

Certificate Of Registration

Taizhou Shubeikang Medical Technology Co., Ltd
200meters east of Oulusha Co., Ltd. Yunhu Village, Hengjie Town, Luqiao District,
Taizhou City, Zhejiang, China

Has Completed With The U.S. Food And Drug Administration Pursuant To 21
CFR Part 807: Establishment Registration And Device Listing

Owner/Operator No.: 10074993

Listing Number	Code No.	Proprietary Name	Model
D406816	KPY	Disposable Face Shield	SP-3224, SP-3228
D406815	HOY	Disposable Goggles	SP-800, SP-900
		KN95 masks	BTG-800, BTG-900
D406808	KHA	Disposable Masks	SP-500, SP-600

Huawin will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration is terminated after issuance of this certificate. Huawin makes no other representations or warranties, nor does this certificate make sole benefit it is issued. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration. Huawin assumes no liability to any person or entity in connection with the foregoing.

The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Huawin is not affiliated with the U.S. Food and Drug Administration.



Manager Guy Su
Issue date: June 4, 2020
Expire Date: Dec. 31, 2020

Shenzhen Huawin Testing Certification Co., Ltd.
Add: 7F, U Center, No.743, Zhoushi Road, Bao'an, Shenzhen, China
Http://www.huawinlab.com E-mail: info@huawinlab.com

CERTIFICATE OF NOTIFICATION

This is to certify that, according to the European Council Directive 93/42/EEC, Riomavix S.L. performed all notification duties and responsibilities as the European Authorized Representative:

MANUFACTURER: Taizhou Shubeikang Medical Technology Co.,Ltd.

ADDRESS: 200m east of oulusha Co., Ltd., Yunhu village, Hengjie Town, Luqiao District, Taizhou City, Zhejiang Province, China

The manufacturer has provided Riomavix S.L. with all the appropriate declaration according to the European Council Directive 93/42/EEC including the EC Declaration of Conformity confirming that its medical device, as stipulated here below, is fulfilling the essential requirements of the European Council Directive 93/42/EEC.

Devices: 1. Disposable Goggles 2. Disposable Face Shield

Classification: I

Where the manufacturer affix the CE mark to the device listed they must ensure that all the essential requirements of European Council Directive 93/42/EEC are met.

The notification of abovementioned device has been completed by the European Authorized Representative in Spain. The Spain Competent Authority is notified of the manufacture's device and has allocated registration. The registration number is RPS/759/2020

Executive Director



Issue date: 11/MAY/2020
Cert. No.: R20200527

