Bid Number: GEM/2020/B/712891
Dated: 16-07-2020

Bid Document

Bid Details	
Bid End Date/Time	27-07-2020 13:00:00
Bid Opening Date/Time	27-07-2020 13:30:00
Bid Life Cycle (From Publish Date)	90 (Days)
Bid Offer Validity (From End Date)	30 (Days)
Ministry/State Name	Jammu & Kashmir
Department Name	Home Department Jammu And Kashmir
Organisation Name	Jammu And Kashmir Police
Office Name	Police Hospital Srinagar Kashmir
Total Quantity	1000
Item Category	VTM KITS (SWINE FLU)
Experience Criteria	3 Year (s)
MSE Exemption for Years Of Experience	Yes
Startup Exemption for Years Of Experience	Yes
Document required from seller	Experience Criteria *In case any bidder is seeking exemption from Experience / Turnover Criteria, the supporting documents to prove his eligibility for exemption must be uploaded for evaluation by the buyer
Bid to RA enabled	No
Inspection Required	No

EMD Detail

Required	No

ePBG Detail

Required	No

Splitting

Bid splitting not applied.

1. Experience Criteria: In respect of the filter applied for experience criteria, the Bidder or its OEM {themselves or through reseller(s)} should have regularly, manufactured and supplied same or similar Category Products to any Central / State Govt Organization / PSU / Public Listed Company for number of years as indicated in the bid document before the bid opening date. Copies of relevant contracts to be submitted along with bid in support of having supplied some quantity during each of the year. In case of bunch bids, the category of primary product having highest value should meet this criterion.

VTM KITS (SWINE FLU) (1000 pieces)

Technical Specifications

* As per GeM Category Specification

Specification	Specification Name	Values	Bid Requirement (Allowed Values)
GENERAL FEATURES	Product Description	VTM Kit - Swine Flu	*
	Items included	Viral transport media with swabs	*
	Purpose	To collect and transport viruses, chlamydiae, mycoplasma and ureaplasma in an active form to the laboratory from collection site for isolation duly maintaining the viability and virulence of the viral sample	*
	Sterile and ready to use	Yes	*
	Method of sterilization for media	0.22 micron filtered sterilized	*
	Method of sterilization for swabs	ETO	*
PRODUCT INFORMATION	Medium should contain protective protein and antibiotic to control microbial and fungal contamination and buffers to control the pH	Yes	*
	Medium should contain cryoprotectant to help preserving the virus for long term frozen storage	Yes	*
	Allows long survival of the present virus and and offers maximum recovery	Yes	*
	Phenol red pH indicator in the medium to ensure medium integrity at the time of specimen collection	Yes	*
	Spontaneous elution of sample into liquid media	Yes	*
	Media filled in self standing tubes which eliminates biohazardous spill	Yes	*
	Swabs suitable for nasopharyngeal,nasal or traceal samples	Yes	*
	Patient Friendly Design of swabs	Yes	*
	pH value of medium at 25°C	7.3 ± 0.3	*
KIT CONTENTS	**Viral transport Medium**	3 ml viral transport medium filled in 15 ml standing polypropylene tube	3 ml viral transport medium filled in 15 ml standing polypropylene tube

	Closure type of tube	Normal cap	Normal cap
	Number of swabs provided with each tube	2	2
	Type of Swabs provided with each tube	Sterile flocked Nylon Swab with breakpoint,Sterile polyester swab with breakpoint	Sterile flocked Nylon Swab with breakpoint, Sterile polyester swab with breakpoint
	Literature with complete directions for collection, storage, transport and carrying provided with the Kit	Yes	*
PACKAGING	The packing and labelling of the kit should be as per Drugs and Cosmetics Act, 1940 and Rules, 1945	Yes	*
	Type of packing for medium	Filled in polypropylene tubes	*
	Type of packing for swabs	Individually packed in easy to open peel pouches	*
	Each tube with swabs packed in a duplex box	Yes	*
	Pack size (Qty/Pkg)	50	*
STORAGE CONDITIONS	Storage temperature for the kit	2º C to 30º C	*
	"The supplier should ensure maintenance of recommended temperature during storage and shipping of Kit "	Yes	*
CERTIFICATIONS & REPORTS	The kit should have approval of the statutory authority in its country of origin	Yes	*
	The Kit should be registered and licensed in India by DCGI (Proof of the same to be submitted to buyer on demand)	Yes	*
	Availability of valid drug license from competent authority defined under Drugs and Cosmetics Act, 1940 (Proof of the same to be submitted to buyer on demand)	For Manufacture to sale,For Sale or Distribution	*
	Drug License Number	-	*
	Drug License Date	-	*
	Manufacturer certifications (Proof of the same to be submitted to the buyer on demand)	GMP	*
	GMP/WHO GMP Certification Number	-	*
	GMP/WHO GMP Certification Date	-	*

	ISO Certified Manufacturer (Proof of the same to be submitted to buyer on demand)	Yes	*
	Product Certifications (Proof of the same to be submitted to buyer on demand)	EU-CE (IVD)	*
	Availability of Test report of Final Product from Central GOVT/NABL/ILAC accredited Lab to prove the conformity to the declared specification(proof of the same to be submitted to the buyer on demand)	Yes	*
SHELF LIFE	**Shelf life from the date of manufacture (months)**	12	12
	The product should not have passed more than 1/6 of the total shelf life at the time of dispatch to the consignee	Yes	*

* Specifications highlighted in bold are the Golden Parameters.
* Bidders may note that In respect of non-golden Parameters, the specifications 'Values' chosen by Buyer will generally be preferred over 'Bid requirement (allowed Values) by the Buyer.

Additional Specification Documents

Consignees/Reporting Officer and Quantity

S.No.	Consignee/Reporting Officer	Address	Quantity	Delivery Days
1	Bashir Ahmad Sofi	190009,BATMALOO BUS STAND	1000	15

Bid Specific Additional Terms and Conditions

1. After award of contract – Successful Bidder shall have to get advance sample approved from buyer before bulk manufacturing / starting bulk supplies. Successful Bidder shall submit 1 samples for Buyer's approval, within 5 days of award of contract. Buyer shall, as per contract specifications framework, either approve the advance sample or will provide complete list of modification required in the sample within 5 days of receipt of advance sample. Seller shall be required to ensure supply as per approved sample with modifications as communicated by Buyer. If there is delay from buyer side in approval of advance sample – the delivery period shall be refixed without LD for the period of delay in sample approval. In case, the sample is found to have major deviations / not conforming to the Contract specifications, the buyer at its discretion may call for fresh samples for approval before allowing bulk supplies or may terminate the contract after notifying the deviations to the seller.
Unless otherwise provided in the contract, all samples required for test shall be supplied by the contractor free of cost. Where under the contract, the contractor is required to submit an advance sample, any expenses incurred by the contractor on or in connection with the production of stores in bulk, before the sample has been approved unconditionally shall be borne by the Seller and he shall not claim any compensation in the event of such sample being found unacceptable by the Buyer / Consignee.

This Bid is also governed by the General Terms and Conditions

---Thank You---