

Management System Certificate

Certificate No.: 10000422254-MSC-NA-CHN

Initial Certification Date: 18 May 2021

Valid Until: 18 May 2024

This is to certify that the quality system of:

ChongQing Sunkingdom Medical instrument Co., Ltd.

No.1 XinMao Road (Free Trade Zone), BeiBei District, ChongQing, China.
(Unicode: 915001097874714975)

and the sites as mentioned in the appendix accompanying this certificate

has been found to conform to the Quality Management System standard:

ISO 13485:2016/NS-EN ISO 13485:2016

This certificate is valid for the following scope:

**DESIGN, DEVELOPMENT, MANUFACTURE, DISTRIBUTION, SERVICE AND SALES OF
FUNDUS CAMERA, PERIMETER, OPHTHALMIC SLIT LAMP MICROSCOPE,
OPHTHALMIC ULTRASOUND EXAMINATION INSTRUMENT, VISUAL
ELECTROPHYSIOLOGY, LENSMETER, OCULAR SURFACE ANALYZER AND
APPLANATION TONOMETER.**

Place and date:
Høvik, 18 May 2021

Check Validity



For the issuing office:
DNV Product Assurance AS



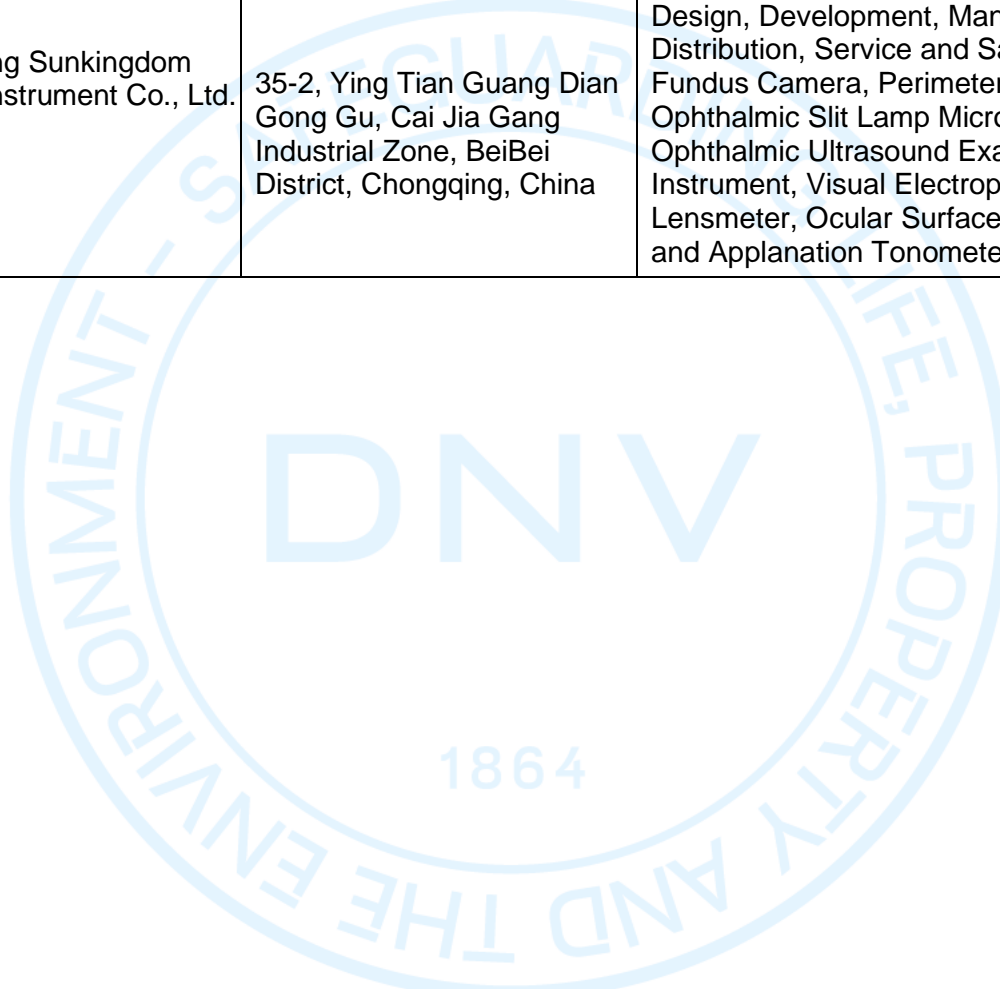
Hazem Tinawi
Technical Reviewer

icate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.

