NOTIFICATION OF REGISTRATION

This is to certify that, according to the Regulation (EU) 2017/745, Riomavix S.L. performed all notification duties and responsibilities as the European Authorized Representative:

MANUFACTURER: Xiamen J-Brace Medical Equipment Co., Ltd.

ADDRESS: 2F, 179# Tong'an Park Tong'an Industrial Concentration Area Xiamen 361100 P.R.C

The manufacturer has provided Riomavix S.L. with all the appropriate declaration according to the European Council Regulation (EU) 2017/745 including the EC Declaration of Conformity confirming that its medical device, as stipulated here below, is fulfilling the essential requirements of the European Council Regulation (EU) 2017/745.

1.Night Splint 2.Wrist Brace 3.Finger Splint 4.Cervical Collar 5.Ankle Brace

6.Shoulder Brace 7.Back Brace 8.Knee Brace 9.Hip Brace

Classification: I

Where the manufacturer affix the CE mark to the device listed they must ensure that all the essential requirements of European Council Regulation (EU) 2017/745 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/643/2021

Executive Director

RIOMAVIX

Issue date: 1/APR/2021
Cert. No.: R20210402

RIOMA

Riomavix S.L.
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