

To Whom It May Concern

For ISO 15189:2012 and ISO 15189:2014 accredited Laboratories -- requirements regarding "Calibration & Verification Procedures" [1]

All In vitro Diagnostics Products which are manufactured and distributed by Roche Diagnostics GmbH and for which a Free-Sales-Certificate is issued, are CE-marked.

The In-Vitro-Diagnostics Directive of the European Union [2] requires for all CE marked products that the manufacturer assures compliance of the products with the requirements of the In-Vitro-Diagnostics Directive. This means that all processes in development and manufacturing of Roche Diagnostics GmbH products are guided by a Quality Management System. Our Quality Management System is in compliance with the requirements from ISO 13485:2012 [3] + AC:2012 and 21 CFR Part 820 [4].

The mentioned regulations and standards require that the production systems and measuring devices used are qualified and the manufacturing and test procedures are validated. This status has to be assured by scheduled maintenance and by regular qualification resp. validation reviews and updates.

All physical quantities, calibrators and controls used in Roche Diagnostic systems are fully traceable to certified standards or reference materials. The performance of all In-vitro diagnostics systems of Roche Diagnostics GmbH at the customer site is assured if regular Quality Control measurements, cleaning and maintenance procedures as described in the instructions for use or service documentation are performed. By having controlled internal procedures and by running the tasks required in the respective user documentation, all In-vitro diagnostics systems of Roche Diagnostics GmbH will be performed as specified during their defined lifetime.

Additional calibration or verification procedures are NOT required of the user in order to assure the specified performance of every system of Roche Diagnostics GmbH. Only if a user deviates from these manufacturer's recommendations, the user have to establish site-specific calibration and verification procedures as part of his accreditation process.

Mannheim, 18 December 2017

Roche Diagnostics GmbH

**Roche Diagnostics GmbH**  
Sandhofer Straße 116  
D-68305 Mannheim

*[Handwritten signature and scribbles]*



c111 V4.3.0.1835 15260  
admin 26.09.2019 14:33

Abs. Air/Water Calibration:

Abs. Air/Water Calibration.

Cuvette segment is moved to handling position 1.

Insert an empty cuvette segment, then press <OK>.

Cuvette segment is moved to handling position 2.

Insert an empty cuvette segment, close the cover, then press <OK>.

Starting the calibration.

Waiting for completion.

Maintenance action complete.

Wavelength	H2O	H2O cuvette
340	-0.0057	0.0777
378	-0.0041	0.0749
409	-0.0034	0.0731
449	-0.0002	0.075
480	0.0009	0.0748
512	0.0025	0.076
520	0.0026	0.0759
552	0.0037	0.0767
583	0.0048	0.0775
629	0.006	0.0783
652	0.0066	0.0789
659	0.0064	0.0786

Outlier statistics:

Air : 0  
Water : 0  
Diff : 0  
Total : 0

Press <OK> to use these values.  
Then press <Cuvettes> and exchange the two segments.

c111 V4.3.0.1835 15260  
admin 26.09.2019 14:51

Check Pipetting Accuracy:

Place sample '@CHK' on the instrument.  
Created 16 CHECK test orders in sample order '@CHK'.

Cuvette segment is moved to handling position 1.

Insert first cuvette segment, then click OK.

Cuvette segment is moved to handling position 2.

Insert second cuvette segment and close the main cover, then click OK.

System changes to operating and runs CHECK tests. Wait for completion.

# of results = 16, Mean = 1.27131Abs  
SD = 0.00893266, CV = 0.702636%

Diagnostic action completed.



# Roche Professional Services (ISO 9001:2015 certified)

Customer Support Center No: 1800-123-7599 /Toll No: +91-44-30413900



Case No.	CAS-0010478754	Instrument Model	cobas c 111
Order No.	ORD-0013607683	Instrument Serial No.	15260
Contract Type		Finance Status	CASH
Lab/Inst./hosp.Name	Dr.N.V Potdar Dr. Potdar Laboratories		
Customer No.	0052060204		
Contact Name :	Nilkanth Potdar		
Contact Number :	+919822323458		
Address :	Potdar Laboratory,		
City :	Sholapur		

Call Received Date/Time:	07.10.2019 16:15	Call Attended Date/Time:	09.10.2019 10:30
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Job Type	Customer training (follow-up)
Job Description	Follow-up training.
Action Summary	<p>Cause: Follow Up training.                  Workdone: Follow-up training completed. calibeations done. Qc was within in range. Maintenance training was shown and performed.                  Verification: system working fine.                  Customer Satisfaction Rating (1-5): 4</p>

