

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 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 9001:2008

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

Effective Date: 2015-09-03
Certificate Registration No.: SY 60103931 0001
An audit was performed. Report No.: 21232272 001
This Certificate is valid until: 2018-09-02

Certification Body

Date 2015-08-20




Dr. H. Lüdemann

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