EU DECLARATION OF CONFORMITY

Slings

Manufacturer	V. Guldmann A/S Graham Bells Vej 21-23A DK-8200 Aarhus N Phone +45 8741 3151 SRN: DK-MF-000003602
Distributor/ Subsidiary	Company
	Address
	Country
	Phone

Hereby declare that the requirements specified in EU Regulation 2017/745 (MDR) regarding medical devices have been fulfilled in relation to the below listed device groups.

The declared medical devices comply where appropriate, with the following European standards
ISO 10535 Hoist for transfer of disabled persons – Requirements and test methods.

Guldmann A/S is certified as meeting quality management systems according with standards ISO 9001 and ISO 14001. Where appropriate, we comply with ISO 13485 Medical Devices – Quality Management Systems – Requirements for regulatory purposes and FDA 21 CFR part 820.

Device Group	**Slings** ABC Slings, Disposable Slings, Repositioning Slings and Lifting Accessories Class I, Rule 1
Basic UDI-DI	15707287slingFE
Intended purpose	The slings are intended for lifting or supporting a person or body parts of a person.

This declaration of conformity is issued under the sole responsibility of the manufacturer.

On behalf of V. Guldmann A/S

Skejby, 2024.01.25

Place and date of issue

Ulrik Møller, Technical Manager

Guldmann™