

Certificate

Quality Management System EN ISO 13485:2016

Registration No.:

SX 1763647-1

Organization:

ScheBo Biotech AG

Netanyastr. 3 35394 Gießen Germany

Scope:

Design and development, manufacture and distribution of immunochemical

in vitro diagnostic devices used in the field of oncology and

gastroenterology

The Certification Body of TÜV Rheinland LGA Products GmbH certifies that the organization has established and applies a quality management system for medical devices.

Proof has been furnished that the requirements specified in the abovementioned standard are fulfilled. The quality management system is subject to yearly surveillance.

Report No.:

1089010-120

Effective date: Expiry date:

2021-09-03 2024-09-02

Issue date:

2021-08-19



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