छायाँनाथ रार्गामार्गरपालिका नगरकार्यपालिकाको कार्यालय गमगढी, मुगु





कर्णाली प्रदेश, नेपाल

For PROPOSAL DOCUMENT

Procurement of RT-PCR Machine and LabEquipment with kits, reagents, accessories

Supplier's Name:
Address:
Contact No
Date:

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छायाँनाथ रारा नगरपालिका नगर कार्यपालिकाको कार्यालय

गमगढी, मुगु कर्णाली प्रदेश, नेपाल

PCR Machine, Lab setup तथा Reagent खरिद सम्बन्धि

विश्व तथा नेपालमा महामारीको रुपमा फैलिएको COVID १९ को उपचार नियन्वण तथा प्रतिकार्यका जागि PCR Machine, Lab setup तथा Reagent खरिद हालको आपतकालिन अवस्थामा सार्वजानिक बारिद ऐन, २०६३ को दफा ६६ तथा नियमावलीको नियम १४४ बर्गोजिम विशेष परिस्थितिमा सरिद गर्नु पर्ने भएकोले इजाजत प्राप्त फर्म वा कम्पनी वा आपूर्तिकताले आ.व २०७५।७६ को करचुक्ता प्रमाणपत्र, मूल्य अभिवृद्धि कर दर्ता प्रमाण पत्र तथा इजाजत पत्र र विश्व कर्ताले आपूर्ति गर्ने उपकरणको स्पेशिफिकेशन र सामान आपूर्ति गर्न लाग्ने समय समेत खुलाई स्वास्थ्य संगठनले मान्यता दिएको आधिकारिक पत्रको छायांकपि सहित यो सूचना प्रकाशित भएको मितिले ७ (सात) दिन (सातौ दिनको १:०० वजे) भित्र छार्यानाथ रारा नगरपालिकाको website. chhayanathraramun.gov.np बाट आर्थिक प्रस्ताव डक्मेन्ट डाउनलोड गरि आपुर्ति प्रस्ताव कार्यालयको इमेल ठेगाना info@chhayanathraramun.gov.np पेश गर्न हुन जानकारी गराईन्छ । सूचनामा मांग गरिएको कागजात अनिवायं रुपमा पंश गर्नु पर्ने छ । अन्यथा त्यस्ता प्रस्तावहरु मूल्यांकनमा समेत समावेश गरिने छैना यस सम्बन्धमा विस्तृत जानकारी चाहिएमा इमेलमा लेखि पठाउन हुन वा फोन नं ९५५५ ३२२३६४ वाट जानकारी लिन सिकनेछ । प्रथम पटक सूचना प्रकाशित मिति : २०७७०४।२९

प्रमुख प्रशासकीय अधिकृत

April

1. Price Quotation and PriceSchedules

Date: To: [name and address of the Purchaser] Gentlemen and/or Ladies: Having examined the Direct Purchase (DP) documents, we the undersigned, offer to supply and deliver [description of goods and services] in conformity with the said DP documents for the sum of [total amount in words and figures] or such other sums as may be ascertained in accordance with the Schedule of Prices attached herewith and made part of this Price Quotation. We undertake, if our Price Quotation is accepted, to deliver the goods in accordance with the delivery schedule specified in the Schedule of Requirements. We agree to abide by this price Quotation for a Period of 45 days from the last date fixed for submission of the Price Quotation.. Until a formal Contract is prepared and executed, this Price Quotation. together with your written acceptance thereof and your notification of award, shall constitute a binding Contract between us. We understand that you are not bound to accept the lowest or any Price Quotation you may receive. Datedthis _____dayof____ [Signature] [In the capacityof] DulyauthorizedtosignPriceQuotationfor andonbehalfof

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RT-PCR Machine and Lab Equipment with kits, reagents,accessories

1 . Price Schedule



				Unit Price in NRs.	2	
	Unit	Quantity	In Figure	In Words	Total	Rem
	set	1			Frice	ark
Bio safety Cabinet Class II		1				
PCR Cabinet or Laminar Flow						
Auto Clave Horizontal		-				
Micro Centrifuge (High Speed)		-				
Refrigerated Centrifuge (High Speed)		-				
	,	1				
		-				
	-	1			/	
Multi-Channel Pipette		w				
Pipette Variable different size (2 each)		10				
		-				
Supply & Installation of Inverter type(wall mount) Air Conditioner 2 ton with all complete work		4				
	Description Machine v Cabinet Class II inet or Laminar Flow ve Horizontal ntrifuge (High Speed) ted Centrifuge (High lixture 20 30 aniable different size (2 trifuge Installation of Inverter ir Conditioner 2 ton omplete work		Chuit Qu	Unit Quantity	Unit Quantity In Figure Office III 1	Unit Quantity In Figure Outcome in Figure set 1

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ACK Quantity ize (500 Test 500 Test	000	1000 vial	1	1	-	,	-	-		-	-		-	1	9	3	10
5 S	Real Time RT-PCR Kits for SARS Cov-2	RNA extraction reagent for manual	3ml VTM (with 2 ewole) one feet	each sample	Pipette Rack	Filter tips with Rack 0.2-10ul (96*10)	Filter tips with Rack 20-200ul	Filter tips with Back 10-1000	inocol of warming of	Storage Rack For microcentrifuge tubes 1.5- 2ml	Neclease free eppendrof tuhe /	microcentrifuge tube 2ml (1000PCs)	Neclease free eppendrof tube /	microcentrifuge tube 0.5 ml (Pcs)	PCR tube (0.2ml) (1000PCS)	PCR strip (0.1ml) (8*125)	Zip lock bag plastic (large)	(Toopes)	(100PCs)
	41	15	16	2	17	18	19	00	24	21	22		23		24	25	26		17

