## File No.: GA-25/2/2025-Gr.Admin-FSSAI Food Safety and Standards Authority of India (A statutory Authority established under the Food Safety and Standards Act, 2006) (General Administration-Central Procurement Unit) FDA Bhawan, Kotla Road, New Delhi – 110002

#### Dated: 09.07.2025

### **Corrigendum**

Reference is invited to the GeM Bid No. GEM/2025/B/6066121 dated 15.05.2025 and the pre-bid meeting held on 29.05.2025.

2. In this regard, the Technical Specifications of the Liquid Chromatography-Mass Spectrometry Mass Spectrometry (LC-MS/MS), UPLC with PDA, Fluorescence and RI Detector have been revised and the same are being placed at Annexure-I & Annexure-II respectively.

3. Further, the Bid End Date is being extended upto 21.07.2025

4. All other Terms and Conditions shall remain the same.

(This issues with the approval of the Competent Authority)

Ashur 2025

(Avinash Kusumakar) Joint Director

### **ANNEXURE 1**

## <u>Technical Specifications of Liquid Chromatography–Mass Spectrometry Mass</u> <u>Spectrometry (LC-MS/MS)</u>

### **General Tender Specification**

The latest Triple/Tandem Quadruple LC-MS/MS Bench-Top System for high sensitivity trace level qualitative and quantitative analysis with complete software control for multi-residue analysis to meet food safety regulations with the following specifications UPLC with Auto-sampler, degasser, column oven, with a splitter to analyze injected sample by MS/MS. Instrument should meet the global food regulation requirements (like CODEX, USFDA, EU, FSSAI, etc.)

S. No.	Parameter		Description
1.	Pump	i.	Binary pump capable of switching between four/two solvents.
		ii.	The system should be capable of being operated both as a HPLC and Fast HPLC.
		iii.	Vacuum degassing capability
		iv.	Operating Flow Rate Range to be 0.01 to 2.00 ml/min,
			in 1µl increments with Gradient profiles.
		v.	Maximum Operating Pressure: 15,000 psi or better.
		vi.	Effective System Delay Volume $\leq 100\mu$ l, independent
			of system backpressure.
		vii.	Compatible to pH 2 to 12
		viii.	Composition Accuracy: +/- 0.5% or better
		ix.	Composition Precision: 0.15% RSD or +/- 0.04 min SD, whichever is greater, based on retention time.
		x.	Flow Precision: 0.075% RSD or +/- 0.02 min RSD, 6
			replicates, based on RT ( $0.500 - 2.000 \text{ mL/min}$ ),
		xi.	Flow Accuracy: +/- 1.0% (0.500-2.00 mL/min) or
			better.
		xii.	Back Pressure: 15,000 psi or more.
2.	Degassing	i.	Online degassing unit
	Unit:		
3.	Auto-	i.	The Auto sampler must accommodate not less than 100 vials
	Sampler/	ii. iii.	Injection precision should be <0.3% RSD or better Vial capacity – up to 1.5- 2ml
	injector	iv.	Needle washing facility – should have needle washing facility
		<b>1 v ·</b>	from internal and external side programmable
		v.	Built in dilution and derivative system facility
		vi.	Syringe size –Should accommodate the injection volume stated
			above.
		vii.	Injector Linearity > 0.999 coefficient of deviation
		viii.	Sample carryover - <0.005%
		ix.	It should have the facility of keeping the sample in a cooling condition (temperature range from 4 to 40°C.
4.	Column	i.	The temperature range should be $\frac{5 \text{ degrees above}}{2}$ ambient to $80 \degree \text{C}$
	Oven:		$/90^{\circ}$ C or better.
		ii.	It should be able to accommodate at least 2 Nos of 25 cm or better columns within the oven.
		iii.	Temperature Stability: ±0.1 °C of set temperature or better.

