

Automatic Bio Chemistry Analyser

Creator : Ms Ritika (TA)

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Measuring and observing and testing instruments : UnSPSC

Sub Category : Clinical and diagnostic analyzer reagents - UNSPSC Code

Automatic Biochemistry Analyser : UNSPSC Code -

S.No.	Parameters	Grouping		Value 1	Value2	Value 3	Value 4	Value 5	Golden	Validation	Whether Filter Required	Remarks	p r i t o y	U n i t
1	Confirmity to Standards	S	E	USFDA	CE	BIS	UL		G					
2	Certification No and Date	S	T											
3	Type of system			Fully Open Random Access system										
4	Purpose	P	E	should be capable of all routine,STAT and special biochemical tests including specific proteins,therapeutic drugs(TDM),Drugs of abuse ,Immunoturbidetric assays and user definable applications in blood,serum or urine and equipment to be configurable as per requirements of consignee										
5	Model type of Biochemistry Analyser	P		Bench top Type	Floor model Type				G					
6	Type of samples	P	E	serum,plasma , urine, CSF and any body fluid samples										
7	Type of Assay	P		End point,Kinetic turbidometric and bichromatic assay										
8	Throughput tests / hour	P	N	200 or more		350 or more		600 or more		G				
9	Calibration facility	P	E	factor	Linear (one, two and multipoint)	Exponential	Logit-log	spline	G					multi select

25	power consumption of lamp in VA	P	N							<1000 VA				
26	Type of sample cups	P	E	Reusable										
27	Number of sample cups provided with machine	P	E	500	1,000	1,200			G					
28	QC programe	P	E	QC Programme with L-J graphs, Print out of reports										
29	The reaction cuvette must be reusable and durable	P	E	Yes										
30	System should have on board cooling system	P	E	Refrigerated										
31	Good quality DI water plant supplied	P	E	Yes	No				G					
32	compatible on line UPS	P	E	YES	No									
33	back up time	P	E	1 hour	2 HOUR									
34	data management software	P	E	equipment to be supplied with compatible programmable windows based comprehensive data processing and management system graphica user interface software .LM Capability complete back up of data basefor calibration control and patient sample results										
35	patient result storage capacity	P	E	10000	15000								G	
36	multitasking facility on computer	P	E	yes										
37	System should have facility for reading results on monitor and print out facility	P	E	Yes										
38	Reagents to be supplied with machine	P	E	urea, uric acid, creatinin, sugar, Cholesterol, TP, albumin, SGOT, SGPT, alkaline phosphatase, Bilirubin (Total & Direct) calibrator 3mlx 4 and quality control 5mlx1 -5 Nos each for normal and abnormal										

63	Availability of test report from central govt/ NABL/ILAC accredited lab covering all parameters	R	Ch	Yes	No								
64	Test report number if test report not available put NA	R	Ch										
65	Test report date if test report not available put NA	R	Ch										
66	Name of test lab if test report not available put NA	R	Ch										
67	Address of test lab if test report not available put NA	R	Ch										
68	Copies of all certifications and reports to be provided to buyer on demand at time of supplies	R	Ch	Yes									

Shahzad
Astt. Director-II(Q/A)

Shri A. V. Muralidharan

DCEO -GeM

tbd : to be declared by vendor

Shahzad
Astt. Director-II(Q/A)

Shri A. V. Muralidharan
DIRECTOR -QA

