clot), Heparin, STAT (PT/PTT).			
5	Accessories, spares and consumables			
5.1	Accessories:			
	 Double Cuvettes: 500 Pcs. Stage auto pipette: 1 Pc. each (25/50/100/200μl) Reagent Adaptor 22,5mm: 1 Pc. Reagent Adaptor 22,8mm: 1 Pc. Reagent Adaptor 24,2mm: 1 Pc. Reagent Adaptor 27,8mm: 1 Pc. Reagent Adaptor 25,2mm: 1 Pc. Stirring magnets: 4 Pcs. Reagent container 22,4mm:100 Pcs. Reagent tubes 16mm:100 Pcs. Thermal Printer with printer cable: 01 no. Thermal Paper: 10 rolls 			
5.0				
5.2	All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above).			
6	Operating Environment			
6.1	The product offered shall be designed to be stored and to operate normally under the conditions of the purchaser's country. The conditions include Power Supply, Climate, Temperature, Humidity, etc.			
6.2	Power supply: 220-240VAC, 50Hz fitted with			
	appropriate plug type D (3 pins). The power cable must be minimum 3 meters long.			
6.3	Suitable UPS with maintenance free batteries, voltage regulation and spike protection for minimum 30 min. back-up shall be supplied with the system.			
7	Standards and Safety Requirements			
7.1	Must submit ISO13485:2003/AC:2007 for Medical Devices AND			
7.2	CE (93/42 EEC Directives) or USFDA approved product certificate.			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/datas heet/manual
7.3	Shall meet IEC 61010-1 safety requirements for electrical equipment for measurement, control, and laboratory use.			
8	User Training			
8.1	Must provide user training (including how to use and maintain the equipment).			
9	Warranty			
9.1	Comprehensive warranty for 2 years.			
10	Maintenance Service During Warranty Period			
10.1	During the warranty period supplier must ensure corrective/breakdown maintenance whenever required.			
11	Installation and Commissioning			
11.1	The bidder must arrange for the equipment to be installed and commissioned by certified or qualified personnel; any prerequisites for installation to be communicated to the purchaser in advance, in detail.			
12	Documentation			
12.1	User (Operating) manual in English			
12.2	Service (Technical / Maintenance) manual in English			
12.3	List of important spare parts and accessories with their part numbers and costing.			
12.4	Certificate of calibration and inspection from factory.			

Specification & Statement of Compliance, Electrolyte Analyser

Bidders are to offer a standard production model most closely matching the specification below and provide details of the offer. The offer must be for brand new equipment.

These specifications are for the **minimum requirement.** Bidders may offer higher specifications but they are to highlight these in the Statement of Compliance column. Units are to be stated using the SI system. For example length in metres (m). Multiples and sub-multiples to be used are: 10^6 mega (M); 10^3 kilo (k); 10^{-3} milli (m) and 10^{-6} micro (μ).

Bidders must enter their offered specifications against each parameter of this Technical Specifications Form (TSF), comment as necessary, and sign and stamp each page. Failure to complete this statement of

compliance may result in the offer being rejected. A Bidder who enters texts such as "Yes", "Complied", "Better", "Refer to catalogue", and directly copying the Purchaser's descriptions, leaving any parameter line blank and/or submit any text or content of this nature may result in the offer being rejected.

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/dat asheet/manua	
	Electrolyte Analyser				
	Manufacturer				
	Brand				
	Type / Model				
	Country of Origin				
1	Description of Function				
1.1	ISE electrolyte analyser (Na+, K+, Ca++, Cl				
) for analysis of serum, plasma, urine, whole blood.				
2	Operational Requirements				
2.1	It shall be open system and based on ISE technology.				
3	System Configuration				
3.1	Electrolyte analyser with integrated printer and with complete accessories.				
4	Technical Specifications				
4.1	Microprocessor controlled electrolyte analyser with the measured parameter of Na+, K+, Ca++ and Cl ⁻ .				
4.2	Sample volume shall be less than or equal to 100ul.				
4.3	Analysing time-less than 60 seconds/test, sample throughout 50-60samples/hour.				
4.4	Shall have fully automatic calibration of all parameters.				
4.5	Maintenance free electrodes with long warranty.				
4.6	Shall have data display on built in LCD display screen.				
4.7	Shall have fully visible measuring chamber.				
4.8	Standby mode facility user controlled and automatic for economical operations.				
4.9	QC data, memory storage and calibration results.				
4.10	Shall have automatic flagging of abnormal result.				
4.11	It shall have only one reagent module for all standards and wash solutions and waste also shall be collected in the same module.				
4.12	It shall have only one cleaning reagents for electrodes and daily maintenance.				

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/dat asheet/manua
4.13	Inbuilt thermal printer for printing patient data and facility to interface withcomputer an external printer.			
4.14	It shall have a memory of at least 20 samples.			
4.15	Shall supply reagent pack for 1000 tests with cleaning solution and one quality control solution and 3 set of printer paper roll.			
5	Accessories, spares and consumables			
5.1	All standard accessories, consumables and parts required to operate the equipment, including all standard tools and cleaning and lubrication materials, to be included in the offer. Bidders must specify the quantity of every item included in their offer (including items not specified above).			
6	Operating Environment			
6.1	The system offered shall be designed to operate normally under the conditions of the purchaser's country. The conditions include Power Supply, Climate, Temperature, Humidity, etc.			
6.2	Power supply: 220 – 240 VAC, 50Hz fitted with appropriate plug. The power cable must be at least 3 metre in length.			
6.3	Shall provide compatible servo voltage stabilizer and UPS of suitable rating with voltage regulation and spike protection for 30 minutes back-up.			
7	Standards and Safety Requirements			
7.1	Must submit ISO13485:2003/AC:2007 for Medical Devices AND			
7.2	CE (93/42 EEC Directives) or USFDA approved product certificate.			
7.3	Shall meet IEC 61010-2-081safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-081: Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes.			
8	User Training			
8.1	Must provide user training (including how to use and maintain the equipment).			
9	Warranty			
9.1	Comprehensive warranty for 2 years afteracceptance.			
10	Maintenance Service During Warranty Period			
10.1	During the warranty period supplier must ensure planned preventive maintenance (PPM) and corrective/breakdown maintenance whenever required.			

S.N.	Purchaser's Specifications	Bidder's Offer	Deviation if any	Page no of catalogue/dat asheet/manua l
11	Installation and Commissioning			
11.1	The bidder must arrange for the equipment to be installed by certified or qualified personnel; any prerequisites for installation to be communicated to the purchaser in advance, in detail.			
12	Documentation			
12.1	User (Operating) manual in English			
12.2	Service (Technical / Maintenance) manual in English			
12.3	List of important spare parts and accessories with their part number and costing.			
12.4	Certificate of calibration and inspection from factory.			

Government of Nepal

Ministry of Health & Population

LUMBINI ZONAL HOSPITAL

BUTWAL

Name of work :- Supply of Equipment for ICU Location :Lumbini Zonal Hospital, Butwal Rupandehi

F/Y:- 2071/072

BILL OF QUANTITY

S.No	Description	Unit Quantity		Rate		Amount	Remark
		Ont	Quantity	In Figure	In Word	Amount	Kemai K
1	Automated Haematolgy Analyzer Blood Cell Counter	Set	1				
2	Electrolyte Analyser	Set	1				
3	Automated Biochemistery Analyser	Set	1				
4	Coagulometer	Set	1				
					Sub Total Rs		
					Add VAT @ 13 %		
					Grand Total Rs		