EU DECLARATION OF CONFORMITY

Portable ramps

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 devices are also in conformity EU Directive RoHS 2011/65/EU and EU Directive REACH - 1907/2006/EC.

The declared medical devices comply where appropriate, with the following European standards DS/EN 12182 Assistive products for persons with disability – General 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	**Portable ramps** Plain ramp, Telescopic ramp, Folding Telescopic ramp, Folding ramp, Wide plain ramp, Wide folding ramp, Wide folding Pro ramp, Telescopic EasyFold ramp, EasyFold Pro ramp, EasyFold Pro3 ramp Class I, Rule 1
Basic UDI-DI	15707287ramp6A
Intended purpose	Portable ramps are intended for helping wheelchair users overcome height differences in the physical surroundings.

On behalf of V. Guldmann A/S

Skejby, 2022.10.10

Place and date of issue

Ulrik Møller, Technical Manager

Guldmann™