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10	vo	200	2	20	10	20	ount	%	ding
Autoclavable Biohazard bag (100pcs)	Almunium foil	falcon tube 15ml	Parafilm	Cryobox	Nitrite gloves-M	Analytical Ethanol 99.99% (500ml)	Total Amount	VAT 13%	Total Including VAT
28 A	29 A	30 fa	31 P	32 C	33 N	34 A			

Total Price (in words)	
Bidder's Name:	Signature of Bidder
	-Address:
	Contact No

जि. प्रमुख प्रशासकीय प्रमुख



Schedule of Requirements

Bidders Offer		
Delivery Schedule	Within 10 days from Contractsign.	
Place of Delivery	Chhayanath Rara Municipality Gamgadhi, Mugu	
Uni t	Set	
Ite m	RT-PCR Machine and Lab Equipment with kits, reagents, accessories	
S.NO.	-	



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Evaluation and Qualification Criteria

Evaluation Criteria

- a) Delivery schedule: Supplier should supply & installation of this package. As per Schedule of Requirement.
- b) Supplier should submit the technical specifications (technical data sheet) of this package with the proposaldocument.



Technical specifications Real Time PCR System

S.N	Purchaser's Specifications	Bidder's	Compliance Sh	eet
•	Real Time PCR System	Yes/N o	Page No. in Catalogue	Remarks
	Manufacturer:			
	Brand:			
	Type/Model:		-	
	Country of Origin			
1	Description of Function			
1.1	Real Time PCR is an instrument that employs precise temperature control and rapid temperature changes to conduct the polymerase chain reaction (PCR) with Real-time Amplification of DNA/RNA from purified samples.			
2	Operational Requirements			*
2.1	Real Time PCR Thermal Cycler Mechanism system along with micro centrifuge tubes and PCR tubes.	ā.	· · · · · · · · · · · · · · · · · · ·	
3	System Configuration			
3.1	Real Time PCR Thermal Cycler Mechanism system with automatic DNA extraction / detection System			
4	Technical Specifications for Real Time PCR			
4.1	Sample Layout: Should be flexible for the tube input volumes having capability of running at least 72 or 96 samples. System must berotor based system or plate based system.			
4.2	Uniformity: Should have uniform temperature distribution. Should maintain optical detection in all wells for maximum uniformity.		*	
	HRM: SystemshouldhaveHRMtoolwithstatisticalan alysissoftware. Should have high resolution SNP screeningcapability.			
4.4	Chemistries: Should come with standard chemistries for gene expression, quantification, miRNA mutation scanning, genotyping, methylation studies etc. Should also be an open system for other kit suppliers.			
4.5	Optical detection: At least 6 filtered Photodiodes			
	Emission: At least 5 targets.			
	LightSourceLifespan: Warrantyonlightsourceformin imum20years			
4.8	Excitation Multiplexing: Minimum upto 5 targets			
	Reaction Volume: 5 μL - 150 μL	V		
	Performance Temperature accuracy: ±0.5°C			6 11

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				1
4.12	Channels: RealtimePCRcyclerandHRMinstrumentwith5channels.			
4.12				
4.13	Detection Device: Photomultiplier Tube (PMT) for higher sensitivity with sensitivity control.			20
1 11	Excitation/emission wavelengths range: 450-715nm			
	Dynamic range: 10 orders of magnitude			
	RampingRate(peak): fastramping of 5°Co10°Csec			2
	Sensitivity: Single copy gene			
		, , , , , , , , , , , , , , , , , , , ,		
4.10	Contamination Protector: The system must have contamination protector so as to avoid DNA			
	carryover contamination.			* .
4 19	Software: Should have user friendly software for			
7.17	data Analysis			
5	Computer Requirement:			
	Laptop Configuration: Must be supplied with suitable			al .
5.1	i5 6 th Generation, 500GB laptop and printer.			
6	Accessories, Spares and Consumables			
	All standard accessories, consumables and parts		10	H 10
	required to operate the equipment, including all			
6.1	standard tools and cleaning and lubrication			,
0.1	materials, to be included in theoffer. Bidders must			
	specify the quantity of everyitem			27
	included in their offer (including items not specified			2
7	above).			
7	Operating Environment			
7.1	3KVA online UPS Backup of suitable rating for at least 60 minutes to be supplied for the entire system.		*	
	The system offered shall be designed tostore and			10
7.2	to operate normally under the conditions of the purchaser's country. The conditions include Power			
	Supply, Climate, Temperature, Humidity, etc.		*	
	Power supply: 220-240V/ 50 Hz AC Single phase			
	fitted with appropriate plugs to meet			
7.3	fitted with appropriate plugs to meet purchaser's country			
	requirements. The power cable must be minimum 3 metr			=
	es			2
	long.			
8	Standards and Safety Requirements			
8.1	Must submit ISO 9001 or ISO 13485 for medical			,
	devices			
8.2	Must submitFDA/CE &CE-IVD certified approved		P	
0.0	product certificate.			
	Must submit Validmanufacturer authorization letter.			
9	User Training			
9.1	Must provide user training (including how to use			
40	and maintain the equipment).			
	Warranty Community warranty for 2 years			
	Comprehensive warranty for 2 years Maintenance Service During Warranty Period	s		
11	Maintenance Service During Warranty Period			

11.1	During the warranty period supplier must ensure corrective/breakdown maintenancewheneverrequired.			
11.1	corrective/breakdown maintenancewheneverrequired.			
12	Installation and Commissioning			
12.1	Supplier must accomplish proper installation &commissioning onsite.			
12.2	Application training must be given on site and minimum 10 test to be done in the lab at the time of training and installation			
13	Documentation			
13.1	User (Operation) manual in English.			*
13.2	Service (Technical / Maintenance) manual in English.			
13.3	List of important spare parts and accessories with their part number and costing.		z.	
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NOTE: -Bidder must completely fill the Technical specification form(TSF). Only Yes/no/all complies should not be written. Page number in 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.

