

#### 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 ख/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

#### नुम्विति अञ्चल अस्पताल बुटवल

#### बोलपत्र आव्हानको सूचना

सूचना प्रकाशित मिति २०७१।१९।२५

यस अस्पतालको लागि आ.व. २०७१।०७२ मा **ICU** संचालन गर्नको निम्ती आवश्यक पर्ने मेशिनरी उपकरणहरु बोलपत्रका माध्यमवाट खरिद गर्नुपर्ने भएकोले निम्न शर्तहरुको 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 वाट खिरद गर्न सिकनेछ विक्रि 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 कुनै बाधा पर्ने छैन ।
- २) बोलपत्र फारम खरिद गर्न लिखित निवेदन साथ आ.व. २०९१।०७२ सम्मको लागि निवकरण गरिएको सम्विन्धत व्यवसाय सञ्चालन गर्न पाउने इजाजत पत्र, फर्म दर्ता प्रमाण पत्र, आयकर दर्ता प्रमाण पत्र तथा मु.अ.कर दर्ता प्रमाणपत्र, आ.व. २०९०।०९ सम्मको कर चुक्ता प्रमाणपत्र नोटरी पब्लीकवाट प्रमाणित गराई संलग्न राखी प्रोप्राइटर स्वयं उपस्थित भै वा Letter Head मा दस्तखत नमूना सिंहत आधिकारिक प्रतिनिधि पठाई वोलपत्र खरिद गर्न सिकिनेव्य ।
- ३) बोलपत्र फारम खरिद तथा दर्ता गर्ने अन्तिम दिन विदा पर्न गएमा सो पछि कार्यालय खुलेको पहिलो 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 नेपाल राष्ट्र बैक भैरहवा स्थित खाता नं १९०१२००.००१.००२.४२४ कार्यालय कोड नं. ३७३७०।०४ मा 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
- उपरकरणहरुको स्पेशिफिकेसन बोलपत्र फाराममा संलग्न गरिएको छ बोलपत्रदाताले उल्लेख भएको स्पेशिफिकेशन अनुशारको मेशिनरी उपकरणहरु उपलब्ध गराउन् पर्नेछ ।
- १०. 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 आपुर्ति गरेको भए सो सम्बन्धी कागजात,
- घ) ISO तथा CE प्रमाणको प्रमाणित पत्र
- 99. प्रत्येक ठे.नं. को प्रत्येक आईटमहरु छुट्टा छुट्टै स्वीकृत गरिनेछ। 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 कर्मचारहिरुलाई 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 कुनै कुरा बुक्तनु 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 सिलवन्दी निवेदन दिनु पर्नेछ।
- १६) यो सूचनामा उल्लेख हुने छुट भएका अन्य कुराहरु सार्वजिनक खिरद ऐन २०६३ र सार्वजिनक खिरद नियमावली २०६४ बमोजिम हुनेछ ।
- १७) बोलपत्र सम्बन्धी सूचना यस अस्पतालको वेभसाईट <u>www.lzhospital@gov.np</u> मा पनि हेर्न सिकनेछ ।

### <u>तपशित</u>

क्र. सं.	ठे. नं.	विवरण	ल.ई रकम (भ्याट बाहेक)	धरौटी	बोलपत्र दस्तुर
٩	१६ <b>"क"</b> /०७१/०७२	आई.सि.यू.को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्य	४३,९८,२३०।० <u>८</u>	१,६८,०००।००	9,000100
२	<b>१६ "स्व"</b> /०७१/०७२	आई.सि.यू.को लागि मेशिनरी उपकरणहरू आपुर्ति गर्ने कार्य	५१,३२,७४३।३६	१,४९,४००।००	૧,૦૦૦ ૦૦
ą	<b>१६ "ग"</b> /०७१/०७२	आई.सि.यू को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्य	४८,६७,२४६।६४	१,४१,४००।००	૧,૦૦૦ ૦૦
γ	१६ <b>"घ"</b> /०७१/०७२	आई.सि.यू.को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्य	३२,४४,२४७।९२	9,09,000100	૧,૦૦૦ા૦૦
x	<b>१६ "ङ"</b> / <b>०७</b> १/०७२	आई.सि.यू.को लागि मेशिनरी उपकरणहरु आपुर्ति गर्ने कार्य	२७,८७,६१०।६६	56,000100	૧,૦૦૦ા૦૦
Ę	१६ <b>"च"</b> /०७१/०७२	आई.सि.यू को लागि मेशिनरी उपकरणहरू आपुर्ति गर्ने कार्य	२७,८३,१८५।८४	द्ध, <u>५</u> ००।००	૧,૦૦૦૧૦૦

मेडिकल सुपरिन्टेण्डेण्ट

## **Technical specification form:**

# Specification & Statement of Compliance, Oxygen Concentrator with Flow Splitter

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 (u).