5.	Workstation	i.	Latest version of software which full fills the requirement of 21
5.		l.	Latest version of software which full fills the requirement of 21 CFR part 11, Food safety compliance. Compatible to LIMS
	software	ii.	The ion ratios fallout with the user defined values
		iii.	Software should have the library database of around 1000
		<u>111</u> .	compounds viz pesticides, antibiotic residues and mycotoxins
			etc.
		iv.	Perform alternating +ve /- ve scan in one run.
		v.	Automated quantification and reporting of acquired samples –
			Batch wise and individual.
		vi.	The software should provide the capability to save data in a user-
			specified folder or subfolder, allowing flexibility in choosing the
			desired storage location.
		vii.	LIMS compatibility software for batch analysis is desirable
		viii.	Any update of software during warranty and CMC period should
			be done free of cost.
MSM	S Detector		
6.	Auto-tuning	i.	Auto-tuning with sensitivity and resolution optimization for both
7.	Mass Range	i.	positive ion and negative ion modes. 10-2000 m/z or better
8.	Scan Speed	i.	15000 Da /sec or better
9.	Resolution	i.	0.7 Da or better at a unit mass resolution
	Mass	i.	
10.	Stability	1.	0.1 Dalton over 24 hours or better
11.	Sensitivity/	i.	MRM ESI +ve 1pg on column reserpine should give
	Detection		chromatographic S/N greater than >15,00,000:1 without
	Limit		smoothening MRM transition 609-195. (Proof of Statement
			must be provided).
		<mark>ii.</mark>	MRM ESI -ve 1pg on column chloramphenicol should give
			chromatographic S/N greater than >15,00,000:1 without
			smoothening MRM transition 321-152. (Proof of Statement
			must be provided)
		iii.	Documentary evidence to be submitted along with quotation.
			For ten injections, $\%$ RSD should be <5%. Chromatograms to
			be provided, with details of mobile phase, column, and injection
			volume. Statistical treatment used to determine S/N ratio is to
			be specified along with raw data.
		iv.	The same data is to be reproduced after the installation of the
		<b>IV.</b>	instrument at the site.
			nistiument at the site.
		v.	Instrument detection limit: Should be 0.5 fg or less (Proof of
			Instrument detection limit: Should be 0.5 fg or less (Proof of Statement must be provided)
12.	Polarity	v. i.	Instrument detection limit: Should be 0.5 fg or less (Proof of Statement must be provided) +ve / -ve polarity switching time between alternate MRM scans:
	Switching	i.	Instrument detection limit: Should be 0.5 fg or less (Proof of Statement must be provided) +ve / -ve polarity switching time between alternate MRM scans: <20 ms or better
12. 13.	Switching MRM		Instrument detection limit: Should be 0.5 fg or less (Proof of Statement must be provided) +ve / -ve polarity switching time between alternate MRM scans: <20 ms or better Must be able to measure minimum 450 MRM/sec in one
13.	Switching MRM channels	i. i.	Instrument detection limit: Should be 0.5 fg or less (Proof of Statement must be provided) +ve / -ve polarity switching time between alternate MRM scans: <20 ms or better Must be able to measure minimum 450 MRM/sec in one acquisition to enable Transition Studies within a single run.
	Switching MRM	i.	Instrument detection limit: Should be 0.5 fg or less (Proof of Statement must be provided) +ve / -ve polarity switching time between alternate MRM scans: <20 ms or better Must be able to measure minimum 450 MRM/sec in one

