Schnitzler Rettungsprodukte GmbH & Co. KG
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EU Declaration of Conformity

(Status: 06_2025-V4)

We,

Schnitzler Rettungsprodukte GmbH & Co KG

Rudolf-Diesel-Str.:7 - 53859 Niederkassel

Manufacturer SRN: DE-MF000005349

declare under their sole responsibility, that the Class I medical devices listed below (non-sterile and without measuring function) according to MDR Annex VIII, Rule 1 with the intended use as:

Stretcher chair/carrying chair Secure the patient in a stretcher chair/carrying chair with a belt system during transport from the accident site to the vehicle

with the basic UDI-DI: 426038867GurtsystemNZ

comply with the relevant provisions of the EU Medical Devices Regulation (EU) 2017/745 and, where applicable, other relevant Union legislation

The product group for **Stryker** carrying chair models: **Stair Pro; Stair Pro-Tread-System** includes the following products:	
Trade name:	**Article no:**
RD-RHS-STUHL-STR Hygiene+	2-428-h+
Die Produktgruppe für Tragestuhl des Hersteller **Stryker** modell: **Xpedition** umfasst folgende Produkte:	
Gurtsatz H+ für Stryker-Tragestuhl, XPEDITION	9-428-H+
Kopfstütze H+ für Stryker XPEDITION-Tragestuhl	9-428-KS-H+

The product group for **Ferno** carry chair models: **Fast R; EZ Glide 59THP** includes the following products:			
Trade name:	**Article no:**	**Trade name**	**Article no:**
Gurtsatz FER Hygiene+ Fast R	3-428-H+	Gurtsatz FER Hygiene+ EZ GLIDE 59TPH	4-428-H+

The product group for stretcher chairs from the manufacturer **Stollenwerk: Stollenwerk stretcher chairs; Dlouhy stretcher chairs; Utila stretcher chairs ALS 103/ALS 328/ALS 428/ALS 300)** includes the following products:	
Trade name:	**Article no:**
Gurtsatz Hygiene+ für Tragestuhl	X-429-H+
RD-RHS-SESSEL Kniegurt f. ALS428 u. ALS300	3-429-30-H+

- The conformity assessment procedure in accordance with Article 19 and Annex IV has been carried out. The technical documentation has been prepared in accordance with Annexes II and III. With this EU Declaration of Conformity, we as a manufacturer of medical devices of the Class I, conformity without involvement of a notified body according to MDR Article 52, paragraph (7)

- Applicable harmonised standards, national standards or other regulatory documents:
  - EN ISO 13485 Medical device quality management systems
  - EN ISO 14971 - Application of risk management to medical devices
  - EN 1865-1 - Ambulance transport equipment in ambulances

- Authorised representative:
  - **[CH] Ambu-Tech AG**, Lettenstrasse: 6c, 6343 Rotkreuz, Reg. No.: CHRN-AR-20002714

Niederkassel, 17 June 2025

Lars Quadt
PRRC according to Art.15 MDR