The DIMENSION EXL 200 s/n DE 2726 58
has been successfully installed as of 18th Feb 2020.
(Date)
ο / Δ l ·
Customer Accepted: Impulse Dragnoshz
Date: 1 1 2020
Title: 10 00 P8

[] EM Checklist attached (Obtain FSR signature)				
[] Calibration and QC Summary Sheets attached				
[] Inventory Summary Sheet attached				
[] Method Calibration / Verification / IMT Reports attached				
[] Rating Form completed and attached				
[] Training Checklists completed and attached				
[] Installation Completion Statement				
[] Software Revision Level				
Signatures:	Felo	18	2010	
Clinical Applications Specialist			date	
13	Feb	18	2020	

Field Service Representative

NOTICE TO INSTALLERS

This package contains new revisions

PLEASE READ PRIOR TO INSTALLATION

date



Instrument Installation Acceptance Statement

Account	Name:	Impulse Diagnostic				
Clty:	Silchar		State:	Assam		
Instrume	nt Installed	i:13/02/2020		S/N:	DE27:	2658
				S/N:		
				S/N:		
				S/N:		
Install Co	ompletion i	Date: 13/02/2020				
Custome	er:	ø				
complete maintena I also sta specifica	d to my sat nc∈ of thIs te that the I tIon and (2)	Ite that the Installation of to isfaction, including training instrument(s). Instrument(s) is/are (1) operated the instrument(s) and conto Siemens protocome.	g of laborat erating in a nsumable(s	ccordance w	on the	facturing
Custome	er's Name:	Impulse Dia	gnostic	8		
Custome	er Signatur	e: Print	Name	=	Date:	18/2/20
Siemens	CAS					
CAS's N	ame:	Da I monmi Iakai Print Na	me		- D-4	19/2/2
CAS SIg	nature:				Date:	11/2/20



Installation Procedure and Checklist – Dimension EXL200.

Models @ Serial No	: Dimension EXL200				
Instrument Sr#	: DE272658				
Customer Name	: Impulse Diagnostic				
Doctor / In-Charge	: Dr. Dipayan Saha				
Address	: Hailakandi, Meherpur				
City	: Silchar				
Phone	: 3482224542				
Inspect Shipper for	or physical damage, then Uncrate.				
a. Physical Da	mage to shipper	:	☐ Yes	I	No
	as per packing List	:	☐ Yes		No
2. Environmental Co	nditions.				
a. Air Condition	ned Environment Available	:	□ Yes		No
b. Room free o	f Rodents/ Insects/ Pests	:	⊡ -∕Yes		No
c. Humidity is l	ess than 80 %	:	G✓Yes		No



3.	Power Requirements.			
	a. Uninterrupted Power Supply Available.			
	 UPS RatingKVA UPS Make Type of batteries Back Up Time Hours Incoming AC Power: V AC. Earth Voltage V. 			
4.	General Cleanliness Around Site : Acceptable Not Acceptable			
5.	Verified all Instruments Components and Accessories :			
6.	Installed all Accessories as per the Instrument Set - Up Procedure Below :			
	Removed all shipping screws from the System			
	2 Assembled the User Interface Modules – PC, Keyboard			
	3 Installed the Printer & Bar Code Scanner			
	4 Installed the Top Cover			
	 Installed the Syringe Assembly Checked all cable connections; 			
	7 Checked all Electronic Boards are properly plugged in			
	8 Checked the Dispense Test for Probes			
	9 Installed the Current Software Version			
	10 Installed the Dade Behring water diluent bottle11 Installed the cuvette cartridge			
7.	Printer.			
	a. Make:			
	b. Installed: TOK			



8. System Calibration and Inspection.

		Verified All Mechanical Configuration Adjustment of the Site and Back up in Disc (Instrument Calibration Diskette provided to the User Department)	□ ⁄
	b.	Checked the Thermal Operations of Reagent compartment and Therma Chamber.	102
	f. g.	Checked the Reagent area and Automatic Flex Loader Checked Sample area. Checked the Barcode reader. Checked the Lamp Calibration Checked the Photometer alignment and mAU Offset calibration. Calibrated the Cuvette and reagent temperature	<u> </u>
9.	Ru perfo	n Preparation: System Check In the System check with CHK Reagent to check the Irmance of Sample Probe assembly, Reagent Probe 1 and 2 Inbly, Photometer alignment.	□ ✓

Attached System check Printout attached



Alignment: print out attached

Voltages:

	Power OF	F	Power ON	Power ON
G-H		V (0-1V)	(90-110V)	(198-264V)
		V (0-1V)	(110-125V)	
H-N		V (0-1V)	(90-110V)	(198-264V)
		V (0-1V)	(110-125V)	
G-N		V (0-1V)	(Max 0.5)	(< 2V)
		V (0-1V)	(< 2 V)	
Gaps: Syringe				
(Glass to	plunger)	Observed	Adjusted	Specification
Sample	Metering			(.005"010")
Sample				(.005"010")
•	1 Metering		_	(.005"010")
				(.005"010")
	1 Flush			(.005"010")
Reagent	1 Metering			
Reagent	1 Flush			(.005"010")
Reagent	1 Flush			(.005"010")
	nstrument ry Wash			(.005"010")
Cuvette	Manufacture	Solenoids		Specification
Top Seal				(.010" *)
Cuvette Form				(0.020" - 0.045"**)
U-Sea 1				(0.020" ± 0.010"*)



Installation Qualifications for Dimension EXL200.