15.	Detector	i.	PMT/EMT/CEM/DDD detector having the highest sensitivity system should bequoted.
16.	Nitrogen	i.	Suitable gas generator where ever available shall be provided
10.	Generator		and cylinders with all accessories such as SS double stage gas
			regulator, gas purification panel with brackets etc. Gas
			cylinders, minimum two nos.
		ii.	Gas Generators capable of supplying all necessary gases with
		11.	the required purity, pressure and flow rate, as required for the
			LC-MS/MS instrument should be provided. It should be
			complete with all necessary accessories.
		iii.	Highly reputed brand of Nitrogen generator with inbuilt
		111.	compressor with low noise should provided. Should be covered
			under two years comprehensive warranty with at least two
			Preventive maintenances along with PM kit each year.
			Satisfactory performance certificate should be given every six
17.	Source	i.	month of preventive maintenance visit.The cleaning of the source should be possible without venting the
1/.	cleaning	1.	system.
18.	Infusion	i.	The infusion device must be an integral component of the instrument
101	Device		and should be operable through the instrument's software, enabling
			the infusion of tuning and calibration solutions into the probe via a
			selection valve.
19.	Ionization	<mark>i.</mark>	Compatible to both ESI and APCI source, with a facility of
	source		interchanging easily by user and auto detection of installed probe by the instrument and software.
		ii.	Ionization ESI and APCI sources to be provided as standard,
		<b>11</b> .	with facility of interchanging easily by the user, and auto-
			detection of installed source by the instrument and software.
			The ionization must be done both in a positive & negative
			mode.
		<mark>iii.</mark>	The source should be easily removable from the system to
			facilitate user cleaning without venting the vacuum, with
			automatic shutdown of system while the source / probe is being removed.
		iv.	The source shall have a flow rate compatibility from 200
			$\mu$ L/min to 2000 $\mu$ L/min, without flow splitting in both ESI and
			APCI modes.
		v.	Temperature for ESI/APCI sources should be 450° Celsius or
			better.
		vi.	Desolvation temperature for ESI and APCI sources should be
		vii.	450° Celsius or better. All source parameters to be adjustable through software.
20.	Operating	<u>vii.</u> i.	Full scan
20.	modes	ii.	SIM scan
		iii.	Product ion scan
		iv.	Precursor ion scan
		V.	Neutral loss/gain scan
		vi.	Multiple Reaction Monitoring (MRM)
21.	MRM	i.	The latest version of the Pesticide and antibiotics MRM
<b>41</b> •	Library	1.	library with more than 850 pesticides having analyst should be
			norary with more than 650 pesticides having analyst should be

		provided along with the system.
22.	LC Columns	i. Columns for pesticides analysis (Multi residue) – 4 nos, along with
		suitable guard column.
		<ul> <li>ii. Column for polar pesticides analysis without derivatization – 2 nos, along with suitable guard column.</li> </ul>
		iii. Column for vitamin analysis (water soluable) – 2 nos, along with
		suitable guard column.
		iv. Column for vitamin analysis (fat soluable) – 2 nos, along with
		suitable guard column.
		v. Column for Steriods and stilbens – 2 nos, along with suitable guard column.
		vi. Column for antibiotics – 2 nos, along with suitable guard column.
		vii. Guard column holder $-4$ set.
23.	Consumable	
	s/ Spares	be provided (total 1000nos each).
	_	ii. PM kit for HPLC and MS sufficient for trouble free operation during
		warranty and CMC period should be provided along with instrument.
		The PM kit to include all consumables of HPLC and MS including O
		rings, probe capillaries, inlet steel tubing, Check valves and seals, rotor
		seals, spray needles, syringe or any other minor consumables other than
		column and peak tubing's. iii. 2 set of 10, 20 and 1 set of 50 and 100µl loop additional shall be provided
		for injection.
		iv. The vendor should provide a PFAS kit so the users can perform
		PFAS applications on the same system in food Matrix (Food of
		Animal Origin) – Fish, Milk, Honey and Water.
		The vendor should also be providing the application method details for PFAS application.
		System should be supplied with suitable accessories required for Sample Preparation/ analysis and
		QUECHERS Kit (as applicable) for PFAS analysis in food Matrix (Food of Animal Origin) – Fish, Milk, Honey and Water. 1000 nos.
		v. System should be supplied with suitable Sample Preparation accessories.
		nos each) for Pesticides etc in following matrices:
		<ul> <li>a) HLB cartridges or equivalent</li> <li>b) High fat containing food.</li> </ul>
		c) High Water content food.
		d) Highly Pigmented foods (eg chlorophyll, lycopene, carotene
		etc).
		e) Vitamin (SPE C18 900mg cartridge).

