TÜVRheinland®

Certificate

The Certification Body of
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization

ScheBo Biotech AG
Netanyastr. 3
35394 Gießen
Deutschland

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

Design and development, manufacture and distribution of immunochemical in vitro diagnostic devices used in the field of oncology and gastroenterology

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:	2018-09-03
Certificate Registration No.:	SX 60131796 0001
An audit was performed. Report No.:	21232272 004
This Certificate is valid until:	2021-09-02

Certification Body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

TÜV Rheinland LGA Products GmbH
TÜVRheinland®
Zertifizierungsstelle

Date 2018-08-24

Dipl.-Ing. Sven Hoffmann

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