EU DECLARATION OF CONFORMITY

Transfer platform

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 REACH - 1907/2006/EC.

The declared medical devices comply where appropriate, with the following European standards DS/EN ISO 21856:2022 Assistive products – 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	**Transfer platform** Guldmann Transfer Platform GTP1, Transfer Belt S, Transfer Belt M, Transfer Belt L Class I, Rule 13
Basic UDI-DI	15707287transferplatPA
Intended purpose	The intended purpose of the GTP1 Transfer Platform product is for a trained helper to manually transfer users with disabilities safely over short indoor distances in standing position.

On behalf of V. Guldmann A/S

Skejby, 2022.11.28

Place and date of issue

Ulrik Møller, Technical Manager

Guldmann™