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24.	UPS System and Pre	i.	10 KVA UPS with a minimum of 60-minute backup time for
	and Pre requisite:	ii.	supporting the instrument. Reputed Branded of online UPS system of 10 KVA capacity with
	requisite.		comprehensive warranty of minimum 5 years inclusive of SMF
			batteries should be provided.
25.	Computer &	i.	Suitable branded computer, i7 processor with 12 GB DDR3
	Printer		Memory, Up to 1 TB SATA hard drive (7200 RPM) with original
			MS office professional lifetime.
		ii.	DVD-RW 24" LCD/LED Monitor with suitable authorized
			operating system
		iii.	Hp LaserJet colour automatic back-to-back printer with scanner.
		iv.	Any update of software during warranty and CMC period should
		1.	be done free of cost.
26.	Software	i.	Latest version of software which full fills the requirement of 21
20.	Software	<b>1.</b>	CFR part 11, Food safety compliance. Compatible to LIMS
		ii.	The ion ratios fallout with the user defined values
		iii.	
		<u>111.</u>	Software should have the library database of around 1000
			compounds viz pesticides, antibiotic residues and mycotoxins etc.
		iv.	Perform alternating +ve /- ve scan in one run.
		<mark>v.</mark>	Automated quantification and reporting of acquired samples –
			Batchwise and individual.
		vi.	The software should provide the capability to save data in a user-
			specified folder or subfolder, allowing flexibility in choosing the
			desired storage location.
		vii.	LIMS compatibility software for batch analysis is desirable
		viii.	Any update of software during warranty and CMC period should
			be done free of cost.
27.	IQ/OQ/PQ	i.	The instrument must be "Qualified" along with the Software.
			Necessary reagents along with Documents must be provided.
			During installation and qualification, Instrument should perform as
			per submitted specification in presence of user.
		ii.	IQ/OQ/PQ to be performed as per OEM Standard protocol should
			be done free of cost with necessary traceable standards (traceable
			to ISO 17034).
		iii.	PM visit along with kit must be supplied every year with the system
			till the warranty period.
		iv.	Documents, Kits & standards etc. as required being supply along
			with the instrument
		v.	To be done free of cost with traceable calibration standards for the
			first 5 years' warranty period (at installation & at every
			maintenance visit of each year) along with PM kits.
		vi.	OQ/IPV should be done free of cost with supply of PM Kits and
			calibration standards at least once in a year during warranty period.
		vii.	Documents, PM Kits & calibration standards etc. to be supply along
			with instrument at every PM visit free of cost during warranty
			period.
		viii.	Satisfactory performance certificate should submit to the laboratory
		, 111.	Page 5

[		after calibration of LC-MSMS before one month of warranty period
		expired with supply and fixation of PM kits of HPLC and MS
		system.
28.	Warranty	i. Minimum 5 years from the date of completion of IQ, OQ and PQ
		of LC-MS/MS to the satisfaction of NFL including Nitrogen
		Generator, PC and all accessories. ii. The date of warranty period for LC-MSMS including Nitrogen
		ii. The date of warranty period for LC-MSMS including Nitrogen Generator, PC, Printer, Gas cylinders with its accessories and all
		associated supply of LC-MSMS, which will start from the date
		completion of IQ, OQ and PQ of LC-MSMS.
		iii. It should cover hardware, software as well as wear and tear
		consumables (except column and sample preparation), Up-
		gradation of software to the latest version (if applicable-), prompt
		service (within 48 hours on-call), training and application support
		during the period without any extra charge. iv. In case of breakdown of the system, the servicing to be done
		immediately by the supplier during the warranty period and
		maximum down time period is 48 hrs, if it's not attended the
		warranty will extend accordingly.
		v. Warranty should be covered for all accessories and 3rd party items
		provided with the system. For delay in attending break-down call
		beyond 2 working days a penalty @ $\gtrless 10,000/$ - per day shall be
		charged. Such amount will be deducted from any amount due or which may become due to the supplier. The warranty period shall
		automatically stand extended by the number of days taken to rectify
		the defects.
29.	After sales	i. Should have a good after sales service/technical support capable of
	service/Post	reaching at short notice the places where LC-MS/MS is proposed
	Warranty	to be installed. Visits and unlimited breakdown calls by
		service/application support, engineers should attend immediately
		without fail for LC-MS/MS including Nitrogen Generator and UPS
		system. ii. Troubleshooting training (Instrumentation/Application) as and
		when required shall be provided free of cost.
		iii. The application and method development support must be rendered
		for minimum 30 days during the warranty period.
		iv. The vendor should also assure supply of spares, accessories,
		consumables and service for at least 10 years including Nitrogen
		generator. v. Terms and conditions for the AMC & CMC, after the warranty
		period has to be specified.
		vi. Quote for AMC & CMC for 6th, 7th, 8th, 9th & 10th years, to be
		submitted separately.
		vii. The CMC shall include parts cover all hardware including detector,
		software as well as wear and tear consumables (except column and
		sample preparation), PM kit (yearly), annual calibration along with
		documentation.
		viii. AMC/CMC price quoted by the vendor will be considered as
		independent price. It will not be considered for finalizing the L1.
30.	Training	i. Basic training for a period of not less than two weeks after
		installation of the equipment to technical personnel and further
		for method development whenever required during warranty and
	1	

