EU DECLARATION OF CONFORMITY

Manufacturer: **SIMON Sp. z o.o.**
ul. Pojezierska 97B
91-341 Łódź

SRN (Single Registration Number): PL-MF-000035415

Hereby declares that:

SIMON

bearing CE mark is a Class I medical device, Rule 1 in accordance with Annex VIII of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices and fulfills the basic requirements specified in this Regulation.

The conformity assessment was done according in Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices.

Medical device is in conformity with the following harmonized standards:

- EN 21856:2022
- EN ISO 14971:2019
- EN ISO 20417:2021
- EN ISO 15223-1:2021
- EN 62336-1:2015

Basic UDI-DI: 59057234387SIMONBT

SIMON is approved to Regulation No. 129 of the Economic Commission for Europe of the United Nations (UNECE) — Uniform provisions concerning the approval of Enhanced Child Restraint Systems used on board of motor vehicles - Supplement 4 to the 03 series of amendments.

EU declaration of conformity is issued under the sole responsibility of the manufacturer.

On behalf of the manufacturer:

Witold Czybis, CEO
Signature:

Manufacturer's seal:

SIMON Sp. z o.o.
ul. Pojezierska 97b
91-341 Łódź
NIP: 727-283-79-16, REG. 383847076
KRS: 0000793362

Łódź, 12th of December 2024