

# Certificate

The Certification Body of  
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization  
**LABOR + TECHNIK**  
**Eberhard Lehmann GmbH**  
**Goerzallee 299**  
**14167 Berlin**  
**Deutschland**

has established and applies a quality management system for medical devices  
for the following scope:

**other in-vitro diagnostics**  
**(see attachment for details)**

Proof has been furnished that the requirements specified in

**EN ISO 13485:2016**

are fulfilled. The quality management system is subject to yearly surveillance.

Effective Date: 2020-02-28  
Certificate Registration No.: SX 60146590 0001  
An audit was performed. Report No.: 21229796 005  
This Certificate is valid until: 2021-08-24

Certification Body



Date 2020-02-28



Dipl.-Ing. (FH) K. Schick

**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**  
Tel.: +49 221 806-1371 Fax: +49 221 806-3935 e-mail: cert-validity@de.tuv.com http://www.tuv.com/safety

**TÜV Rheinland**  
**LGA Products GmbH**  
**Tillystraße 2, 90431 Nürnberg**

**Attachment to  
Certificate**

**Registration No.:** SX 60146590 0001  
**Report No.:** 21229796 005

**Organization:** LABOR + TECHNIK  
Eberhard Lehmann GmbH  
Goerzallee 299  
14167 Berlin  
Deutschland

**Scope:** Manufacturing and Distribution of in-vitro-diagnostics, reagents, controls and calibrators for clinical chemistry, staining solutions for hematology/cytology/bacteriology and specialised trade of photometers as well as other laboratory accessories



**Date:** 2020-02-28

**Certification Body**



*K. Schick*

**Dipl.-Ing. (FH) K. Schick**