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$\underline{\textbf{1.}} \quad \underline{\textbf{TechnicalSpecification of Biological Safety Cabinet (BSC)}}$

S.N.	Purchaser's Specifications 90	Bidder's Offer	Yes	No	Pg No of Catalog
	BSC Class II type A2				
	Manufacturer				
	Brand				1
	Type / Model				
	Country of Origin				
1	Description of Function				
1.1	BSC Class II used for safely working with materials contaminated with or potentially contaminated with pathogens requiring a defined biosafety level with integrated temperature-compensated airflow monitoring system.				
2	System Configuration				
2.1	BSC Class II type A2 with complete accessories ensuring both the safety of sample and environment.				
, 3	Technical Specifications				
3.1	External Dimensions (W x D x H) with optional base stand Size: 136*79*220 (cms) Approx.				
3.2	Internal Work Area Dimensions (W x D x H) Size: 116*61*68 (cms) Approx.			2	a.
3.3	Tested Opening: 203 mm (8") Approx.				
3.4	Material: Main Body Should be made of mild steel duly epoxy coated powder and inner chamber is totally made of joint less stain less steel 304 grade.				e e
	Main filter: High-efficiency Particulate Air (HEPA) filter with 99.999% efficiency at 0.3 micron				
3	Light and UV Lamp with LED indicator, LCD Display, UV Timer	o e			
	Motorized front window consist of frameless laminated UV resistance shattered proof glass door.				

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3.8	Low heat and low energy consumption			
3.9	offering 70% more energy savings Average airflow velocity (Inflow) of 0.53			
3.7	m/s (105 fpm) at initial set point			
3.10	Air flow system is 70% air recirculation and			
;	30% exhaust.			
3.11	Noise Level: 55 dB Approx.			
3.12	Audio visual alarms			
3.13	Power: 220-240V/ 50 Hz AC with appropriate plug to meet purchaser's country requirements.			
3.14	Normal Power Consumption.			·
3.15	Angle arm rest for comfortable working position			
4	Accessories, spares and consumables		-	:
4.1	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).			
5	Operating Environment	8		
5.1	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.			
5.2	Power supply: 220-240V/ 50 Hz AC Single phase fitted with appropriate plug to meet purchaser's country requirements.			
6	Standards and Safety Requirements			
6.1	Must submit CE and ISO 14644-1 Class 3 and			
	quality			
7	User Training and Technician Training			
7.1	The Supplier shall conduct onsite and offsite			× .
	user and Biomedical technician training for			
	this equipment to enable operators and			
	Biomedical technician to use the equipment	¥		2
	properly. The training shall include the use of			,
	all operational functions of the equipment, as			





	well as routine checks and maintenance				
	expected by users and Biomedical				a.
	Technician.				
8	Warranty				
8.1	Comprehensive warranty for 2 years.				
9	Maintenance Service during Warranty Period				*
9.1	During warranty period supplier must ensure corrective/ breakdown maintenance whenever required.				
10	Transport, Installation and				100
10.1	The bidder must arrange for the equipment to be installed and commissioned by certified or qualified personnel at different 5 laboratorieslocated				
11	Documentation				
11.1	User (Operating) manual in English				
11.2	Service (Technical / Maintenance) manual in English				
11.3	List of important spare parts and accessories with their part numbers			2	
11.4	Company is must.			-	g.
	Note: Bidder must completely fill the Techn complies should not be written. Page parameters must be clearly mentioned so may lead to rejection of bid from to	number in the ca d and highlighted	talogue of a I. Failure in	ll the re	Yes/no/all quired

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<u>i.</u> <u>Technical Specification of PCR Cabinet or Laminar flow</u>

S.N.	Purchaser's Specifications	Bidder's Compliance
	PCR Cabinet or Laminar flow	
•	Manufacturer	
	Brand	
	Type / Model	
	Country of Origin	
1	Description of Function	
1.1	PCR cabinet used for safely working with potentially contaminated with pathogens requiring a safety of contamination less work with safety of operator.	
2	System Configuration	
2.1	PCR cabinet with complete accessories ensuring both the safety of sample and environment with interlock function.	
3	Technical Specifications	
3.1	External Dimensions (W x D x H) size: 1500 x 695 x 1700 (mm) approx. or equivalent size.	
3.2	Internal Dimensions (W x D x H) Size: 1400 x 580 x 520 (mm) approx. or equivalent size.	
3.3	Airflow Velocity: 0.2 – 0.5 m/s with adjustable speed	
3.4	Main filter: High-efficiency Particulate Air (HEPA) filter with 99.999% efficiency at 0.3 micron.	
3.5	Motorized front window consist of frameless laminated UV resistance toughened glass	
3.6	Light and UV Lamp with LED indicator,	, v
3.7	Illuminating lamp should be LED and Illumination should be≥500Lux.	
3.8	LCD or LED Display.	
3.9	Noise Level: ≤65dB Approx.	
3.10	Material: Work zone should be made of stainless steel, Main body: cold-rolled steel with anti-bacteria powder coating.	
	Power: 220-240V/ 50 Hz AC with appropriate plug to meet	
3.11	purchaser's country requirements.	
4	Accessories, spares and consumables	
4.1	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. Biddersmust	
	specify the quantity of every item included in their offer	

