

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 [2A.] which is currently switching to IVD Regulation 2017/746/EU (final timeline: May 26, 2022) [2B.] requires for all CE marked products that the manufacturer assures compliance of the products with the requirements of the mentioned directive or regulation. 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 13485:2016 [3] 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 by 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.

- [1] ISO 15189:2012/ ISO 15189:2014 Medical laboratories — Requirements for quality and competence
- [2] A. Directive 98/79/EC of the European Parliament and of the Council of the 27 October 1998 on vitro diagnostics medical devices;
B. IVD Regulation 2017/746/EU of the European Parliament and of the Council of 5 April 2017 on in vitro diagnostic medical devices and repealing Directive 98/79/EC and Commission Decision 2010/227/EU
- [3] EN ISO 13485:2016 Medical devices — Quality management systems-Requirements for regulatory purposes
- [4] CFR Part 820, Quality System regulations 21 Regulations on medical devices

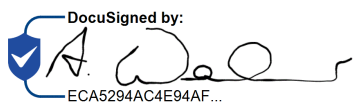
Mannheim, 10. August 2021

Sincerely,

Roche Diagnostics GmbH

i.V./on behalf of the company

ppa/on behalf of the company


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Andrea Weber
Manager Global Regulatory Affairs
Centralised and Point of Care Solutions


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Ralf Zielenski
Head Q&R Compliance, PRRC RDG
Centralised and Point of Care Solutions

Roche Diagnostics GmbH
Sandhofer Straße 116
D-68305 Mannheim



Case No.	CAS-0013934445	Instrument Model	cobas 6000 core unit 150
Order No.	ORD-0017985909	Instrument Serial No.	1288-04
Contract Type	IN-COMPREHENSIVE	Finance Status	RENT
Lab/Inst./hosp.Name	Bioline Laboratory		
Customer No.	0052611292		
Contact Name :	Jasheera Rahman		
Contact Number :	+919961878447		
Address :	21/829, CKS Building,		
City :	Kozhikode		

Call Received Date/Time:	12.08.2021 16:30	Call Attended Date/Time:	22.10.2021 07:00
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Job Type	PM Visit
Job Description	
Action Summary	Cause:Preventive maintenance cobas 6000 core unit Workdone:PM has been performed as per the checklist Verification:Rebooted the instrument.Performed mechanism check Customer Satisfaction Rating (1-5):5

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

Time Report	
Effective Visit Date : 22.10.2021	Complete Date : 22.10.2021

Date	Type	Time
2021.10.22	Travel Time--Standard	0.5
2021.10.22	Working Time--Standard	2
Total:		2.5

Customer's Signature Name : Mrs. Jasheera Rahman	Service Engineer/Application Specialist Name : Thomas Antony
Date: 22.10.2021	Date: 22.10.2021

Disclaimer
<p>1. This Service report has been signed by the authorized representative of your organization.</p> <p>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 report will be final and conclusive.</p> <p>3. This report shall be governed by the laws of India and the courts at Mumbai shall have the exclusive jurisdiction to deal with the disputes arising hereunder.</p>



Case No.	CAS-0013934443	Instrument Model	cobas 6000 c 501 module
Order No.	ORD-0017985906	Instrument Serial No.	1294-14
Contract Type	IN-COMPREHENSIVE	Finance Status	RENT
Lab/Inst./hosp.Name	Bioline Laboratory		
Customer No.	0052611292		
Contact Name :	Jasheera Rahman		
Contact Number :	+919961878447		
Address :	21/829, CKS Building,		
City :	Kozhikode		

Call Received Date/Time:	12.08.2021 16:30	Call Attended Date/Time:	22.10.2021 09:00
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Job Type	PM Visit
Job Description	
Action Summary	Cause:Preventive maintenance cobas c501 Workdone:PM has been performed as per the checklist Verification:Run Qc and samples Customer Satisfaction Rating (1-5):5

Spare Part Replaced					
Part No	Parts Description	Batch No	Batch Expiry Date	Qty	Invoice Type
07783175001 ⁽¹⁾	KIT MAINTENANCE 4 501/502 ⁽¹⁾			1	Free of charge
05521521001 ⁽¹⁾	KIT MAINTENANCE 1 501/502 ⁽¹⁾			1	Free of charge

(1): Customer owned

Time Report	
Effective Visit Date : 22.10.2021	Complete Date : 22.10.2021

Date	Type	Time
2021.10.22	Travel Time--Standard	0.5
2021.10.22	Working Time--Standard	4
Total:		4.5

Customer's Signature Name : Mrs.Jasheera Rahman	Service Engineer/Application Specialist Name : Thomas Antony
Date: 22.10.2021	Date: 22.10.2021



Disclaimer

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