		-	CMC period should be provided, free of cost.
		ii.	Application support for method development for antibiotics,
			pesticides, mycotoxins analysis should be provided during
			warranty and as well as during CMC period, whenever required
			shall be imparted free of cost.
		iii.	Trouble shooting training as and when required shall be imparted
			free of cost.
		iv.	The instrument supplier has to perform on site method
			development and validation as per the laboratory /FSSAI
			regulatory requirements/ DG SANTE/ protocols for all pesticide,
			aflatoxin, vitamin and antibiotic in least four matrices as selected/
			preferred by the lab.
31.	Experience	i.	The supplier should have executed at least Minimum 20 or more
	-		of the model/series of model quoted successful installations
			among which 5 should be installed in Government institutes.
		ii.	The Complete users list for the quoted model in India, with
			contact addresses, emails and phone numbers should be provided
32.	General conditionsof	i.	The instrument and all its sub units should operate on 240 volts
	supply	ii.	50 Hz powersupply. All the operation and maintenance manuals, circuit diagrams,
	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	11.	application notes and application software's to be supplied should
			be in English language.
33.	Servicing of	i.	The servicing of LC-MS/MS, Nitrogen generator, Vacuum pump
	LC-MS/MS,		& UPS System shall be carried out only by the OEM authorized
	Nitrogen		service center/ engineer only, during the warranty period/ CMC
	generator,		period.
	Vacuum		
	pump & UPS System		
<mark>34.</mark>	System System	i.	Repeatability:
	performance		
	requirements		should be able to demonstrate RSD $< 5\%$ for 10µl injection of
			.5µg/kg standard solution containing group of pesticides without
			nternal standard should be provided by supplier along with tender
			ocuments in Matrix- Tea and spices for Pyridalyl, Spinosad, Dinotefuran.
			inoterurur.
		(	Calibration data, Chromatograms, Experimental data, Injection
		v	olume, COA of standard/ column used to be provided).

# **Technical Specifications of UPLC with PDA, Fluorescence and RI Detector**

# **General Tender Specification**

Quaternary solvent system with degasser, Auto sampler, Column oven and detectors. Total system (including pump, detector and auto sampler) should be capable of operation ranging minimum 15000 psi or better. Instrument should meet the global food regulation requirements (like CODEX, USFDA, EU, FSSAI, etc.)

Sl. No.	Main component		Desired Specifications
1	Pump	i.	High-Performance Binary gradient pump, capable of switching between four/ two solvents.
		ii.	Flow range: 0.01 to 2.0 mL or better
		iii.	<b>Flow rate accuracy:</b> $\pm 1.0$ % or better
		iv.	Flow rate precision: ≤0.1 % RSD or better
		V.	<b>Compositional accuracy:</b> ±0.5 % or better
		vi.	Back Pressure: 15,000 psi or more.
2	Degassing Unit	i.	Online degassing unit should be part of main system.
3	Auto-Sampler/ injector	i.	The Autosampler design should have variable injection volume capability.
		ii.	Temperature setting ranges from 4 °C to 35 °C or better.
		iii.	The Carryover must be below 0.005 % or better.
		iv.	Suitable sample rack for handling a minimum of 90 Nos. of 1.5-
			2 ml vial.
		v.	Injection-volume Precision <mark>: &lt;0.3% RSD</mark> or better
4	Column Oven	i.	The temperature operating range should be ambient to 80 °C or
			better.
		ii.	It should be able to accommodate at least 2 Nos of 150 mm
			columns within the oven.
		iii.	Temperature Stability: $\pm 0.1$ °C of set temperature or better.
5	Diode Array	i.	Wavelength range should be <mark>190 – 700 nm</mark> or more.
	Detector or Detection	ii.	Detection type should be with 1024 element photodiode array.
	Photodiode Array Detector	iii.	Light source - Deuterium and/ or tungsten lamps/ xenon.
	(DAD/PDA)	iv.	Data rate should be up to 80 Hz (points/sec) or better
	()	v.	Wavelength accuracy = $\pm 1$ nm.
		vi.	Flow cell: <1 μL or better volume, 10 mm cell path length
		vii.	Noise: < 5 x 10-6 AU
		viii.	Drift: < 0.5 x 10-3 AU/Hr
		ix.	Peak Purity analysis
6	Fluorescence	i.	Light source: Helium/ Xenon lamp Excitation Wavelength
	detector (FLD) with Post-		range: 200 nm to 850 nm or better
	Column	ii.	Emission Wavelength Range: 210nm to 850 nm or better
	Derivatisation unit	iii.	Spectral bandwidth: 20 nm

