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: Hunan Cofoe Medical Equipment Co., Ltd.

ADDRESS: Jinlong Industrial Park, Furong North Road, Xiangyin County, Yueyang City, Hunan, China

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.

Walkers (Walkers Frames/ Rollators)

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/1704/2021

Executive Director

Issue date: 16/JUL/2021
Cert. No.: R20210712-4

Riomavix S.L.
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