

# नेपाल सरकार

#### स्वास्थ्य तथा जनसंख्या मन्त्रालय

## कोशी अस्पताल

बिराटनगर

दररेट पेश गर्ने बारेमा सूचना प्रथम पटक प्रकाशित मिति २०८२।०२।१६ गते विषय:-दरभाउपत्र पेश गर्ने बारे।

प्रस्तुत विषयमा सार्वजिनक खिरदिएन २०६३ को दफा४१ तथा सार्वजिनक खिरद नियमावली,२०६४को नियम८५को१(क)प्रावधान बमोजिम तपिशल बमोजिका सलग्न स्वीकृत स्फेसिफिकेसन बमोजिमको अस्पतालको बच्चाबिभाग एन.आइ.सि.यु.का तथा अन्य वार्डका लागि उपकरण खिरद गर्नुपर्ने भएकोले यो सूचना प्रकाशन भएको मितिले ७ सातौ दिनको दिन १२.० वजे भित्र यस अस्पतालको खिरद इकाइ शाखामा कार्यलय समय भित्रमा शिलबन्दी खाममा पेश गर्नु हुन सम्बन्धित सबैको लागि यो सूचना प्रकाशन गिरएको छ।

दरभाउपत्र सुचना प्रकाशन भएको सातौ दिनमा सोदिन (सार्बजनिक बिदा परेमा) वा भोलीपल्ट सो दरभाउपत्र सोही दिन अपरान्ह २.०बजे खोलिनेछ ।दरभाउपत्र दररेटपेश गर्दा उक्त फर्मको (खरिद कारोवार सिमा १०(दशलाख)यस आ.ब.मा ननाघेको हुनु पर्नेछ) । तपिशल

### क. सूचि दर्ता गर्न पेश गर्नु पर्ने कागजातहरु

- v. व्यवसाय दर्ता प्रमाणपत्र (निवकरण भएको)
- vi. व्यवसाय इजाजत प्रमाणपत्र (निवकरण भएको)
- vii. मु.अ.कर दर्ता प्रमाणपत्र ।
- viii. आ.ब. २०८०।८१ को कर चुक्ता प्रमाणपत्र

#### ख. दररेटपेश गर्नुपर्ने सामानको विवरण

S.n	Item Name and Description	Specifi cation	Name of Company/Countr y/Brand/Model	Qty	unit	Per unit Rate	Total	VAT	Total
1	Infant Warmer with sensor and Phototherapy unit(led type)	सलग्न छ		2	set				
2	ECG Machine Protable (12 channel)	सलग्न छ		2	set				
								G Total	

#### ग. दररेटको मूल्ययाङ्गन -विधि

- सामानको स्फेसिफकेसन मिलेको तथा आइटम वाइज न्यूनतम कवोल अंक पेश गर्नेबाट खरिद गरिनेछ i
- खिरद गिरने सामान अस्पतालले प्रयोग गर्न उपयूक्त हुन नदेखिएको खण्डमा सामान फिर्ता गिरनेछ ।सामानहरु खिरद गर्ने वा नगर्ने वा
- आशिकरुपमा खरिद गर्ने सम्पूर्ण अधिकार यस अस्पतालमा निहित हुनेछ ।सामानको क्याटलग,फोटो उपलब्ध गराउनु पर्ने छ ।
- दररेट पेश गर्नेको विवरण फर्मको नाम /फर्मको ठेगान :

प्रोपाईटर / संचालकको नाम :

- 1	1	
ਟਸਕ	ठेगाना	
5414	וויוויט	

मिति:

हस्ताक्षर:

ह्याप:

S.N.			ortable (12 Channel)  Bidder's Compliance sheet				
	ECG Machine, Portable (12 Channel)	Yes/No	Page No. in	Remark			
100	Manufacturer	k	Catalogue				
116	Brand	864					
	Type / Model	- A					
	Country of Origin						
_ 1	Description of Function						
1.1	ECG Machine is primary equipment to record ECG Signal in various configurations.						
2	Operational Requirements						
2.1	Portable digital ECG machine must be able to acquire all 12 Leads simultaneously.		ľ				
3	System Configuration						
3.1	Portable digital ECG machine with complete accessories						
4	Technical Specifications						
4.1	Simultaneous recording of 12 standard leads: aVR, aVL, aVF, I, II, III	— Т					
4.1	and V1-6 pre-cordials.	7					
4.2	Internal memory for data storage.						
4.3	Splash-resistant alphanumeric keyboard with function keys.			у.			
4.4	With zeroing reset, auto-base-line correction (0.5Hz) and 1mV test/calibration signal.						
4.5	Filter setting for line-frequency (50 or 60Hz).						
4.6	Continuous check on the quality of electrodes connection, audio						
4.0	visual alert on loss of signal						
4.7	Appropriately protected for operation during defibrillation.						
4.8	shows ECG-curves, heart rate, patient name and ID, time, speed and						
4.9	filter setting.  Machine Should have lead reversal Facility, it should display message.			-			
4.1	ECG should have colored waveform display for the quality of ECG waveform.			,			
	Machine Should have print preview option before taking printout.						
.12	Machine should have interpretation based on gender and age.						
.13	Front panel provides indication of system and battery status, electrode.			· · · · · · · · · · · · · · · · · · ·			
.14	Built-in or integrated high-resolution thermal printer.						
15	Number of channels printed is user selectable: 3, 6 or 12						
16	Combination of channels printed is standard and user selectable and with copy function.						
17 F	Paper speed, user adjustable: 5, 25 and 50mm/sec.						
18	Data interface: RS232 or equivalent. Self-test is performed each time the device is switched on.						
	Sensitivity, automatic or user selectable: 5, 10 and 20 mm/mV.						
Α	Accessories, spares and consumables						
	Accessories:						
	Patient cable-1 no.						
	Reusable chest electrodes, suction ball-type- 6 nos.	-					
	Extremity clamp electrodes, reusable- 4 nos.			18			