		iv.	Wavelength accuracy: $\pm 3$ nm or better
		v.	Wavelength repeatability: $\pm 0.2$ nm or better
		vi.	Signal to Noise ratio for Water Raman peak should be > 500
		vii.	Flow cell volume: < 2 uL
		viii.	Data rate: 70Hz or better
		ix.	Post Column Derivatisation (Photo chemical or electro
			chemical) unit shall be supplied with FLD
7	Refractive Index	i.	RI Range: 1.00 to 1.75 RIU
	<b>Detectors (RID)</b>	ii.	Measurement range: $\pm 6 \times 10-4 \text{ RIU}$
		iii.	Noise level should be $\pm 2 \ge 10-9 \text{ RIU}$
		iv.	Drift: 2 x 10-7 RIU/h
		v.	Compatible with flow rates up to 5 mL/min
		vi.	Flow cell volume: < 2 uL
		vii.	Safe leak detection
		viii.	Temperature control 30°C /ambient to 55 °C or better
		ix.	Data collection rate: 60Hz or better
8	Computer platform and Software	i. ii. iii. iv. v. v. vi. vii. vii. ix. x. x. xi.	Computer platform Software and operating system Suitable branded computer, i7 or equivalent Advanced processor with 16 GB DDR3Memory, Upto 1 TB SATA hard drive (with 5 TB External Drive) (7200 RPM) or better for software requirements of HPLC DVD-RW 32" LED Monitor with suitable authorized operating system, 4 USB Port or higher configuration. Licensed MS Office and PDF editor to be included along with the system. Software must be Multitasking type. It must acquire and process the data simultaneously. Workstation must be able to control the UHPLC with FLD RID and PDA/DAD & acquire, store, process and reproduce the data by the same computer. Software must have automated calibration and Quantitative optimization. Automated Quantitation and reporting of acquired samples. Data may be processed as it is being acquired. Chromatography software with integrated Oracle/SQL/ other database for the easy retrieval of data in case of disaster. Suitable multi-functional LaserJet B/W Printer should be supplied. Software should comply with 21 CFR Part 11 regulations and Support GLP operations.

