Signed by /
Podpisano przez:

Robert Łupicki

Date / Data:
2023-11-07
14:26

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer:

Ningbo MFLAB Medical Instruments Co., Ltd.
No. 508, Yindong Road(N), Yinzhou Economic Development Zone, 315145 Ningbo, China

Name and address of European Representative:

Shanghai International Holding Corp. GmbH(Europe)
Eiffestrasse 80, 20537 Hamburg, Germany
Tel:+49-40-2513175 Fax: +49-40-255726
E-mail:shholding@hotmail.com

We declare under our sole responsibility that the medical device:

1	**PVC free Anesthesia Mask**	**IIa, rule 2**
2	**Anesthesia Masks**	**IIa, rule 2**
3	**Laryngeal Mask Airways**	**IIa, rule 2**
4	**Resuscitation Masks**	**IIa, rule 2**
5	**Silicone/SEBS/PVC Manual Resuscitators**	**IIa, rule 2**
6	**Suction Catheters**	**IIa, rule 5**
7	**Oxygen Masks for Single Use**	**IIa, rule 2**
8	**Oxygen Masks with Reservoir Bags for Single Use**	**IIa, rule 2**
9	**Nasal Oxygen Cannulas**	**IIa, rule 5**
10	**Nebulizer Masks for Single Use**	**IIa, rule 2**
11	**Silicone/SEBS/PVC Manual Resuscitators (Handle for resuscitator is one part of this item)**	**IIa, rule 2**

according to annex IX of directive 93/42/EEC

meets the provisions of the directive 93/42/EEC Annex IV and its transpositions in national laws which apply to it. The declaration is valid in connection with the "final inspection report" of the device.

Conformity assessment procedure: **93/42/EEC Annex V**

Registration No.: **DD 60149456 0001**

Notified Body:

TÜV Rheinland LGA Products GmbH
Tillystraße 2
90431 Nürnberg
Deutschland
CE 0197

Ningbo 2021-5-15

Place, date

Li Yazeng (General manager) Li Yazeng

Name and function