5	Operating Environment		
5.1	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.		
5.2	Power supply: 220-240V/ 50 Hz AC Single phase fitted with appropriate plug to meet purchaser's country requirements.		
6	Standards and Safety Requirements		
6.1	Must submit CE and ISO 14644-1		13534946
7	User Training		
7.1	Must provide user training to the Staff (including how to use and maintain the equipment).		
8	Warranty	9	
8.1	Comprehensive warranty for 1 years.		
9	Maintenance Service during Warranty Period		
9.1	During warranty period supplier must ensure corrective/ breakdown maintenance whenever required.		
10	Transport, Installation and Commissioning		
10.1	The bidder must arrange for the equipment to be installed and commissioned by certified or qualifiedpersonnel.		=
11	Documentation		
11.1	User (Operating) manual in English		
11.2	Service (Technical / Maintenance) manual in English	***	
11.3	List of important spare parts and accessories with their part numbers and costing.	***************************************	
11.4	Authorization Letter from the Company is must.	-	





i. Technical Specification of Autoclave(50L)

S.N.	Purchaser's Specifications	Bidder's Offer	Deviatio n if any	Page no. of catalogue/ datasheet/ manual
	Autoclave			
	Manufacturer			`
	Brand			
-	Type / Model			
	Country of Origin			
1	Description of Function			
1.1	Autoclaves are required for sterilizing an objectin high temperature and high pressure steam.			
2	Operational Requirements			
2.1	Vertical autoclave, universal basic version for microbiological standard laboratory to sterilize liquids, instruments, glassware, plastic articles or general infectious waste.			

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S.N.	Purchaser's Specifications	Bidder's Offer	Deviatio n if any	Page no. of catalogue/ datasheet/ manual
3	System Configuration			
3.1	Autoclave with complete accessories			
4	Technical Specifications			
4.1	Triple walled construction; chamber, door, doorframe, bolts made ofcorrosion-resistant material and able to prevent stresscracking.			
4.2	Compact, portable easily moveable on non-rusting, non-marking castors from one place to another place. The wheels/castors shall have brakes.			
4.3	Sterilizing For water, culture media, reagents and other fluids. After completion and cooling to a selected temp., air is expelled automatically through the exhaust valve. Sterilizing temp.: 115°C to 135°C Timer: 1 to 300 min. Exhaust temp.: 0°C to 45°C			
4.4	Instrument Sterilizing For flasks, beakers, test tubes, other lab instruments. When completed, the exhaust valve opens and the temp. drops to 100°C. Thus, cool down period can be shortened. Suitable for equipment that can withstand sharp drops in pressure and for sterilizing waste. Sterilizing temp.: 115°C to 135°C Timer: 1 to 300 min.			
4.5	Sterilizing/Keep Warm After sterilizing culture media, reagents and other liquids, and cooling down naturally to a selected temp., air is expelled automatically from the exhaust valve. High temp. prevents solidifying. Sterilizing temp.: 115°C to 135°C Timer: 1 to 300 min.			

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S.N.	Purchaser's Specifications	Bidder's Offer	Deviatio n if any	Page no. of catalogue/ datasheet/ manual
	Exhaust temp.: 0°C to 45°C Incubation temp.: 45°C to 60°C			
4.6	Melting/Keep Warm		 	
4.6	To melt or keep culture media at			
	a fixed temp. (This function is not			
	for sterilizing but prevents			*
	solidifying).			
	Melting temp.: 60°C to 114°C			8
	Timer: 0 to 300 min., 72 hrs.			
	Incubation temp.: 45°C to 60°C			
4.7	Chamber volume: ≥50 litres.			
4.8	Exhaust tank: 2-liter polyethylene tank			
4.9	Chamber material: SUS304 (Austenitic stainless steel)			
4.10	Keep warm timer: 72 Hrs. Fixed			
4.11	Program Timer: 1 week (Designation: Year,			
-	month, day, hour and minute)			
4.12	Fast safety lid lock.			
4.13	Lid lock by a circumferential, durably heat- and pressure-resistant seal.			
4.14	Control lock-out switch that prevents starting a cycle if the door is not locked safely.			
4.15	Control that prevents opening the door until chamber is depressurized.			
4.16	Temperature-dependent door-locking system according to international standard.			
4.17	Maximum operating pressure: 0.240MP bar. Maximum operating temperature: 135 °C			
4.18	Sterilisation timer: 1–300 minutes.			
4.19	Instrument sterilization timer: up to 72 hours.			
4.20	Melting timer: 1–300 minutes.			
4.21	Exhaust valve open temperature setting			
4.22	Microcomputer control system.			
4.23	The control panel to be mounted so that the			,
	components sensitive to steam and heat are			- E
	protected.			
4.24	display showing:			
4)	• temperature			
	• steampressure			
	sterilizationtime			
	stage ofcycle			



S.N.	Purchaser's Specifications	Bidder's Offer	Deviatio n if any	Page no. of catalogue/ datasheet/ manual
	 alarminformation 			
4.25	Lid interlock.		- X1 - X	
4.26	Alarm: audible, with display on dysfunction.			
4.27	All information on alarm to be in full writing and not based on a code.			
4.28	Safety devices: Pressure safety valve, over- temperature limiter, anti-scorch limiter, door interlock, over-pressure limiter, current fuse			
4.29	Pressure vessel type: Small-scale pressure vessel			
4.30	A manual control that can run a complete cycle manually in case of system failure.			
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 must be designed to store and be operated normally under the condition of the purchaser's Country. The conditions include Power supply, Climate, temperature and relative humidity.			
6.2	Power supply: 220-240V/ 50 Hz AC Single phase or 380-400V AC 50 Hz Three phase fitted with appropriate plugs to meet purchaser's country requirements. The power cable must be minimum 3 metres long			
7	Standards and Safety Requirements			
7.1	Must submit ISO 9001 and CE			
8.1	User Training Must provide user training (including how to use and maintain the equipment).			
9	Warranty			
9.1	Comprehensive warranty for 1 years.			
10	Maintenance Service during Warranty Period			
10.1	During warranty period supplier must ensure corrective/breakdown maintenance whenever required.			