Carried out all the Installation Steps as well as the Necessary Checks and Alignments of all Robotics were done for Dimension EXL200 located in Impulse Diagnostic, Silchar bearing serial No. DE272658 as per Installation Procedure and Checklist.

Checked the System check report and the obtained values acceptable. (Printout attached)

Performed all due maintenance activities such as Daily, Weekly Maintenance, Automated System Prime.

Handed over the Instrument for Operations Training & Qualifications.

For Siemens Ltd.

Name

: Mr. Kamal Kumar Baishya

Signature

Designation: Field Service Engineer

Date

: Carestya.

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Installation Certificate

This is to certify that the Dimension EXL200 Clinical Chemistry System, Instrument Serial DE272658 has been successfully Installed and Commissioned in Impulse Diagnostic, Meherpur, Silchar, Assam as per the Installation Procedure & Checklist.

Siemens Ltd.

Impulse Diagnostic

Name: Kamal Kumar Baishya

Name: Jantu Das

Designation: Field Service Engineer

Designation: Technician In-charge

Signature: Checistyn

Date: 18 | 02 | 20

Signature: Jantu \mathbb{R}^{as} ,
Date: 18 | 02 | 20



Operational Qualification for Dimension EXL200

Operator Qualification: Conducted the operator Training on the following Topic	
1, Component Overview	
a, System Components	
b, Keyboard, Touchscreen and Alert Keys	
2, Calibration:	
a, Calibrated Linear Method and verify Enzyme Method	
3, Maintenance:	\Box
a, Daily, Weekly, Monthly Maintenance and Periodic Maintenance	
b, Replace Cuvette Nozzle Diaphragm	
c, Replace Cuvette film cartridge	
d, Replace Reagent and Sample Probe tip.	
4, Sample Processing	
a, Running sample using Sample cup, primary tube.	
b, Manual dilution and respond to system needs.	
c, Determine Segment status and delete Segment.	
d, Review use of System status key	
e, Edit samples including adding and deleting tests, rerunning test and deleting S	Samples
f, Review use of these keys: Pause, Exit, Shift, Reset, Backspace, Backslash, Run and Arrow keys.	
g, Review Interpreting test report messages.	
5, Customization	
a, Set Password	
b, Enable automatic cartridge removal, and automatic repeat for panic	
c, Enable Automatic Flex reagent cartridge testing.	



- d, Select Plumbing configuration
- e, Define panel
- f, Define QC Status and QC ranges
- g, Review method QC results from method review screen.
- h, Enter Panic values
- i, Configure barcode choice
- j, Touchscreen and alert features
- k, Configure QC Alerts, QC ranges and QC Panels.
- I, Define calibration products and calibration alert.
- m, Setting calibration.

6, Problem Resolution

ACTIVITY	Reference
Review response to alarm ON/OFF	Operator's Guide, Introducing,
Review response to error messages using ALT M	Operator's Guide, Introducing
Review using Reset key to clear error messages	Operator's Guide, Introducing
Review active and resident error logs; including More Info and See Minor functions	Operator's Guide, Troubleshooting
Review troubleshooting, emphasizing system check troubleshooting guidelines	Operator's Guide, Troubleshooting
Review icons and using CTL Help to respond to icons	Operator's Guide, Appendix



REAGENT INVENTORY SUMMARY

Attached printout

THIS INVENTORY SUMMARY CONSISTS OF ALL METHODS CALIBRATED AND INTENDED FOR USE AT TIME OF INSTALL.

Moum Laken

CAS

CUSTOMER



Dimension EXL200 Performance Evaluation, Annexure1

Following Procedure was Carried out as part of the Performance Qualification:-

1. Calibration of Assay

Checked and found all calibrations within the acceptable CV limits and in range.

See Print out Attached.

2. Internal Quality Control Performance

Two Level Biorad Lyphocheck Assayed Chemistry Control. Checked and found all level Controls to be within the acceptable limits. See print out Attached.

3. Precision Study

A Within Run Precision of replicates were carried out and CV % obtained are within the acceptable limit for the assay as stated in the IFU.

See print out attached.

4, Linearity Study

Linearity study done for AST, BUN, Creatinine, GGT, Glucose.

See Print out attached.

4000



Performance Qualification for Dimension EXL200

With reference to the Annexure 1 and studies carried out in the Laboratory have determined that the analyzer meets all performance criteria and has passed Performance

The System is ready for specific usage.

Protocol performed by

Siemens Ltd.

Signature

Monmi Lakan

Name

: Da I Monmi lakai

Designation: Application Service

Customer Authorization :

Impulse Diagnostic

Signature

: Janta Dos

Name

: Jantu Das

Designation: Technician In-charge

Date: 18/02/20