

EU - Declaration of Conformity



We, **X-CEN-TEK GmbH & Co. KG**, Westerburger Weg 30 - D-26203 Wardenburg, declare under our sole responsibility that the medical devices of the product group

PAX Treatment Systems

Basis UDI-DI: 426074548 01625019 A6

complies with the relevant provisions of Regulation (EU) 2017/745 (MDR) on medical devices and, where applicable, other relevant Union legislation

The product group includes the following medical devices

Commercial name	Article No.	Commercial name	Article No.
PAX-Extremities-Tourniquet P-E-T – fluorescent yellow	138103908	PAX-Extremities-Tourniquet P-E-T – black	138103903
PAX-Extremities-Tourniquet P-E-T – olive	138103921		

Intended use of the product group: Treatment of injuries and illnesses - to tie off the blood flow in cases of severe, life-threatening bleeding of the upper and lower extremities that cannot be controlled in any other way or that, depending on the situation, represent the best alternative therapy

According to Annex VIII, Rule 1 MDR, all devices in the product group are class 1 medical devices and the applicable essential safety and performance requirements according to Annex I MDR are fulfilled. A conformity assessment procedure according to Annex II & III MDR has been carried out.

Applicable harmonized standards, national standards, or other regulatory documents:

EN ISO 13485 – Medical devices – Quality management systems – Requirements for regulatory purposes

EN ISO 14971 – Medical devices – Application of risk management to medical devices

This EU Declaration of Conformity is valid until **25.05.2029**

Wardenburg, 22.05.2026

Dr. med. Thomas Castner

Dep. PRRC according to Art.15 MDR

Manufacturers SRN: DE-MF-000009521

Version 1.1	BasDok erstellt: TC-05.04.2021	Freigabe QMB: TC-10.08.2023	Dok erstellt: ME	Freigabe VP: TC - 22.05.2026
Datei: PAX CE KE-EN TQ 05-26.docx			Anlage: 18.05.2026	Stand: 22.05.2026
Firma X-CEN-TEK GmbH & Co.KG – 26203 Wardenburg			QM-System nach EN ISO 13485	Seite 1 von 1