

### Bid Document

Bid Details	
Bid End Date/Time	24-07-2020 17:00:00
Bid Opening Date/Time	24-07-2020 17:30:00
Bid Life Cycle (From Publish Date)	90 (Days)
Bid Offer Validity (From End Date)	75 (Days)
Ministry/State Name	Ministry Of Labour And Employment
Department Name	Employees State Insurance Corporation
Organisation Name	Employees State Insurance Corporation
Office Name	Model Hospital Basaidarapur
Total Quantity	6240
Item Category	HCV (ELISA) Test Kits
MSE Exemption for Years of Experience and Turnover	No
Startup Exemption for Years of Experience and Turnover	No
Bid to RA enabled	No
Inspection Required	No

### EMD Detail

Required	No
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### ePBG Detail

Required	No
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### Splitting

Bid splitting not applied.

### HCV (ELISA) Test Kits ( 6240 pieces )

### Technical Specifications

\* As per GeM Category Specification

Specification	Specification Name	Values	Bid Requirement (Allowed Values)
PRODUCT INFORMATION	<b>Type of Kit</b>	4 th generation	4 th generation
	Detects	Antibodies specific to Hepatitis C Virus	*
	<b>Detection Type</b>	Qualitative	Qualitative
	Microplate ELISA Coated with recombinant and/or synthetic peptide antigens for core NS3, NS4 and NS5	Yes	*
	<b>Test can be performed on</b>	Whole Blood, Serum, Plasma	Whole Blood, Serum, Plasma
	Assay Procedure Time (minutes)	90	*
	The Assay should have sensitivity of $\geq 99\%$ and specificity of $\geq 98\%$	Yes	*
	The assay component should include reactive and non reactive control with each kit	Yes	*
	Storage temperature	2°C to 8°C	*
	The supplier should ensure maintenance of cold chain during storage and transportation of Kits at 2°C to 8°C	Yes	*
	Cold Chain indicator provided with the kits	Yes	*
	A cumulative time/temperature indicator should indicate the exposure to temperature in the range of 2°C to 8°C	Yes	*
	The cumulative time temperature indicator technology used should be pre qualified by WHO	Yes	*
	Adequate documents in english detailing the principle, components, methodologies, validity	Yes	*

	criteria, performance characteristics, bio-safety, limitations of assay, storage condition, manufacturing and expiry dates and methods of disposal provided with each kit		
	The Kit Should be compatible to both semi automated and fully automated Elisa analyzers	Yes	*
	The volume of all the chemicals used should be adequate enough for automated Elisa analyze	Yes	*
	The volume should cover the dead volume for automated ELISA system	Yes	*
	The kit should comply with all provisions of Drugs and cosmetics Act, 1940 and applicable rules there under	Yes	*
GENERAL FEATURES	Product Description	HCV ELISA Test Kit	*
	Clinical Purpose	To provide diagnosis of Hepatitis C virus infection	*
PACKAGING	<b>Pack Size</b>	96 tests / kit	96 tests / kit
	The packing and labelling of the kit should be as per Drugs and Cosmetics Act, 1940 and applicable rules there under	Yes	*
CERTIFICATIONS & REPORTS	The kit should have approval of the statutory authority in its country of origin	Yes	*
	The Kit is registered and licensed in India by DCGI in case of imported kits (Proof of the same to be submitted to buyer on demand)	NA for domestically manufactured kits	*

Indigenous manufacturers should be licensed by the competent authority defined under Drugs and Cosmetics act 1940	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 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 13485 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)	*
Four digit number of notified body If product is EU-CE certified	-	*
Certificatiion Number	-	*
Certificatiion Date	-	*
Certification Issuing Authority	-	*
Availability of Test report 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	Yes	*

	on demand)		
	Availability of Performance Evaluation Report (Proof of the same to be submitted to the buyer on demand)	Yes	*
	Performance Evaluation Report issuing body	Other	*
	Name of the Performance Evaluation Report issuing body if other than specified institute	-	*
SHELF LIFE	<b>Shelf Life from the date of manufacture (in months)</b>	24	36, 24, 18, 30
	The product should have atleast 3/4 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	Vivek Kumar	110015,ESIC MODEL HOSPITAL BASAIDARAPUR NEW DELHI	6240	15

### Bid Specific Additional Terms and Conditions

1. Scope of supply (Bid price to include all cost components) : Only supply of Goods

[This Bid is also governed by the General Terms and Conditions](#)

---Thank You---