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.

The Statement of Compliance must be substantiated with authenticated catalogue/data sheet/ manual with the page number of original catalogue/datasheet/manual of the relevant

S.N.	Purchaser's Specifications	Bidder's Offer	Devi ation if any	Page no of catalogu e/datashe et/manua
	Oxygen Concentrator			
	Manufacturer			
	Brand			
	Type / Model			
	Country of Origin			
1	<b>Description of Function</b>			
1.1	Oxygen concentratorproduces oxygen from ambient air.			
2	<b>Operational Requirements</b>			
2.1	Integrated Oxygen sensing device (OSD) measures concentration at flowmeter entrance.			
3	System Configuration			
3.1	Oxygen Concentrator set complete with Flow Splitter.			
4	<b>Technical Specifications</b>			
Ι	Oxygen Concentrator			
4.1	Output flow: max 5 LPM (Litre per minute).			
4.2	Flow meter range: 1 to 5 LPM.			
4.3	Output pressure: 60 kPa.			
4.4	Oxygen concentration: 95% +/- 3% at			

S.N.	Purchaser's Specifications	Bidder's Offer	Devi ation if any	Page no of catalogu e/datashe et/manua l
	1-3 LPM, 92% +/- 3% at 4 LPM, 90% +/- 3% at 5LPM.			
4.5	Time to reach 95% the specified			
	performance: 5 minutes.			
4.6	Four-step filtering (coarse, pre, inlet and bacterial) of air-intake.			
4.7	All filters replaceable, coarse filter washable/reusable.			
4.8	Continuous monitoring, with visual and audible alert on:  • Low and high output pressure  • Low oxygen concentration  • Oxygen monitor: amber light on the front illuminates when oxygen concentrator is below 85%. If concentration remains below 85% for morethan 15 minutes, an audible alarm sounds.  • Power failure  • Battery test.			
4.9	Temperature operating range: 20 to 60 °C.			
4.10	Sound level produced: 40 to 50 dB(A).			
4.11	Shall have 4 antistatic swivel casters, 2 with brakes and with integrated handle allows for easy moving and positioning.			
II	Flow Splitter for Oxygen Concentrator			
4.12	Five way split of oxygen flow			
4.13	Each flow can be adjusted individually via its flow meter, range: 0.125to 2 LPM (Litre per minute).			
4.14	The output nozzle can either be fit with tubing or left blank.			
4.15	Input pressure: approx. 50 to 350 kPa. Flow splitter allows precise distribution of the oxygen output of aconcentrator towards 2, 3, 4 or 5 patients, i.e. neonates and infants.			
5	Accessories, spares and consumables			
5.1	Accessories:  • 2 x Adult cannula, with 2m tubing.  • 4 x Infant/Paediatric cannula, with 2m tubing.  • 4 x New-born cannula, with 2m tubing.			

