

Certificate

The Certification Body of
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization
**Tianjin MNCHIP Technologies
Co., Ltd.**
2102,2202, Building 2
No.19 Xinhuan Rd. West, TEDA
300457 Tianjin
P.R. China

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

(See attachment for Scope)

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: 2020-11-19
Certificate Registration No.: SX 60150610 0001
An audit was performed. Report No.: 50334778 002
This Certificate is valid until: 2023-09-02

Certification Body



Date 2020-11-19



TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Tel.: +49 221 806-1371 Fax: +49 221 806-3935 e-mail: cert-validity@de.tuv.com <http://www.tuv.com/safety>

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

**Attachment to
Certificate**

Registration No.: SX 60150610 0001
Report No.: 50334778 002

Organization: Tianjin MNCHIP Technologies
Co., Ltd.
2102,2202, Building 2
No.19 Xinhuan Rd. West, TEDA
300457 Tianjin
P.R. China

Scope: Design and Development, Manufacture and
Distribution of In-vitro Diagnostic Chemistry
Analyzers, In-vitro Diagnostic Test Kits used in the
Detection of Cardiac Markers, Disease Status,
Protein Metabolism and Electrolytes including Home Use

Certification Body



Date: 2020-11-19

Jing Zhang