Spare Part Replaced					
Part No	Parts Description	Batch No	Batch Expiry Date	Qty	Invoice Type

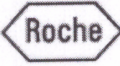
Time Report	
Effective Visit Date : 09.10.2019	Complete Date : 10.10.2019

Date	Type	Time
2019.10.09	Travel Time--Standard	8
2019.10.09	Working Time--Standard	6.5
09.10.10	Travel Time--Standard	2
2019.10.10	Working Time--Standard	8
Total:		24.5

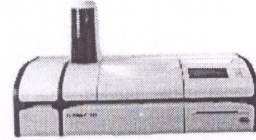
Customer's Signature Name : Potdar Nilkanth	Service Engineer/Application Specialist Name : Shivajirao Mohite
Date: 10.10.2019	Date: 10.10.2019

Disclaimer
<p>1. This Service report has been signed by the authorized representative of your organization.                  2. Discrepancies if any must be intimated within a period of seven (7) days of the receipt of the email, in the absence of which the</p>





cobas® c111



### General Information

Country: INDIA  
Customer Name: Dr. Potdar Laboratories  
Customer Address: First Floor, Shashwat Heights, 519, Shukrawar Peth, Solapur  
Person Responsible for Quality Assurance: Dr. Potdar

### System Information

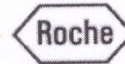
cobas C111 : S/N 15260  
Host provider:  
Software Version: 3.0.185

### Installation Information

Installation Start Date: 26/9/2019  
First Installation: yes  
Reconfiguration: From: To:  
Relocation: From: To:

Roche Responsible Representative : Pravin Kankure





## Installation Qualification:

This document forms the basis of the Qualification Services Certificate. It certifies that the instrument is installed according to the manufacturer's specifications. The report presents and documents the test procedures, the documentation, reference and acceptance criteria used to verify that the system is installed according specifications. The report demonstrated that all installation qualification criteria have been met satisfactorily.

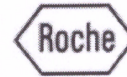
**Notice:** The following tests are to be carried out by trained Roche personnel only.

**Purpose:** The purpose of this test is to confirm that the instrument was delivered undamaged and installed correctly.

Test #	Test	Pass Fail	Signature Date
IQ.1.1	Operator's Manual available	Pass	26/9/2019
IQ 1.2	Environmental parameters met	Pass	26/9/2019
IQ 1.3	Instrument delivered undamaged and complete	Pass	26/9/2019
		Pass	26/9/2019
IQ 1.4	Transport locking successfully removed	Pass	26/9/2019
IQ 1.5	All connections correctly installed	Pass	26/9/2019
IQ 1.6	Instrument positioned according to Installation Manual	Pass	26/9/2019
		Pass	26/9/2019
IQ 1.7	Instrument boot process successfully	Pass	26/9/2019
IQ 1.8	Checksum according to specification	Pass	26/9/2019
IQ 1.9	Mechanical adjustments complete	Pass	26/9/2019
IQ 1.10	Auxiliary components positioned	Pass	26/9/2019
IQ 1.11	Instrument installation check	Pass	26/9/2019
IQ 1.12	Host communication settings checked	Pass	26/9/2019

Test #	Test	Pass Fail	Signature Date
IQ.2	Installation Qualification for cobas c111	Pass	26/9/2019





## Operational Qualification:

This document is the basis of the Qualification Service Certificate. It certifies that the instrument is operating according to the manufacture's specifications. This report presents and documents the test procedures, documentation, references and acceptance criteria used to verify that the specified system is operating according the specifications. The report demonstrates that all operational qualification criteria have been met satisfactorily.

**Purpose:** The purpose of this test is to check that the modules are operating in accordance with the

Test #	Test	Pass Fail	Signature Date
OQ.1	Calibration successfully	Pass	9/10/2019
OQ.2	Quality Control successfully	Pass	9/10/2019
OQ.3	Accuracy check successfully	Pass	9/10/2019

**Deviation Report:** Any discrepancies found during the installation must be documented in the space below. Roche personnel will then investigate the deviation and decide upon the most appropriate action to be taken.

