

EU – Declaration of Conformity

- Name/address of the manufacturer: MAPA GmbH
Industriestraße 21 – 25
27404 Zeven, Germany

SRN: DE-MF-000017639
- We declare under our sole responsibility that for the designated product, which has been manufactured in accordance with the Technical Documentation

TD 032 Revision 5

and which is documented in the batch documentations, complies with the provisions of the following directives / regulations:

(EU) 2017/745 Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC

(EU) No 10/2011 Commission Regulation (EU) No 10/2011 of 14 January 2011 on plastic materials and articles intended to come into contact with food

(EC) No 1907/2006 Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH) and establishing a European Chemicals Agency

- Basic-UDI-DI: 400860 MIPUMA 0000 01GW
- Product and trade name: NUK Manual Breast Pump Soft&Easy
Article number: 10.252.086
- Medical device class: I



- The conformity of the listed products with the essential protection requirements of the Directives/Regulations is demonstrably and fully in compliance with the following harmonized standards:

MAPA GmbH

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County Court Tostedt HRB 120049 · General Manager: Dr. Ralf Holschumacher, Sean Beckstrom



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| 7. | DIN EN ISO 10993-1
:2021 | Biological evaluation of medical devices - Part 1:
Evaluation and testing within a risk management
process |
| | DIN EN ISO 15223-1
:2017 | Medical devices - Symbols to be used with medical
device labels, labelling and information to be
supplied
- Part 1: General requirements |
| | EN 14350 | Child use and care articles - Drinking equipment |
| 8. | Notified body | Not applicable for medical devices class I |
| 9. | Additional information: | Document validity until [yyyy-mm-dd]: 2025-05-31 |
| 10. | Place of issue, Date [yyyy-mm-dd]: | Zeven, 2021-05-31 |

 i.A. Guenter STEITZ (Quality Management)
 Signed for and on behalf of Alexander Du Chesne (Director Quality Management)

