



**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

Warsaw, 2015-09-01

**CERTIFICATE OF FREE SALE No. 369/2015**

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
Laboratory centrifuge	MPW-54

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*



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Warsaw, 2015-09-01

**CERTIFICATE OF FREE SALE No. 370/2015**

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
Laboratory centrifuge	MPW-55

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



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Warsaw, 2015-09-01

**CERTIFICATE OF FREE SALE No. 371/2015**

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
Laboratory centrifuge	MPW-56

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 Informatyki i Wyrzobach Medycznych

*Etielista-Medolewska*



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Warsaw, 2016-09-20

**CERTIFICATE OF FREE SALE No. 463/2016**

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 laboratory centrifuge MPW-150R	-

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 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 as amended) 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

*[Signature]*  
Elżbieta Mioduska



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NIP 521-32-14-182 REGON 015249601

Warsaw, 2015-09-01

**CERTIFICATE OF FREE SALE No. 372/2015**

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
Laboratory centrifuge	MPW-215

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

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Elzbieta Maciejewska



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Warsaw, 2015-09-01

**CERTIFICATE OF FREE SALE No. 373/2015**

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
Laboratory centrifuge	MPW-223a

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.

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*Eldzista Maciejewska*





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Warsaw, 2015-09-02

**CERTIFICATE OF FREE SALE No. 374/2015**

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
Laboratory centrifuge	MPW-223c

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.

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Warsaw, 2015-09-02

**CERTIFICATE OF FREE SALE No. 375/2015**

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
Laboratory centrifuge	MPW-223e

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.

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Warsaw, 2015-07-09

**CERTIFICATE OF FREE SALE No. 301/2015**

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 medical device listed below:

Name of the medical device	Type
Laboratory centrifuge MPW-223es	-

manufactured by:

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

on the basis of the statement of the manufacturer the aforementioned medical device is CE marked at the sole responsibility of the manufacturer. The medical device CE marked in conformity with the act of 20<sup>th</sup> May 2010 on medical devices (Official Journal of Laws, No. 107, item 679 as amended) 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.

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Warsaw, 2015-09-02

**CERTIFICATE OF FREE SALE No. 376/2015**

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
Laboratory centrifuge	MPW-251

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.

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Warsaw, 2015-09-02

**CERTIFICATE OF FREE SALE No. 377/2015**

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
Laboratory centrifuge	MPW-260

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

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*Etielista Maciejewska*



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Warsaw, 2015-09-02

**CERTIFICATE OF FREE SALE No. 378/2015**

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 laboratory centrifuge	MPW-260R

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  
*Ełżbieta Maciejewska*



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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



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Warsaw, 2015-09-02

**CERTIFICATE OF FREE SALE No. 380/2015**

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
Laboratory centrifuge	MPW-351e

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





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Warsaw, 2015-09-01

**CERTIFICATE OF FREE SALE No. 366/2015**

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
Laboratory centrifuge MPW-352, catalogue number 10352	-

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 Madzińska*



**Office for Registration  
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Warsaw, 2015-09-01

**CERTIFICATE OF FREE SALE No. 367/2015**

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 laboratory centrifuge MPW-352R, catalogue number 10352R	-

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

*Elzbieta Maciejewska*



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Warsaw, 2015-09-01

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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-352RH, catalogue number 10352RH	-

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

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DYREKTOR  
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Eżbieta Maciejewska



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Warsaw, 2015-09-02

**CERTIFICATE OF FREE SALE No. 381/2015**

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
Laboratory centrifuge	MPW-380

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

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DYREKTOR  
Departamentu Informacji o Wymaganiach Medycznych  
  
Elżbieta Maciejewska



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Warsaw, 2015-09-02

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**„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 laboratory centrifuge	MPW-380R

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

*Ełżbieta Maciejewska*