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: Sharps Management (refer to see the attached 1)

Basic UDI-DI: refer to see the attached 1

Intended purpose: The purpose of GRI Sharps Management devices is to provide the clinician with a physical barrier to microbial and other contamination to protect both the clinician and the patients from contamination transmission during surgical procedures, when sharp devices must be sercured during transfer between uses. It is used sterile.

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

Notified Body Name: TUV SUD

Notified Body Identification Number: 0123

Conformity Assessment Procedure: Annex IX Chapters I and III, MDR 2017/745

CE Certificate: G11 056820 0042 Rev.02

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 2026/03/06

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

Place: Jiaxing, China

Attached 1

Part #	Description	Basic-UDI
30-0003-S	Scalpel Transfer Device_Bule_Sterile_120ea/cs	69319181F012KQ