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

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NIP 521-32-14-182 REGON 015249601

Warsaw, 2017-09-19

CERTIFICATE OF FREE SALE No. 660/2017

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

Przedsiębiorstwo Elektroniki Profesjonalnej IGEL Jerzy Kowaliński
(applicant for certificate of free sale)

it is concluded that the medical device listed below:

Name of the device	Type
Patient Monitor	ICARD M

manufactured by:

Przedsiębiorstwo Elektroniki Profesjonalnej IGEL Jerzy Kowaliński
ul. Chorzowska 64, 44-100 Gliwice, Poland
(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 20th May 2010 on medical devices (Official Journal of Laws from 2017, item 211) which implements Directive 93/42/EEC 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.

On behalf
President of the Office

VICE-PRESIDENT
for Medical Devices
Sebastian Migdalski

Certificate of Free Sale is issued pursuant to the provisions of the art. 67 of the Act on Medical Devices of 20th May 2010 (Official Journal of Laws from 2017, item 211)