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, 2022-02-28

CERTIFICATE OF FREE SALE No. 114/2022

President of the Office for Registration of Medicinal Products, Medical Devices and Biocidal Products based on Article 60 of the Regulation (EU) 2017/745 of the European Parliament and of the Council pursuant to art. 67 of the Act of May 20, 2010 on medical devices (Journal of Laws of 2021, items 1565) in connection with the application for a certificate of free sale made by:

MyWam Kupiec Bartold Angres Sp. J.
(*applicant for certificate of free sale*)

certifies that the medical device listed below :

Name of the device	Type
Specialist stroller Grizzly	Not applicable
Notified body certificate number	Not applicable
Basic UDI-DI code	5901122279wozekinw-specC5

manufactured by :

MyWam Kupiec Bartold Angres Sp. J.
Ul. Szczecińska 10, 41-516 Chorzów, 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) 2017/745 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.

President of the Office

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

Sebastian Migdalski

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, 2022-02-28

CERTIFICATE OF FREE SALE No. 118/2022

President of the Office for Registration of Medicinal Products, Medical Devices and Biocidal Products based on Article 60 of the Regulation (EU) 2017/745 of the European Parliament and of the Council pursuant to art. 67 of the Act of May 20, 2010 on medical devices (Journal of Laws of 2021, items 1565) in connection with the application for a certificate of free sale made by:

MyWam Kupiec Bartold Angres Sp. J.
(*applicant for certificate of free sale*)

certifies that the medical device listed below :

Name of the device	Type
Specialist stroller PEGAZ	Not applicable
Notified body certificate number	Not applicable
Basic UDI-DI code	5901122279wozekinw-specC5

manufactured by :

MyWam Kupiec Bartold Angres Sp. J.
Ul. Szczecińska 10, 41-516 Chorzów, 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) 2017/745 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

President of the Office

On behalf of the President
Vice-President for Medical Devices

Sebastian Migdalski

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, 2022-02-28

CERTIFICATE OF FREE SALE No. 115/2022

President of the Office for Registration of Medicinal Products, Medical Devices and Biocidal Products based on Article 60 of the Regulation (EU) 2017/745 of the European Parliament and of the Council pursuant to art. 67 of the Act of May 20, 2010 on medical devices (Journal of Laws of 2021, items 1565) in connection with the application for a certificate of free sale made by:

MyWam Kupiec Bartold Angres Sp. J.
(*applicant for certificate of free sale*)

certifies that the medical device listed below :

Name of the device	Type
Specialist stroller MOUSE	Not applicable
Notified body certificate number	Not applicable
Basic UDI-DI code	5901122279wozekinw-specC5

manufactured by :

MyWam Kupiec Bartold Angres Sp. J.
Ul. Szczecińska 10, 41-516 Chorzów, 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) 2017/745 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 I PRODUKTÓW BIOBÓJCZYCH WYROBÓW MEDYCZNYCH

President of the Office
On behalf of the President
Vice President for Medical Devices
Sebastian Migdalski

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, 2022-02-28

CERTIFICATE OF FREE SALE No. 116/2022

President of the Office for Registration of Medicinal Products, Medical Devices and Biocidal Products based on Article 60 of the Regulation (EU) 2017/745 of the European Parliament and of the Council pursuant to art. 67 of the Act of May 20, 2010 on medical devices (Journal of Laws of 2021, items 1565) in connection with the application for a certificate of free sale made by:

MyWam Kupiec Bartold Angres Sp. J.
(*applicant for certificate of free sale*)

certifies that the medical device listed below :

Name of the device	Type
Specialist stroller MEWA	Not applicable
Notified body certificate number	Not applicable
Basic UDI-DI code	5901122279wozekinw-specC5

manufactured by :

MyWam Kupiec Bartold Angres Sp. J.
Ul. Szczecińska 10, 41-516 Chorzów, 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) 2017/745 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.

President of the Office

On Behalf of the President
Vice-President for Medical Devices

Sebastian Migdalski

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, 2022-02-28

CERTIFICATE OF FREE SALE No. 117/2022

President of the Office for Registration of Medicinal Products, Medical Devices and Biocidal Products based on Article 60 of the Regulation (EU) 2017/745 of the European Parliament and of the Council pursuant to art. 67 of the Act of May 20, 2010 on medical devices (Journal of Laws of 2021, items 1565) in connection with the application for a certificate of free sale made by:

MyWam Kupiec Bartold Angres Sp. J.
(*applicant for certificate of free sale*)

certifies that the medical device listed below :

Name of the device	Type
Specialist stroller YETI	Not applicable
Notified body certificate number	Not applicable
Basic UDI-DI code	5901122279wozekinw-specC5

manufactured by :

MyWam Kupiec Bartold Angres Sp. J.
Ul. Szczecińska 10, 41-516 Chorzów, 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) 2017/745 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.

President of the Office
On behalf of the President
Vice-President for Medical Devices
Sebastian Migdalski