Declaration of conformity No. EU-01

To the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC

Manufacturer *Brand name*	PI KRYVORUCHKO ROMAN PRO STERIL
Address	Vinnytsia, str. K. Vasylenko, building 6A, sq. 44, Ukraine
Manufacturing address	21000, Vinnytsia, str. Keletska, bldg. 117 b, Ukraine
EU Authorized Representative, address	**GRONADA LTD** Eisiskių ave. 47-312 Vilnius, Lithuania +370 609 52385 / www.pro-steril.eu
SRN	LT-CA01/MDD/1-00048/23 LT-CA01/MDD/1-00044/23 LT-CA01/MDD/1-00045/23 LT-CA01/MDD/1-00046/23 LT-CA01/MDD/1-00047/23
Medical Device	Disposable sterilization pouches (UDI: 482027015SP0012J) Disposable sterilization rolls (UDI: 482027015SR0012Y)
Classification (MDR, VIII, Rule I)	*Class I (non-sterile, without measurement function)*
Applied Technical Regulation	*Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC Annex VIII, Rule 1*
GMDN	*13735*
Date of signing, version:	*24 December 2022, Version 1*
Declaration of conformity is valid until:	*23 December 2027*

LIETUVOS RESPUBLIKA UAB „Gronada" JSC REPUBLIC OF LITHUANIA

24 December 2022, Vilnius, LT
(Place and date of issue)

Irena Urbanavičienė Director
First name, second name, position (signature)

CE