Certificate No.: 10000422254-MSC-NA-CHN Rev. 0.0
Place and date: Høvik, 18 May 2021

Site name	Site address	Site specific scope
ChongQing Sunkingdom Medical instrument Co., Ltd.	No.1 XinMao Road (Free Trade Zone), BeiBei District, ChongQing, China	Legal Registered Address
	35-2, Ying Tian Guang Dian Gong Gu, Cai Jia Gang Industrial Zone, BeiBei District, Chongqing, China	Design, Development, Manufacture, Distribution, Service and Sales of Fundus Camera, Perimeter, Ophthalmic Slit Lamp Microscope, Ophthalmic Ultrasound Examination Instrument, Visual Electrophysiology, Lensmeter, Ocular Surface Analyzer and Applanation Tonometer.

End of Certificate



Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.

Accredited Body: DNV Product Assurance AS, Veritasveien 3, 1363 Høvik, Norway, Tel +47 67 57 88 00, www.dnv.com

ICP-4-5-i1-ISO13485-f1,

rev.0

EC CERTIFICATE

Full Quality Assurance System

Certificate No.:
10000326476-PA-NA-CHN Rev.0.0

Project No.:
PRJN-176150-2019-PA-CHN

Valid Until:
27 May 2024

This is to certify that the quality system of:

Chongqing Sunkingdom Medical Instrument Co., Ltd.

No.1 XinMao Road (Free Trade Zone), BeiBei District, Chongqing, China

For design, production and final product inspection/testing of:

OPHTHALMIC ULTRASOUND EXAMINATION INSTRUMENT, APPLANATION TONOMETER

Has been assessed with respect to:

THE CONFORMITY ASSESSMENT PROCEDURE DESCRIBED IN ANNEX II EXCLUDING SECTION 4 OF COUNCIL DIRECTIVE 93/42/EEC ON MEDICAL DEVICES, AS AMENDED

and found to comply.

Further details of the product(s) and conditions for certification are given overleaf.

Place and date:
Høvik, 06 October 2020

For:
DNV GL PRESAFE AS

Notified Body No.: 2460

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Sholeh Gheissar

The certificate is digitally verified by blockchain technology. For more info, see
www.dnvgl.com/assurance/certificates-in-the-blockchain.html



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Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift om Medisinsk Utstyr" by the Norwegian Ministry of Health and Care Services.

Certificate history:

Revision	Description	Issue Date
0.0	Original Certificate	06-10-2020

Products covered by this Certificate:

Product Description	Product Name	Class
Ophthalmic Ultrasound Examination Instrument	SK-3000A, SK-2000AP	IIa
Applanation tonometer	SK-T, SK-Q, SK-R	IIa

The complete list of devices is filed with the Notified Body

Sites covered by this certificate

Site Name	Address
Chongqing Sunkingdom Medical Instrument Co., Ltd.	Legal Registered Address: No.1 XinMao Road (Free Trade Zone), BeiBei District, Chongqing, China
	Manufacturing Address: 35-2, Ying Tian Guang Dian Gong Gu, Cai Jia Gang Industrial Zone, BeiBei District, Chongqing, China

EU Representative

MedNet EC-REP GmbH, Borkstrasse 10, 48163 Muenster, Germany

Certificate No.:
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Project No.:
PRJN-176150-2019-PA-CHN

Valid Until:
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Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform Presafe of any intended updating of the quality system and Presafe will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. Presafe reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of Presafe.

End of Certificate