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S.N.	Purchaser's Specifications	Bidder's Offer	Deviatio n if any	Page no. of catalogue/ datasheet/ manual
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 becommunicated to the purchaser in advance, in detail.	z		
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.			

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

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ii. Technical Specification of Micro Centrifuge

S.N.	Purchaser's Specifications		ce Sheet	
5.11	Micro Centrifuge	Yes/No	Page No. in Catalogue	Remarks
	Manufacturer			
	Brand			
	Type / Model			
	Country of Origin			
1	Description of Function			
1.1	Micro centrifuge is a piece of equipment, generally driven by a motor that puts an object in rotation around a fixed axis, applying force perpendicular to the axis. The centrifuge works using the sedimentation principle. Where the centripetal acceleration is used to separate substances of greater and less density.			
2	Operational Requirements			
2.1	Lightweight and Compact in size.			
3	System Configuration			2
3.1	Micro centrifuge with digital display. The centrifuge bodyis made of high quality steel, stainless steel chamber,			
	safe and reliable.			
4	Technical Specifications Matthews May Speed 16 000 PPM			
4.1	Must have Max Speed 16,000 RPM		я.	
4.2	RCF 17940 x g			
4.3	Must be maintenance free brushless motor.			
4.4	Must have Acc / Dec of at least 10 types.		1	
4.5	LCD display for RCF, Time and Speed.			
4.6	Micro controller based program			
4.7	Hold at least 12 tubes of 1.5 / 2.0 ml.			
4.8	Timer up to 0 ~ 99min 59sec			
	Electric lid lock, super speed, imbalance protection.			
5.0	Noise level shall be less than 55dB			
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 theoffer.			
6	Operating Environment			
6.1	The system offered shall be designed to store 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 - 240 VAC, 50Hz fitted with appropriate plug. The power cable must be at least 3 metre in length.			
		2	1 AND	<u> </u>

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7	Standards and Safety Requirements		
7.1	Must submit ISO13485:2003/ AC:2007 AND		
7.2	CE approved product certificate.		
8	User Training		
8.1	Must provide user training (including how to use and maintain the equipment).		
9	Warranty		
9.1	Comprehensive warranty for 1 years after acceptance.		
10	Maintenance Service During Warranty Period		
10.1	During the warranty period supplier must ensure corrective/breakdown maintenance wheneverrequired.	e n o	
11	Installation and Commissioning	3 L	
11.1	Supplier must accomplish proper commissioning of the equipment on site.		
12	Documentation		
12.1	User (Operation) 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.	8 9 1	,
12.4	Certificate of calibration and inspection from factory.		

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

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iii. Technical Specification of Refrigerated Centrifuge

S.N.	Purchaser's Specifications	Bidde	er's Complia	nce Sheet
	RefrigeratedCentrifuge	Yes/NO	Page No. in Catalogu e	Remarks
	Manufacturer			
	Brand			,
	Type / Model			
	Country of Origin			
1	Description of Function			
1.1	Centrifuges are required in the Laboratory to separate various components of Blood and any other liquid sample for analysis.			
2	Operational Requirements			
2.1	Refrigerated centrifuge is melt housing, safe & compact, it is of microprocessor control, less noise & fast cooling			
2.2	Table top version, maintenance free brushless motor.		5 H V	
3	System Configuration			
3.1	Centrifuge complete with Swing out or angle rotors.			
4	Technical Specifications		20 10	
4.1	Tube Capacity: No. 12: Size 1.5 – 2 ml			
4.2	Must be micro controller based program.			
4.3	Must have LCD display for RCF, time and speed.			2
4.4	Must be made of strong fabricated & corrosion resistant steel inside.			
4.5	Timer up to 0 ~ 99hr 59min			
4.6	Must have safe lid lock with insert alarm & over-speed protection			
4.7	Maintenance-free brushless drive motor with exact speed pre selection and display.			
4.8	Max speed 16,000 RPM and Max RCF 17940 × g with speed accuracy ±20 RPM			
4.9	Temperature range –20 °C to 40°C		AMBER	
5.0	Noise level shall be less than 55dB			
5	Accessories, spares and consumables			
5.1	Accessories: 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			
. 20	offer (including items not specified above).			



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6	Operating Environment		
6.1	The system offered shall be designed to store 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 - 240 VAC, 50Hz fitted appropriate plug. The power cable must be at least 3 metre in length.		
7	Standards and Safety Requirements		х х
7.1	Must submit ISO13485:2003/ AC:2007 for Medical Devices AND		
7.2	CE (93/92 EEC Directives) approved product certificate.		
8	User Training		
8.1	Must provide user training (including how to use and maintain the equipment).		
9	Warranty	 :	
9.1	Comprehensive warranty for 1 years after acceptance.		
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 and commissioned bycertified or qualified personnel; any prerequisites for installation to becommunicated to the purchasers in advance, in detail.		
12	Documentation		
12.1	User (Operation) manual in English		
12.2			
12.3	List of important spare parts and accessories with their part number and costing.		
12.4	Certificate of calibration and inspection from factory.		7

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

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iv. Technical Specification of Vortex Mixer