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			
The second	Bidders must specify the quantity of every new (including items not specified above).			
6	. r			
	The system offered shall be designed to stole and 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 fitted with			
6.2	Power supply: 220 - 240 VAC. Solid			
ATAMA	appropriate plug.			
7	Standards and Safety Requirements  Must submit ISO 13485:2003/AC:2007 for Medical Devices  Miss Submit ISO 13485:2003/AC:2007 for Medical Devices			
7.1	Must submit ISO 13485:2003/AC:2007 for integration of USFDA (510K) Must submit Valid CE (93/42 EEC Directives) or USFDA (510K) approved product certificate for both warmer and phototherapy unit.			
8	User Training Must provide user training (including how to use and maintain the			
	Must provide user training (including now to use and			
	equipment).			
9	Warranty confirmation of the agreemence.			
	Comprehensive warranty for 2 years after acceptance.			
10	Maintenance Service during Warranty Period  During warranty period supplier must ensure corrective/breakdown maintenance whenever required.			
11	Installation and Commissioning  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.	r		
12	Documentation	1		
12.1	- in a manual in English.			
12.2				
12.3	List of important spare parts and accessories with the		hand as	t he written Po

Note: Bidder must completely fill the Technical Specification Form (TSF). Only Yes/no/all complies should not be written. Page number in the catalogue of all the required parameters must be clearly mentioned and highlighted. Failure in doing so may lead to rejection of bid from technical committee.

En brovinda Mishra NEC NO: 79325

	Technical Specification of Infant Warmer with Sensor and Phototherapy unit (LED type)  Purchaser's Specifications (Koshi Hospital, FY: 81/82)  Bidder's Compliance Sh							
S.N	Infant Warmer with Sensor and Phototherapy unit (LED type)	Yes/No	Page No. in Catalogue	Reamr				
	Manufacturer							
	Brand							
	Type/Model							
	Country of Origin							
1	Description of Function							
	An Infant warmer is used to keep the patient's core temperature stable							
	at 37 degree C. Phototherapy units are used to treat hyperbilirubinemia		=					
	a condition characterized by high bilirubin concentrations in the blood.							
2	Operational Requirements							
	It shall be microprocessor controlled Infant warmer with manual and							
	servo options and inbuilt baby bassinet trolley. It should have							
	provision for operating with double sided phototherapy as stated in		1					
	Technical specification.	- 1						
3	System Configuration							
	Infant Warmer with Sensor, 2 unit of phototherapy either inbuilt or							
	configurable with warmer and complete standard accessories for both							
	warmer and phototherapy unit.	_						
4	Technical Specifications							
4.1	It must have facility to display skin temperature.							
4.2	It should have visually coded control panel and color-coded safety							
	alarms for simple understanding.							
	Shall have user friendly control panel with large easy to read LED							
4.3	displays for real time patient skin temperature and set temperature							
	separately and heater power.							
4.4	It should have +/- 12 Degree Continuous Bed tilting mechanism.		-					
45	In manual mode, the heater output must be directly controlled by a							
	setting on the front panel.							
4.6	The desired temperature range from 30 to 40°C or better.		100					
4.7	Observation light must be provided for observing the baby.							
1.8	Shall have standard IV pole, monitor tray and storage drawers							
	underneath the bassinet							
	It should have provision for double surface led phototherapy ( each unit			-				
1.9	from upper and lower side of bassinet ). Base for holding Phototherapy		1					
- !	unit should be strong and height adjustable.	-	1					
	Two unit of Phototherapy can be inbuilt with radient warmer or can be							
.10	configurable with provided radient warmer.If configurable, then	. 11						
- '	warmer and phototherapy unit should be able to run at the same time.							
11	Phototherapy unit Should have blue power LEDs. Life of LED should							
t	be at least 20,000 hours of use or more.							
15	Shall have alarms with audio and visual indicators for radient warmer	_						
a	s following:							
.12	Temperature high/ Low							
	The state of the s							
E	Power failure							
L	Heater failure							
11	Control board failure	1						



	All standard accessories, consumables and parts	T	_	
l	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			1
	(including items not specified above).			1.
6	Operating Environment			i i
	The system offered shall be designed to store and to	(6)	T	
6.1	operate normally under the conditions of the purchaser's country.		1	
6.1	The conditions include Power Supply, Climate, Temperature,		1	
	Humidity, etc.			
6.2	Power supply: 220 - 240 VAC, 50Hz fitted with			
25000	appropriate plug.			
7	Standards and Safety Requirements			4
7.1	Must submit ISO 13485:2003/AC:2007 for Medical Devices			
	Must submit Valid CE (93/42 EEC Directives) or USFDA (510K)			-
7.2	approved product certificate for both warmer and phototherapy unit.			
8	User Training			-
	Must provide user training (including how to use and maintain the		T -	
	equipment).			
9	Warranty			
	Comprehensive warranty for 2 years after acceptance.			
10	Maintenance Service during Warranty Period			
	During warranty period supplier must ensure			
9.4	corrective/breakdown maintenance whenever required.			
11	Installation and Commissioning		-	
	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.			

**Note:** Bidder must completely fill the Technical Specification Form (TSF). Only Yes/no/all complies should not be written. Page number in the catalogue of all the required parameters must be clearly mentioned and highlighted. Failure in doing so may lead to rejection of bid from technical committee.

NECNO: 79329