S.N.	Purchaser's Specifications	Bidder's Offer	Devi	Page no
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	3 x Connector for above.			-
	• 4 x Humidifiers.			
	• 4 x 50' tubing.			
	• 4 x tubing adapter kit.			
	• 6 x Spare coarse filters.			
	• 3 x Spare pre-filters.			
	• 3 x Spare inlet-filters.			
	• 3 x Spare bacterial-filters.			
5.2	All standard			
	accessories/consumables/parts			
	required for the proper operation of			
	the above item shall be included in			
	the offer. Bidders shall specify, in a			
	separate Excel worksheet, the			
	quantity and details of any items			
	included in this offer which have not			
	been specified in this Technical			
6	Specifications Form.			
6.1	Operating Environment The product offered shall be designed			
0.1	to be stored and to operate normally			
	under the conditions of the			
	purchaser's country. The conditions			
	include Climate, Temperature,			
	Humidity, etc.			
6.2	Power supply: 220-240VAC, 50Hz			
	fitted with appropriate plug. The			
	power cable must be minimum 3			
	metres long.			
	Power consumption, approx.: 500 W.			
7	Standards and Safety			
7.1	Requirements			
7.1	Must submit			
	ISO13485:2003/AC:2007 for Medical			
7.2	Devices <b>AND</b> CE (93/42 EEC Directives) or			
1.2	USFDA approved product certificate.			
8	User Training			
8.1	Must provide user training.			
9	Warranty			
9.1	Comprehensive warranty for 1 year			
	after acceptance.			
10	Maintenance Service During			
	Warranty Period			
10.1	During warranty period supplier must			
	ensure corrective/breakdown			
	maintenance whenever required.			
11	Installation and Commissioning			
11.1	The supplier must accomplish proper			
	installation and commissioning of			
	equipment onsite.			
12	Documentation			

S.N.	Purchaser's Specifications	Bidder's Offer	Devi ation if any	Page no of catalogu e/datashe et/manua l
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.			

## **Technical specification form:**

### Specification & Statement of Compliance, Bedside Monitor

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 ( $\mu$ ).

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.

The Statement of Compliance must be substantiated with authenticated catalogue/data sheet/ manual with the page number of original catalogue/datasheet/manual of the relevant

S.N	Purchaser's Specifications	Bidder's Offer	Deviatio n if any	Page no of catalogue /datashee t/manual
	Bedside Monitor			
	Manufacturer			
	Brand			
	Type/Model			
	Country of Origin			
1	<b>Description of Functions</b>			
1.1	A bedside patient monitor to monitor physiological parameters of patients in the critical care units or operating theatres.			
2	Operational Requirements			
2.1	It shall operate on AC power supply as well as built-in battery.			
3	System Configurations			
3.1	Monitor Patient Bedside 4 chl. colour with ECG/Resp., SpO2, NIBP, Temp, 2IBP, ETCO2.			
3.2	All accessories, consumables and etc. required for monitoring of physiological parameters specified herein.			
4	<b>Technical Specifications</b>			
4.1	High resolution colour flat panel non-reflective screen: > 10" display size			

S.N	Purchaser's Specifications	Bidder's Offer	Deviatio n if any	Page no of catalogue /datashee t/manual
	for at least 4 channel waveforms display.			
4.2	Display of up to 4 physiological parameter modules without the need for external devices.			
4.3	Display waveform: ECG, IBP, SpO2, pulse wave and respiration.			
4.4	Numeric data display: heart rate / pulse rate, respiration rate, NIBP (Systolic, Diastolic, Mean), SpO2 and current time of NIBP measurement.			
4.5	Use interaction via integrated touch screen, press pad/button or rotary knob.			
4.6	With storage of at least 24 hours of trend data in 30-second sampling resolution for all monitored parameters to be displayed graphically and in tabular form.			
4.7	Data resolution shall be minimum of 30 second sampling.			
4.8	Display of trend:			
4.9	a) Trend tables in at least with 1, 5, 15, 30 or 60 -minute display formats; and			
4.10	b) Trend graphs in at least 1, 2, 4, 8, 12 or 24 -hour display formats.			
4.11	With storage of events for event recalling, review and documentation. It shall be able to store and record at least 10 events.			
4.12	The monitor shall be protected against the interference from the electric cautery and other electrical equipment.			
4.13	Despite the technical requirements of the networking capability, the networking works shall not be included in this offer.			
4.14	All parameters modules shall work in all monitors within the network and shall be easily interchangeable by the user. There shall be no restriction on the combination of them.			
4.15	Parameter required:  ECC/Pagningtion with 5 system with			
4.16	ECG/Respiration with 5 system with cable (1 set) and complete reusable ECG electrodes for Adult & paediatric, 1 set each.			
4.17	ECG cable and patient cable 5 leads for disposable electrodes, 1 set.			
4.18	Disposable electrodes for adult, child and infant, 50 pcs each.			
4.19	Shall come with at least a 2-lead (channel) <b>ST analysis.</b>			