S.N.	Purchasers Technical Specification	Bidder's Offer
2	Vortex Mixer	
	Manufacturer:	
	Brand:	
	Model:	
	Country of Origin:	
	Description ofFunction	
1.	A vortex mixer, or vortexer, is a simple device used commonly in laboratories to mix small vials of liquid. Designed for mixing liquids for samples and chemicals.	
2.	Operational Requirements: A vortex mixer with speed changeable from regulator knob provided on control panel.	
3.	SystemConfiguration: Vortex mixer with complete accessories for mixing different tubes and vials.	
4.	TechnicalSpecifications: Shaking Movement should be orbital	
	Orbital diameter should be at least 4 mm	
	Motor type shall be shaded pole motor	
20	Permissible ON time shall be 100 % power of 30 mins	
	Speed range shall be from 0 – 2500 RPM	
	Run type shall be continuous or touch operation	
9	Dimension shall	
	Tube adaptor shall have at holes for mixing tubes of 10mm diameter	
	Operation mode selection should be through bi-directional switch	
5.	Accessories, spares and consumables	
-	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: The product offered shall be designed to be stored and to operate normally under the conditions of the purchaser's country. The conditions	

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	include Power Supply, Climate, Temperature, Humidity, etc.	8
	Power supply: 220-240V/ 50 Hz AC Single phase fitted with	
	appropriate plug to meet purchaser's country requirements. The power	
	cable must be	
	minimum 3 meters long.	
2	Certification:	
	Must submit ISO 13485:2003/AC:2007 for Medical Devices AND	
7	CE (93/42 EEC Directives) approved product certificate.	
	User Training:	
- T	The Supplier shall conduct onsite user training for this equipment to	
8	enable operators to use the equipment properly. The training shall	
	include the use of all operational functions of the equipment, as well	
	as	
	routine checks and maintenance expected by users.	
	Warranty:	
9	Comprehensive warranty for 1 years after acceptance.	
	Maintenance Service During Warranty Period:	
10	During the warranty period supplier must ensure planned preventive	
	maintenance (PPM) along with corrective/breakdown maintenance	
	whenever required.	
	Installation and Commissioning:	
11	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.	
	Documentation:	
12	User (Operating) manual in English.	
7	Service (Technical / Maintenance) manual in English.	
	List of important spare parts and accessories with their part number and	× , a
	costing.	
	Certificate of calibration and inspection from factory.	





V. Technical specification of Deep Freezer (-20°C)

S.N.	Purchaser's Specifications		Bidder's Offer			
·	Laboratory Freezer (-20 ° C)	Yes	No	Page No. in Catalogue	Remarks	
	Manufacturer:					
	Brand:					
	Type/Model:				¥	
	Country of Origin:					
1	Description of Function					
1.1	Deep Freezers or Laboratory freezers are required to preserve blood and blood products, vaccines, plasma etc. at specified temperatures.					
2	Operational Requirements					
2.1	Microprocessor controlled frost-freeFreezer					
3	System Configuration					
3.1	Ultra Low Deep Freezer (-20 °C), Vertical type.					
4	Technical Specifications					
4.1	A microprocessor controlled upright -20 0C deep freezer					
4.2	Capacity: Approx 100 litres.	10			2 2	
4.3	Freezer construction:					
,	Outer panels and interior panels shall be made of corrosion resistant material, preferably stainlesssteel.					
4.5	Door:					
	Locking door supplied with minimum two keys.					
4.6						
	High quality lockable castors for easy mobility.					
4.7						
	Adjustable trays/racks (preferably stainless steel) with perforated design.		e.			
. "	(Bidder to indicate the number off tray/racksoffered.)					
4.8	Alarm:					
	Audio-visual high & low temperature, door open/lock alarm.					
4.9	To be supplied complete with:			ancie.		
	Mains electric, voltage stabilizer unit. Unit is to stabilize power supply, along with UPS minimum 1 hour backup.				,	
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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 be stored and to operate normally under the conditions of the purchaser's country. The conditions include Power Supply, Climate, Temperature, S.N. Purchaser's Specifications, Humidity, etc.				
6.2	Power supply: 220–240VAC, 50Hz fitted withappropriate plug type D round 3 pins. The power cable must be at least 3 metre inlength.			-	
7	Standards and Safety Requirements				
7.1	Must submit ISO 9001 or ISO 13485:2003/AC: 2007 AND				a.
7.2	CE or USFDA approved product certificate.				
8	User Training				
8.1	Must provide user training (including how to use and maintain the equipment).				
9	Warranty				
9.1	Comprehensive warranty for 1 year and extra 1 year free AMC.				
10	Maintenance Service During Warranty Period				
10.1	During the warranty period supplier must ensure corrective/breakdown maintenance wheneverrequired.				
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.	2			4
12	Documentation				
12.1	User (Operating) manual in English and/or				
	Service (Technical / Maintenance) manual in English				- 1
12.2	List of important spare parts and accessories with their part numbers and costing available in stock with				
	the supplier.				

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.

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Vi. Technical specification of Laboratory Freezer upright, Low (-80 °C)

S.N.	Purchaser's Specifications	Bidder's Offer
	Laboratory Freezer upright, Low (-80 ° C)	
2	Manufacturer	
	Brand	
	Type/Model	
	Country of Origin	
1	Description of Function	
1.1	Deep Freezers or Laboratory freezers are required to	
1.1	preserve blood and blood products, vaccines, plasma	y .
	etc. at specified	
	temperatures.	
2	Operational Requirements	
2.1	Microprocessor controlled frost-freeFreezers,	
2.2	Separate chamber racks that can be pulled out for easy	
2.2	handling	
2.3	User friendly, non-CFC refrigerant	
3	System Configuration	
3.1	Ultra Low Deep Freezer (-80 °C):	
	CFC freerefrigerant	
	 Microprocessor controlled 	
	Alarmfacility	
4	Technical Specifications	
4.1	A microprocessor controlled upright -80 °C deep freezer.	
4.1	Capacity: approximately 200 litres or more.	
4.2	Refrigeration system: CFC-free refrigerant cooling	
4.3	system.	
4.4	Freezer construction: Outer panels and interior	
	panels shall be made of corrosion resistant material,	
	preferably stainless steel.	
4.5	Door: Foam sealed door system or vacuum insulated	
	glass door.	
4.6	Insulation: High density polyurethane foam.	
4.7	Castor or wheels: High quality lockable castors or omni	
	wheel for	
4.8	easy mobility. Shelves: 2 shelves must be available	
4.9	Alarm: Audio-visual high & low temperature, door	
4.9	open/lock	
*	alarm.	
4.10	Condenser: EBM condenser fan motor.	
5	Accessories, spares and consumables	
5.1	All standard accessories, consumables and parts	
J.1	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).	2
6	Operating Environment	. \



