EU DECLARATION OF CONFORMITY

Manufacturer: GRI Medical & Electronic Technology Co., Ltd.
1805 Honggao Road
314031 Jiaxing, Zhejiang
PEOPLE'S REPUBLIC OF CHINA

MF's SRN: CN-MF-000004203

EU Authorized Representative: Emergo Europe B.V.
Westervoortsedijk 60,
6827 AT Arnhem,
The Netherlands

AR's SRN: NL-AR-000000116

Device: Patient securement (refer to see the attached 1)

Basic UDI-DI: refer to see the attached 1

Intended purpose: This product family is intended to be used as positioning devices for the patient during various procedures to assist in the positioning and securement of the patient. These are single-use products and are used non-sterile.

Classification: Class I non-sterile, according to MDR 2017/745 Annex VIII, Rule 1

Notified Body Name: N/A

Notified Body Identification Number: N/A

Conformity Assessment Procedure: Article 19,Annex II, Annex III , MDR 2017/745

CE Certificate Number and Validity Period: N/A

Applicable standards or Common Specifications: N/A

The device covered by the present EU declaration is in conformity with the (EU) MDR 2017/745. All supporting documentations are retained under the premises of the manufacturer. We are exclusively responsible for the declaration of conformity.

Signature/Date: Sidonia Dong 2026/07/02

Print Name/Position: Sidonia Dong VP RA&QA Asia Pacific

Place: Jiaxing, China

Attached 1

Part #	Description	Basic-UDI
50-582	Trendelenburg Positioning System_35"x20"x1"_ Non-sterile_6set/cs	69319181F050KY