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	With lethal arrhythmia detection: at			
4.20	least with detection & monitoring of			
	asystole, ventricular, fibrillation, and ventricular tachycardia and			
	bradycardia.			
4.21	Pulse oximetry SpO2 with adult and			
4.21	child finger transducer, 1 each.			
4.22	SpO2 reusable sensor for infant, 1pc.			
4.23	Non-invasive blood pressure, NIBP			
4.24	with reusable NIBP Starter Kit.  NIBP connection hose, 1 set.			
4.24	NIBP cuff & tubing for both adult &			
	child (At least 2 different sizes for			
4.25	adult and 4 different sizes for child/			
	infant/ neonate).			
4.26	<b>Temperature</b> : 2 type of probes			
	required.  Core temperature probe adult, child &			
4.27	infant, 1 pc each.			
4.20	Skin Temperature probe, adult/child &			
4.28	infant, 1 pc each.			
4.29	Invasive blood pressure,IBP for			
	monitoring of 2 IBP.			
4.30	Shall come with one complete set of IBP reusable accessories.			
	EtCO2, preferably microstream but at			
4.31	least must be able to perform			
	mainstream and side stream EtCO2			
	monitoring.			
	Come with one complete set of EtCO2 flow sensor and accessories for			
4.32	mainstream and side stream			
	monitoring, 1 set each.			
	In the case of microstream system, it			
4.33	shall come with one complete set of EtCO2 flow sensor and accessories for			
	side stream monitoring, 1 set.			
	Come with internal rechargeable			
4.34	Lithium battery complete with built-in			
	charger.			
4.35	Monitor shall be operated by the battery for at least 60 minutes.			
	Come with Alarms for all monitored			
	parameters including: exceeding user-			
4.36	selectable upper and lower limits, life			
	threatening alarms, lead/ probe/ sensor			
<u> </u>	disconnection, system failure or error.			
4.37	Alarm shall have at least 3 levels: Crisis, Warning, and Advisory.			
	Alarm notification shall be given by			
4.38	Audible and Visual.			
4.39	With networking capability to			
7.37	interface with the central monitor.			

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	RS232 port with interface with			t/manual
4.40	computer.			
	System architecture shall be designed			
	such that deactivation or failure of any bedside or central station device on			
4.41	the network shall not disable, inhibit			
	or degrade communication functions			
	among any other devices in the system.			
_	Accessories, Spare Parts and			
5	Consumables			
	All standard accessories, consumables			
	and parts required to operate the equipment, including all standard tools			
5.1	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			
-	The system offered shall be designed			
6.1	to operate normally under the			
	conditions of the purchaser's country.			
	The conditions include Power Supply,			
	Climate, Temperature, Humidity, etc.  Power supply: 220 – 240 VAC, 50Hz			
6.2	fitted with appropriate plug. The			
	power cable must be at least 3 metres			
	in length.			
7	Standards & Safety Requirements			
7.1	Must submit ISO13485:2003/AC:2007 for Medical			
/.1	Devices AND			
7.2	CE (93/42 EEC Directives) or USFDA			
7.2	approved product certificate.			
7.2	Shall meet IEC-60601-1-2:2001			
7.3	General Requirements of Safety for Electromagnetic Compatibility.			
	Shall meet the safety requirements as			
	per IEC 60601-2-27:1994—Medical			
7.4	electrical equipment—Part 2:			
	Particular requirements for the safety of electrocardiographic monitoring			
	equipment.			
8	User Training			
	The Supplier shall conduct user			
	training for this equipment to enable			
	operators to use the equipment properly. The training shall include			
8.1	the use of all operational functions of			
	the equipment, as well as routine			
	checks and maintenance expected by			
	users.			

S.N	Purchaser's Specifications	Bidder's Offer	Deviatio n if any	Page no of catalogue /datashee t/manual
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 bycertified or qualified personnel; any prerequisites for installation to becommunicated 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.			

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	Qty	Unit		Rate	Amount	Remark
	Description	20	Circ	In Figure	In Word	2 Milount	
1	Bed Side Monitor	6	Set				
2	Oxygen Concentrator	4	Set				
					Sub Total Rs		
					Add VAT @ 13 %		
					Grand Total Rs		