S.N.	Purchaser's Specifications	Bidder's Offer
6.1	The system 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 round 3 pins. The power cable	*
	must be at least 3 metre in length.	
7	Standards and Safety Requirements	
7.1	Must submit ISO 9001 or ISO 13485:2003/AC: 2007 AND	
7.2	CE (93/42 EEC Directives) approved product certificate.	
8	User Training	
8.1	Must provide user training (including how to use and	
	maintain the	
	equipment).	
9	Warranty	
9.1	Comprehensive warranty for 1 years.	
10	Maintenance Service During Warranty Period	
10.1	During the warranty period supplier must ensure	
29	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	9
ä	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 available in stock with the supplier.	on form(TSE) Only Ves/no/all complie

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



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VII. Technical Specifications of Micropipette Single and multi channel

S. No.	Specifica	ation Required Quantity	Bidder's Offer
	Manufacturer:		-
2	Country of Origin:		
	Model:		
	Brand:		3
1	Description of Function		5
	Laboratory Micro pipette to use for	lab sampling preparation.	
2	Operational Requirements		
<u> </u>	Different size autoclavable micropip	репе	
3	System Configuration		
	Single channel micropipette		1 0
1	8 channel micropette Technical Specification		
4.1	Ingle Channel Micro Pipette		
4.1	Fullyautoclavable		
	Ergonomic design provides	excellentonerating	
	experience	excelentoperating	
	Easy-to-read volume displa	v	
	• Easy calibration andmainte		
	 provides excellent operating 		22
	Large display window allow		
	identification	,	
	Easy calibration andmainter	enance	
4.1.1	Micropipette • SingleChanne		
	• Capacity: 0.1	-2.5µl 5	
	• Increment:0.	5µl	
	• Inaccuracy%	s: At 2.5 μl : 2.50, At 1.25 μl :	
	3.00, At 0.25	μl :12	
	Variablevolun	e e	8
	• fullyautoclava	ıble,	
4.1.2	Micropipette • Single Chann	nel 5	
	• Capacity: 0.5	-10µl	
	• Increment:0.		
		5: At 10 μl : 1.00, At 5 μl :	=
	1.50, At 1µl :2		
	Variablevolur		
	fullyautoclava		
4.1.3	Micropipette • Single Chann		
	• Capacity: 2-2		
	• Increment:0.	•	6
. ·		5: At 20μl-0.90, At 10μl-	
8	1.20, At 2 µl-		9
	Variablevolur		
2	fullyautoclava	ıble,	

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4.1.4	Micropipette	• SingleChannel	5	
		• Capacity: 10-100μl		
	*	• Increment:0.1μl		
		• Inaccuracy: At 100 μl ±0.80, At 50μl		
		± 0.50 , At 10 μ l ± 0.30		
		Variablevolumes,		
*		fullyautoclavable,		
4.1.5	Micropipette	Single Channel	5	
		• Capacity: 20-200μl		
	8	• Increment: 0.1 µl		
		• Inaccuracy%: At 200 μl- 0.60, At 100μl-		
		0.80, At 20μl-3.00		. =
8		 Variablevolumes, 		
		• fullyautoclavable,		
4.1.6	Micropipette	Single Channel	5	
1.1.0	Micropipette	• Capacity: 100-1000μl		
		• Increment:5.0µl		
	39	• Inaccuracy%: At 1000 μl- 0.60, At 500μl-		
	9	0.70, At 100 μl-2.00		-
		Variablevolumes,		y
		• fullyautoclavable,		s 2
4.2	8 Channel Pipe			8 <u>.</u>
4.2		ellplates		
		ing head rotates foreffortless		* *
		gconvenience		
4		al piston and tip coneassemblies		
		easy repair andmaintenance		
		ind material-made tip cone secures high		
	sealing p	performance		
	 Compat 	ible with most universal tipbrands		
4.2.1	Micropipette	Multi Channel (8channel)	5	
		• Capacity: 0.5-10μl		
	4	• Increment:0.1μl		*
		• Inaccuracy%: At 10 μl- 1.50, At 5μl-		· · · · · · · · · · · · · · · · · · ·
		2.50, At 1 μl-4.00		
		Variablevolumes,		
		fullyautoclavable,		
4.2.2	Micropipette	Multi Channel (8channel)	5	*
		• Capacity: 5-50μl		2
		• Increment: 0.5μl		-
	,	• Inaccuracy%: At 50 μl- 1.00, At 25μl-		
	er er	1.50, At 5 µl-3.00		
	*	Variablevolumes,		
		• fullyautoclavable,		
5	Accessories, sn	ares and consumables		2
5.1	All standard acc	essories to be included in the offer. Bidders must		
5.1	specify			-
	the quantity of e	every item included in their offer (including items not		<u> </u>
	specified above).		\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \
6	Operating Env	ironment	-	A A A A A A A A A A A A A A A A A A A
				PX Gledona nec.

