

CERTIFICATE OF CONFORMITY

No.: EBO1808103-V262

The following product has been tested by us with the listed standards and found in conformity with the European directive EMC 2014/30/EU and LVD 2014/35/EU.

Applicant: ZENITH LAB (JIANGSU) CO., LTD

Address: No.12, Hongshan Road, Jincheng Industrial Area, Jintan

District, Changzhou, Jiangsu Province, China

Manufacturer: ZENITH LAB (JIANGSU) CO., LTD

Address: No.12, Hongshan Road, Jincheng Industrial Area, Jintan

District, Changzhou, Jiangsu Province, China

EUT: STIRRER

Brand Name: Zenith, Zhengji

Model No.: MHS-C, 85-1, CJ-881, 78-1, 79-1, 78HW-1, HJ-3A, DF-II,

HJ-4, MHS-A, MHS-B, JJ-1, JJ-3, JJ-4, GZ-20, HS-160C,

A-88, FSH-2A, ZJ-100C, CJ series, DF series, HJ series,

MHS series, JJ series, GZ series, HS series, ZJ series

Test Report No.: EBO1808103-E260, EBO1808103-E261

Sufficient samples of the product have been tested and found to be in conformity with

Test Standards: EN 61326-1:2013

EN 61010-1:2010

The test report was carried out from submitted type samples of a product in conformity with the specification of the respective standards. The CE mark as shown below can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of Conformity and compliance with all relevant EC Directives.

Kevin Wang

Laboratory Manager

Issue Date: August 27, 2018





Shenzhen EBO Testing Center

A506, Financial port building, Xin'an Sixth Road, 82th District, Bao'an, Shenzhen, China. Tel: 86-755-33126608 ebo@ebotest.com www.ebotest.com



MANAGEMENT SYSTEM CERTIFICATE

Certificate No.: 117 19 Q0M 0008 R1S

This is to certify the quality management systems of

Zenith Lab (Jiangsu) Co., Ltd.

Unified Social Credit Code 91320413720511570J

Location No. 12 Hongshan Road, Industrial Park, Jincheng Town, Jintan District, Changzhou, Jiangsu

has been assessed and registered as meeting the requirements of

ISO13485: 2016 < Medical device-Quality management

systems - Requirements for regulatory purposes>

Scope of approval

Design, Production and Sales of Medical Centrifuge

Signed by:



First Certification: 28 Jul. 2016 Recertification Date: 15 Jan. 2019 Expiry Date: 27 Jul. 2022

Shanghai Ingeer Certification Assessment Co.,Ltd.

Certification and Accreditation Administration of PRC:CNCA-R-2003-117
Tel: 400-182-9001/+86 21-51114700

Web: www.icas.org.cn

Add: Room 801, HuaDing Mansion, 2368# West Zhongshan Rd.,

Xuhui District, Shanghai, China, 200235



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The ownership of the certificate belongs to Shanghai Ingeer Certification Assessment Co., Ltd. The information & validation of this certificate can be checked on the CNCA website: WWW.CNCA.GOV.CN and ICAS website, or by calling ICAS's clients services Dept. The certificate is only valid when used together with related permits when appropriate. If the organization can't effectively maintain the above management system, ICAS has the right to withdraw the qualification certificate.



CERTIFICATE OF CONFORMITY

No.: EBO1808103-V265

The following product has been tested by us with the listed standards and found in conformity with the European directive EMC 2014/30/EU and LVD 2014/35/EU.

Applicant: ZENITH LAB (JIANGSU) CO., LTD

Address: No.12, Hongshan Road, Jincheng Industrial Area, Jintan

District, Changzhou, Jiangsu Province, China

Manufacturer: ZENITH LAB (JIANGSU) CO., LTD

Address: No.12, Hongshan Road, Jincheng Industrial Area, Jintan

District, Changzhou, Jiangsu Province, China

EUT: INCUBATOR

Brand Name: Zenith, Zhengji

Model No.: DNP-9025A, SPX-150B, BS-1E, SPX-150G, MJ160, BLX-250,

YLX-150B, IB-9052A, DNP-9052A, GNP-9080, LHP-160, CHP-80, PQX-250B, YQX-II, DHG-9030A, DHG-9053A, OV-9053A, DZF-6050, DZF-6090, PPH-140A, GW-070A, SPX Series, BS Series, MJ Series, BLX Series, YLX Series, IB Series, DNP Series, GNP Series, LHP Series, CHP Series,

PQX Series, YQX Series, DHG Series, OV Series, DZF Series,

PPH Series, GW Series, GDW Series, ZJ Series

Test Report No.: EBO1808103-E263, EBO1808103-E264

Sufficient samples of the product have been tested and found to be in conformity with

Test Standards: EN 61326-1:2013

EN 61010-1:2010

The test report was carried out from submitted type samples of a product in conformity with the specification of the respective standards. The CE mark as shown below can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of Conformity and compliance with all relevant EC Directives.

Kevin Wang

Laboratory Manager

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