Deviation #1

NONE

Investigation

Action taken

Deviation resolved satisfactorily? specify

Deviation #2

NONE

Investigation

Action taken

Deviation resolved satisfactorily? specify





## Conclusion

All test results are acceptable.  yes

Any deviation or non-conformances observed have been recorded as a deviation and the relevant forms completed.  yes

All acceptance criteria have been met. This equipment is deemed acceptable and the unit is approved for its intended use.  yes

### Comments

ALL RESULTS WITHIN ACCEPTABLE RANGES

Completed by Roche Representative \_\_\_\_\_ Date 9/10/2019

Print Name Shivajirao Mohite Signature *Mohite*

Reviewed by Customer Contact \_\_\_\_\_ Date \_\_\_\_\_

Print Name Dr. Potdar Signature \_\_\_\_\_

Reviewed by Customer Quality Assurance \_\_\_\_\_ Date \_\_\_\_\_

Print Name \_\_\_\_\_ Signature \_\_\_\_\_





## Installation Qualification for cobas<sup>®</sup> c111

### Description

#### IQ.1.1 Operator's Manual available

Check that a copy of the latest version of the Operator's Manual is available. Pass

#### IQ 1.2 Environmental parameters

Ambient temperature in the lab is between 15° and 32 °C Pass

Relative Humidity maximum of 50% at 32 °C and non-condensing Pass

Bacteria free, deionized water < 10 cfu/ml Pass

Water conductivity 1.0 µS/cm or less Pass

Dust and Vibration free Pass

Instrument is not exposed to direct sunlight Pass

#### IQ 1.3 Instrument delivered undamaged and

All covers are undamaged Pass

All accessory boxes are delivered Pass

Instrument does not show any external damage Pass

#### IQ 1.4 Transport locking successfully removed

Unpacking of the Analyzer and accessories without damage to units Pass

#### IQ 1.5 All connections correctly installed

Power supply voltage at the customer facility: YES

UPS system available: yes

Voltage fluctuation less than 230 ±5V Pass

Grounding less than 1.0 V Pass





## Operational Qualification:

### Notice:

The steps described in OQ.1 have to be carried out after a new system installation and after any repair action which requires additional calibration.

If the service action does not affect the measurement performance, only apply steps OQ.2 and OQ.3 of the Operation Qualification.

### Description

#### OQ.1 Calibration

Calibration of all photometric parameters successful	yes
Calibration of all ISE parameters successful (attached printout)	not applicable

#### OQ.2 Quality Control

Specify the type of control used:

BIORAD CHEMISTRY ASSAYED CONTROL

QC of all photometric parameters within acceptable range	yes
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QC of ISE parameters within acceptable range (attached printout)	not applicable
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#### OQ.3.1 Accuracy check for ISE

Perform test with analytical reagents

	Number of det.
Na	21
K	21
Cl	21

Sample solution: PNU (code 300 / Cat. No. 10171735, 10171743, 10651257).  
Fill 21 Hitachi cups with PNU (code: 300) and perform Na, K and Cl tests. Calculate the CV.

Accuracy check for ISE was within acceptable range not applicable



## Carry over

	Nacl	Sample	Nacl	% carryover
Albumin	0.1	2.83	0.12	0.0088
Alk.phosphatase	-0.7	357.9	-0.4	0.014
ALTL	-0.7	83.2	-1	0.02
ASTL	0.7	173.9	0.7	0.003
Bili Direct	0	1.5	0	0
Bili Total	0	3.6	0	0
Cholesterol	0.29	90.8	-0.16	0.0116
Creat	-0.1	4.6	-0.2	0.012
Glucose	-0.21	274.6	-0.26	0.0126
HDL	0.06	22.5	-0.14	0.0114
T.Protein	-0.1	3.9	-0.1	0.011
Triglyceride	0.18	91.9	0.32	0.0068
Uric Acid	0	9.5	0	0
Urea	-0.27	92.04	-0.32	0.0132



Assay	ALB	ALP	ALTL	AST	BIL - D	BIL - T	CHOL	CREAT	GLUCO SE	HDL	TP	TGL	UA	UREA
1	4.27	100.1	26	40	0.4	0.8	236	2	81	59	6.5	190	4.8	32
2	4.38	104	26	41	0.4	0.8	241	2	81	60	6.7	191	4.9	32
3	4.38	103	26	40	0.4	0.8	239	2	81	59	6.6	193	4.9	32.7
4	4.38	103	26	40	0.4	0.8	239	1.9	81	59	6.6	192	4.9	32
5	4.38	104	26	40	0.4	0.9	237	2	81	60	6.6	193	4.9	32
6	4.25	97	26	40	0.4	0.8	232	2	78	58	6.4	184	4.7	32
7	4.33	101	25	41	0.4	0.8	236	1.9	80.8	60	6.6	190	4.8	33
8	4.38	101	27	41	0.4	0.8	240	1.9	81	59	6.7	193	4.9	33
9	4.37	105	26	40	0.4	0.8	239	2	81	59	6.7	194	4.9	33
10	4.38	105	26	41	0.4	0.8	239	1.9	80	59	6.6	191	4.9	33
Mean	4.35	102.31	26	40.4	0.4	0.81	237.8	1.96	80.58	59.2	6.6	191.1	4.86	32.47
SD	0.05011	2.5309	0.4714	0.5164	6E-17	0.03162	2.61619	0.05164	0.95893	0.63246	0.09428	2.84605	0.06992	0.50343
CV	1.15198	2.47375	1.81309	1.27821	1E-14	3.90405	1.10016	2.63468	1.19004	1.06834	1.4285	1.4893	1.4387	1.55045

