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, development, production and sales of in vitro diagnostic medical devices based on immunochemical technologies

Proof has been furnished that the requirements specified in

EN ISO 13485:2012
EN ISO 13485:2012/AC:2012

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

Effective Date:	2015-09-03
Certificate Registration No.:	SX 60103930 0001
An audit was performed. Report No.:	21232272 001
This Certificate is valid until:	2018-09-02

Certification Body

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

TÜV Rheinland LGA Products GmbH
TÜVRheinland®
Zertifizierungsstelle

Date 2015-08-20

Dr. H. Lüdemann

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