9	Branded UPS System	i.	5 KVA online UPS with batteries with 60 mins backup suitable for complete system should be offered.
10			
10	Instrument Performance	i.	Automated system generated OQ and PQ report as per the end
	Verification/IPV		user requirement should be submitted.
	( <b>IQ,OQ, PQ</b> )	ii.	To be done free of cost with ISO 17034 traceable calibration
			standards for the first 2 years (at installation & at every
			maintenance visit of each year) with PM kits for UHPLC
			including detectors with lamp, only consumables will be cover in
			CMC, UPS with its accessories and all associated supply of
			UHPLC.
		iii.	OQ/IPV with report should be done free of cost with supply of
			PM Kits and calibration standards during warranty period.
		iv.	Satisfactory performance certificate should submit to the
			laboratory after calibration of UHPLC including detectors with
			lamp, Post column derivatiser unit, UPS with its accessories and
			all associated supply of UHPLC before one month of warranty
			period expired with supply and fixation of PM kits of UHPLC
			system. UPS with its accessories and all associated supply of
			HPLC before one month of warranty period expired with supply
			and fixation of PM kits of HPLC system.
11	Warranty	i.	Minimum 5 years from the date of completion of IQ, OQ and PQ
			of UHPLC to the user satisfaction including PDA, FLD Detectors, POST COLUMN DERIVATISER (Photo Chemical or
			electrochemical) for mycotoxin analysis and UPS all associated
			accessories of UHPLC.
		ii.	The date of warranty period for UHPLC including detectors with
			lamp, POST COLUMN DERIVATISER unit, UPS and all associated supply of UHPLC will start from the date completion
			of IQ, OQ and PQ of UHPLC.
		iii.	It should cover hardware, software as well as wear and tear
			consumables (except column and sample preparation), prompt
			service (within 48 hours on-call), training and application support
		iv.	during the period. In case of breakdown of the system, the servicing to be done
			immediately by the supplier during the warranty period and
			maximum down time period is 48 hrs, if it's not attended the
			warranty will extends accordingly.
		v.	For delay in attending break-down call beyond 2 working days a penalty @ ₹5,000/- per day shall be charged. Such amount will
			be deducted from any amount due or which may become due to

		the supplier. The warranty period shall automatically stand extended by the number of days taken to rectify the defects (beyond 2days).
12	After sales service/Post Warranty	<ul> <li>i. Should have a good after sales service/technical support capable of reaching at short notice the places where UHPLC is proposed to be installed. Visits and unlimited breakdown calls by service/application support, engineers should attend immediately without fail for UHPLC including PDA, FLD with POST COLUMN DERIVATISER (Photo chemical or electro chemical) unit and UPS system.</li> </ul>
		<ul> <li>ii. Troubleshooting training (Instrumentation/Application) as and when required shall be provided free of cost.</li> <li>iii. The application and method development support must be rendered for minimum 30 days during the warranty period.</li> <li>iv. The vendor should also assure supply of spares, accessories, consumables and service for at least 10 years.</li> </ul>
		<ul> <li>v. Terms and conditions for the AMC &amp; CMC, after the warranty period has to be specified.</li> <li>vi. Quote for AMC &amp; CMC for 6th, 7th, 8th, 9th &amp; 10th years, to be submitted separately.</li> <li>vii. The CMC shall include parts cover all hardware including detector, software as well as wear and tear consumables (except</li> </ul>
		<ul> <li>column and sample preparation), PM kit (yearly), annual calibration along with documentation.</li> <li>viii. AMC/CMC price quoted by the vendor will be considered as independent price. It will not be considered for finalizing the L1.</li> </ul>
13	Spares and accessories	<ul> <li>i. The following consumables, but not limited to should be supplied along with instrument:</li> <li>a) C18 (2.1×100MM) - 2 Nos</li> </ul>
		<ul> <li>b) Sugar analysis columns- 2 Nos</li> <li>c) Certified Vials (1.5 mL) with cap and septa 1000 Nos each.</li> <li>d) Poly Spring inserts or glass insert vials-150uL 1000 Nos each.</li> <li>e) High Recovery Vials (2 mL) with septa cap 1000 nos each.</li> <li>f) Amber Vials (2 mL) with septa cap 1000 nos each.</li> <li>g) PM kit for HPLC for trouble free operation during warranty period should be provided free of cost every year along with instrument.</li> <li>h) Injection Loop of 100uL for the auto sampler should be offered. Qty-1</li> <li>i) UHPLC Peak tubing (5 meter), (UHPLC round tight ferrules- 40 Pcs).</li> <li>i. The supplier should have executed at least Minimum 20 or more of</li> </ul>

15	Delivery	i.	The instrument supplied to the site of address provided. If any permit such as road permit/way bill, customs/excise duty, octroi or any taxes should be borne by the supplier. If any documents required for the above purpose, the office may consider to provide on request prior intimation
16	Training Component	i.	The supplier will have to carry out the successful installation at our laboratory premises (wherever the system will be installed) and provide on–site comprehensive training for scientific personnel operating the system and support services for 5/7 days every year with the system and training at the supplier's lab premises is also required.