Unrestricted



	ALB	ALP	ALT	AST	BIL D	BIL T	CHOL	CREAT	GLUC	HDL	TP	TGL	UA	UREA
RUN 1	4.29	103.4	25.3	39.1	0.4	0.9	236.3	2	81.35	58.39	6.5	190.73	4.9	32.6
	4.13	98.4	24.4	39.1	0.4	0.7	226.4	1.9	78	55.9	6.3	183.1	4.7	31.5
	4.17	98.2	25.6	39.1	0.4	0.8	232.8	1.9	82.4	57.5	6.4	191.8	4.7	31.4
	4.3	104.9	25.7	39.1	0.4	0.8	236.1	2	81.2	58.3	6.4	191.2	4.8	31.4
	4.3	102.1	24.1	39	0.4	0.9	237.8	2	80.9	58.5	6.4	191.8	4.8	32.9
RUN 2	4.24	105.5	25	38.7	0.4	0.8	234.9	2.1	81.5	58.2	6.4	191.7	4.8	31.9
	4.4	106	25.6	38.9	0.4	0.8	243.2	2.2	82.7	60	6.6	193.2	4.9	32.2
	4.4	103.8	26.8	42.1	0.4	0.8	245.7	2.2	84.9	61.8	6.6	193.7	5	32
	4.5	104.2	26.6	42.1	0.5	0.8	251.9	2.2	84.2	53.6	6.6	200.1	5	32.9
	4.36	100.2	26.8	39.7	0.5	0.8	241.5	2.2	84.3	62.6	6.5	194	4.9	32
RUN 3	4.33	89.1	28.2	41.7	0.4	0.8	238.6	2	80.9	60.8	6.3	185	4.7	31.6
	4.26	88.7	27.4	41.3	0.4	0.8	237.8	2	79.9	60.6	6.4	185.3	4.7	30.1
	4.4	91.8	27.4	41.1	0.4	0.8	241.6	2	81.9	61.1	6.4	188.3	4.8	30.7
	4.4	93.2	28.6	42.2	0.5	0.8	243.5	2.1	82.3	61.1	6.4	191.6	4.8	31
	4.4	93.7	28.2	42.5	0.5	0.9	246.6	2.1	82	62.1	6.5	190	4.8	31.4
RUN 4	4.2	83.8	27.2	40.6	0.4	0.8	229.7	2	78.8	58.7	5.9	180.4	4.6	30.4
	4.2	86.5	27.5	41.4	0.4	0.8	235.9	2	89.7	60.2	6.2	180.6	4.7	30.8
	4.1	86.1	28.4	42.2	0.4	0.8	239.2	2.1	80.4	60.8	6.1	181.2	4.7	31.4
	4.3	89	27.8	42.1	0.4	0.8	238.5	2.1	79.7	62.4	6.4	183.3	4.9	31.2
	4.2	86	27.9	42.3	0.5	0.9	239.5	2.1	80.4	60.7	6.2	188.9	4.8	30.4
RUN 5	4.2	85.5	27.4	41	0.4	0.8	237.3	2	80.1	60.2	6.3	183	4.7	31.7
	4.3	85.6	26.2	40.8	0.4	0.8	239.2	2.1	81	60.7	6.3	185.2	4.7	29.8
	4.3	86.3	26.6	42.1	0.4	0.9	242.3	2.1	80.5	60.8	6.2	187.4	4.8	30.4
	4.1	83.2	26	41.1	0.4	0.7	232.1	1.9	78.9	59	5.9	180.1	4.6	29.3
	4.3	86.6	25.7	40.8	0.4	0.7	236.8	1.9	80.3	59.1	6	182.3	4.7	30.6
MEAN	4.284	93.672	26.656	40.804	0.42	0.8083	238.61	2.048	81.53	59.724	6.328	187.76	4.78	31.264
SD	0.108	8.074	1.259	1.315	0.041	0.058	5.446	0.096	2.391	2.067	0.197	5.275	0.108	0.926
CV	2.520	8.620	4.724	3.224	9.720	7.220	2.282	4.700	2.933	3.460	3.111	2.810	2.260	2.962