NOTIFICATION OF REGISTRATION

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

MANUFACTURER: TIANJIN GRAND PAPER INDUSTRY CO.,LTD

ADDRESS: HONGGUANG FARM BEICHEN DISTRICT TIANJIN CHINA

The manufacturer has provided Riomavix S.L. with all the appropriate declaration according to the European Council Regulation (EU)2017/745 including the 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.

Medical Device: Medical Record Paper

Classification: Class 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/177/2024**

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Issue date: 17/Jan/2024
Cert. No.: R20240104-1

Digitally signed by Morozov Maria
Date: 2024.07.19 16:31:38 EEST
Reason: MoldSign Signature
Location: Moldova

CE