Certificate of CE-Registration

This is to certify that, in accordance with Regulation (EU) 2017/745 on medical devices, mdi Europa GmbH, SRN DE-AR-000006218 agree to perform all duties and responsibilities as the Authorized Representative for

C-A-T Resources, LLC
483 Lakeshore Parkway
SC 29730 Rock Hill
USA
US-MF-000017612

as stipulated and demanded by the afore-mentioned Regulations. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

CND Code	Description	Classification	Registration Number
C900103	**Basic UDI-DI: 08603620024CR00770XQX** Tourniquet - Arterial Access Haemostasis, Percutaneous Systems	I	DE/CA09/00070170

mdi Europa verified the conformity assessment procedure and that the manufacturer has drawn up all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the general safety and performance requirements of Regulation (EU) 2017/745.

Signed on 14 April 2023

Werner Sander
President