

EC DECLARATION OF CONFORMITY

Number: PSEN0025
Version: 03



1. Product - instrument Type / Model:

Electrically operated hospital bed – *Mimi* / 131

2. Name and address of the manufacturer:

Commercial name	LINET spol. s r.o.
Registered address	Želevčice 5, 274 01 Slaný, Czech Republic
Reg. No.	00507814
Telephone	+420 312 576 111
Fax	+420 312 522 668

3. This declaration of conformity is issued under the sole responsibility of the manufacturer.

4. Object of declaration:

Product:	Mimi
Description and function designation:	Mechanically operated infant crib intended for neonatal and/or postneonatal departments (standard care, acute care and long term care). This EC conformity declaration also covers all applicable accessories approved by manufacturer.
Classification of the product as the medical device:	Class I non sterile, without measuring function, according to annex IX of Government Order No.54/2015 Coll. (MDD 93/42/EEC) – rule 1

5. The object of the declaration described above is in conformity with the relevant Union harmonization legislation:

- Act No. 268/2014 Coll., on Medical Devices (Directive 93/42/EEC)
- Act No. 350/2011 Coll., on chemical substances and mixtures (Regulation (EC) No 1907/2006)
- Government Order No.54/2015 Coll., with is specifies technical requirements for medical devices (Directive 93/42/EEC)
- Applicable requirements of Government Order No.176/2008 Coll., on machinery devices (Directive 2006/42/EC)

6. References to the relevant harmonized standards used or references to the other technical specifications in relation to which conformity is declared:

EN ISO 14971:2012, EN ISO 10993-5:2009, EN ISO 10993-10:2013

Place and date of declaration issue: Slaný, 20.07.2018

Signed for and on behalf of LINET spol. s r.o.

Ing. Tomáš Kolář, Managing Director