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6.1	The system offered must be designed to store and be operated	
	normally under the condition of the purchaser's Country. The	
e e	conditions include	8.
27 14	Power supply, Climate, temperature and relative humidity.	(148.117
7	Standards and Safety Requirements	
7.2	Must submit ISO and CE certificates	3
8	User Training	
8.1	Must provide user training (including how to use and maintain	
	the equipment).	
9	Warranty	
9.1	Comprehensive warranty for 1 years.	
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 bidder must arrange for the equipment to be installed by certified	
	or	
	qualified personnel; any prerequisites for installation to be	
er	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	Certificate of calibration and inspection from factory.	

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

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Real Time RT-PCR Kits for SARS Cov-2

Ñame of ítem	Purchaser's Specifications	Bidder's Offer/ Statement of Compliance	Deviation if any	Page no. of catalogue/ datasheet/ manual
Purpose	RT-PCR testing (COVID 19)			
Description	Must target at least two genes (E, RdRP, N, ORF 1ab.) should include positive control and internal controlfor both targets, should be compatible with ABI 7500, Biorad (CFX 96) and rotorgeneplatforms The kit should include RT PCR enzyme and Buffer Sensitivity atleast 95%			
Other requirements	The manufacturer should be certified by WHO for emergency use			

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RNA extraction reagent for manual extraction

Item name	RNA extraction reagent for manual extraction	Bidder's Offer/ Statement of Compliance	Deviation if any	Page no. of catalogue/ datasheet/ manual
Purpose	RNA extraction from swab sample			
Description	Spin column based suitable for manual extraction of body fluid, oro and nasophayringeal swab, blood samples serum or plasma samples. Sample input: Up to 400 microliter Elusion Volume: More than 6 microliter Purity: High quality RNA ready for Real Time PCR Should not require healting step. Extraction steps should not take more tan twenty minutes. Should contain reagents for RNA binding, nonenzymatic lysis (washing and elusión)			
Other requirements	The manufacturer should be certified by WHO or USFDA or CE or should have been listed by USFDA/CDC or WHO for emergency use			

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Virus Transport Medium (VTM)

Item name	Virus Transport Medium (VTM)	Bidder's Offer/ Statement of Compliance	Deviation if any	Page no. of catalogue/ datasheet/ manual
Purpose	Swab collection (COVID 19)			
Tube Requirement s	Suitably prepared sterile media for use in collecting throat and nasal swabs from human. Should contain virus inactivator, should stablized with RNA. Should be contained in airtight plastic tubes with cap. There must be sticker for labeling			
Medi a volu me	1ml or 3ml			*,
Swab collection sticks	Along with the tubes there should be two swab sticks (one for oropharangeal swab and another for nasophyrangealswab). There should be provision of break lines to allow to feed into thecontainer. Both the sticks should have fiber acrylic swab. The stick for nasophyrangeal swab should be flexible enough for ease in collection of swab simple from nasopharynx. The tube and stick should have been blistered.			
Others	The item should be CE and USFDA approved.	¥		

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Other Terms and Conditions of the Proposal:

- (i) The supplier should submit document stating the stock evidence of required quantity or the evidence with purchase order of required quantity in given time frame.
- (ii) The Supplier should submit ISO,CE or USFDA certificate as mention in Specification of required item.
- (iii) The supplier should quote the price of package item and Municipality evaluates packagewise & award the lowest amount of package. Supplier shall quotes in more packages.
- (iv) The supplier should supply the item according to municipality purchase order (partiall yor fully) as per municipality need. The quantity mention in Price Schedule will be increased or decreased as per municipality need.
- (v) Municipality has right to fully/partially accept or decline the proposal submitted by the supplier.
- (vi) Bids must be valid for a period of 90 days after the bid closing dedline. Bids may be accompanied by a bid security amount to a minimum of 2.5 percent of total amount or either in the form of bank voucher in the name of rastriya banijya bank limited gamgadhi mugu branch, dharauti account number 4060100303000002 or irrevocabale and unconditional bank gurentee issued by A class commercial bank with a minimum amount of 2.5 percent of total amount. The bid security shall be valid for a 30 days beyond the validity period of the bid.

विविवेजेंड्र मल्ल इ. वमस प्रशासकीय प्रमुख







गमगढी, मुगु

प.सं.: ०७७/०७८

च.नं.:

कर्णाली प्रदेश, नेपाल

Form of Agreement

WHEREAS the Purchaser invited Priced Quotation for certain goods and ancillary services, viz., [brief description of goods and services] and has accepted a Price Quotation by the Supplier for the supply of those goods and services in the sum of [contract price in words and figures] (hereinafter called "the Contract Price").

NOW THIS AGREEMENT WITNESSETH AS FOLLOWS:

- In this Agreement words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to.
- $\bullet \ \, \text{The following documents shall be deemed to form and be read and construed as part of this Agreement, viz.:} \\$
- Price Quotation Form and the Price Schedule submitted by the Supplier;
- The Schedule of Requirements;
- The Technical Specifications;
- The Conditions of Contractand
- The Purchaser's Notification of Award.
- In consideration of the payments to be made by the Purchaser to the Supplier as hereinafter mentioned, the Supplier hereby covenants with the Purchaser to provide thegoods and services and to remedy defects there in inconformity in all respects with the provisions of the Contract.
- The Purchaserhere by covenants to pay the Supplier inconsideration of the provision of the goods and services and the remedying of defects therein, the Contract Price or such other sum as may become payable under the provisions of the contract at the times and in the manner prescribed by the Contract.

IN WITNESS whereof the parties hereto have caused this Agreement to be executed in accordance with their respective laws the day and year first above written.

On behalf of the Purchaser

Name:

Designation

Sign:

Seal:

ललितजङ्ग मल्ल मख प्रशासकीय प्रमुख On behalf of the Supplier

Name:

Designation:

Sign:

Seal: