

## EU – Declaration of Conformity

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

SRN: DE-MF-000017639

2. 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

3. Basic-UDI-DI: 400860 MIPUMA 0000 02GW

4. Product and trade name: NUK JOLIE Manual Breast Pump

Article numbers:  
10.252.090  
10.252.116  
10.252.131  
10.252.133  
10.749.078



**MAPA GmbH**

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



5. Medical device class: I
6. 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:
7. EN 62366-1:2017 Medical devices -Part 1: Application of usability engineering to medical devices
- DIN EN ISO 10993-1:2021 Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process
- EN 14350 Child use and care articles - Drinking equipment
- 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
8. Notified body Not applicable for medical device of class I
9. Additional information: Document validity until [yyyy-mm-dd]: 2025-04-13
10. Place of issue, Date [yyyy-mm-dd]: Zeven, 2021-04-13

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

