Office for Registration
of Medicinal Products, Medical Devices and Biocidal Products

Al. Jerozolimskie 181C, 02-222 Warsaw, Poland; Phone +48 22 492-11-00, fax +48 22 492-11-09
NIP 521-32-14-182 REGON 015249601

Warszawa, 01 -08- 2023

CERTIFICATE OF FREE SALE No. 530/2023

President of the Office for Registration of Medicinal Products, Medical Devices and Biocidal Products based on Article 55 of the Regulation (EU) 2017/746 of the European Parliament and of the Council of 5 April 2017 on *in vitro* diagnostics medical devices and repealing Directive 98/79/EC and Commission Decision 2010/227/EU (OJ L 117 z 5.5.2017, p. 176) pursuant to art. 30 of the Act of April 7, 2022 on medical devices (Journal of Laws of 2022, items 974) in connection with the application for a certificate of free sale made by the

"MPW MED. INSTRUMENTS" SPÓŁDZIELNIA PRACY
(*applicant for certificate of free sale*)

certifies that the in vitro diagnostics medical device listed below :

Name of the device	Type
Refrigerated laboratory centrifuge MPW-380R	MPW-380R
Notified body certificate number	Not applicable
Basic UDI-DI code	590538636-IVD-CEN-012-65

manufactured by :

"MPW MED. INSTRUMENTS" SPÓŁDZIELNIA PRACY
ul. Boremlowska 46, 04-347 Warszawa, Poland
(*identification of the manufacturer*)

on the basis of the statement of the manufacturer is CE marked at the sole responsibility of the manufacturer. The medical device CE marked in conformity with the Regulation (EU) 2017/746 of the European Parliament and of the Council can be placed on the market and put into service in the European Union. Export of the product is not prohibited.

PREZES URZĘDU REJESTRACJI PRODUKTÓW LECZNICZYCH WYROBÓW MEDYCZNYCH I PRODUKTÓW BIOBÓJCZYCH * 2 *

President of the Office

On behalf of the President ,
Vice-President for Medical Devices

Sebastian Migdalski