

Certificate

The Certification Body of
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization

Shenzhen Watmind Medical Co., Ltd.
8th Floor, Building A, No.16-1,
Jinhui Road, Jinsha Community
Kengzi subdistrict, Pingshan District
518118 Shenzhen
P.R. China

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

Design and Development, Manufacture and Distribution of
in-vitro diagnostic Medical Devices
(see attachment for products and additional sites included)

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: 2020-12-30
Certificate Registration No.: SX 60150107 0001
An audit was performed. Report No.: 17054604 003
This Certificate is valid until: 2022-07-03

Certification Body



Date 2020-12-30



Wenxiang Zhang

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TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

**Attachment to
Certificate**

Registration No.: SX 60150107 0001
Report No.: 17054604 003

Organization: Shenzhen Watmind Medical Co., Ltd.
8th Floor, Building A, No.16-1,
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518118 Shenzhen
P.R. China

Scope:

Products:

In-vitro diagnostic analyzers and in-vitro diagnostic test kits used in the diagnosis and quantitative detection of cardiac markers, immune status, Thyroid Functions, Auto-Immune Diseases, Fertility testings, Coagulation and Infectious Diseases including point of care in-vitro diagnostic medical devices

Sites included:

Shenzhen Watmind Medical Co., Ltd.
8th Floor, Building A, No.16-1, Jinhui Road, Jinsha Community, Kengzi subdistrict, Pingshan District, 518118, Shenzhen, China

Manufacture of the a.m. products

Shenzhen Watmind Medical Co., Ltd.

Room 106-1 & 107, Shenzhen IC Design & Application Industrial Park, No.1089 Chaguang Road, Nanshan District, Shenzhen, China

Design and Development and Distribution of the a.m. products

Certification Body



Date: 2020-12-30


Wenxiang Zhang

