ZERTIFIKAT ■ CERTIFICATE ■ СЕРТИФИКАТ ■ CERTIFICADO ■ CERTIFICAT

RIOMAVIX

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: Hangzhou Shanyou Medical Equipment Co., Ltd

ADDRESS: No. 138, Louta Development Zone, Guancun Village, Louta Town, Xiaoshan District, Hangzhou, Zhejiang 311266, 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- Medical Face Mask 2- Surgical Face Mask

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 notifed of the manufacture's device and has allocated registration. The registration number is RPS/745/2020

Executive Director

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Issue date: 9/MAY/2020
Cert. No.: R20200521

CE

Riomavix S.L.
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