



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

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NIP 521-32-14-182 REGON 015249601

Warsaw, 2015-09-02

**CERTIFICATE OF FREE SALE No. 379/2015**

In reference to application for a free sale certificate made by the

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

it is concluded that the in vitro diagnostic medical device listed below:

Name of the medical device	Type
Refrigerated and heated laboratory centrifuge	MPW-260RH

manufactured by:

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

on the basis of the statement of the manufacturer the aforementioned in vitro diagnostic medical device is CE marked at the sole responsibility of the manufacturer. The in vitro diagnostic medical device CE marked in conformity with the act of 20<sup>th</sup> May 2010 on medical devices (Official Journal of Laws from 2015, item 876) which implements Directive 98/79/EC can be placed on the market and put into service in the territory of Republic of Poland. Export of the above product is not prohibited.

President of the Office

z upoważnienia Prezesa  
DYREKTOR  
Departamentu Informacji o Wyrobach Medycznych

  
Elżbieta Maciejewska