



EU DECLARATION OF CONFORMITY according to Regulation (EU) 2017/745

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|---|---|
| Manufacturer: | FIAB SpA |
| Registered address: | Via Costoli 4, 50039 Vicchio (FI), Italia |
| Single Registration Number: | IT-MF-000005988 |
| Basic UDI-DI: | 803300326101000004K7 |
| Product name/ Intended Purpose | Medium concentration oxygentherapy mask |
| Models: | See list in Attachment |
| Technical Documentation File | TDF 101 |
| Risk Class (MDR Annex VIII): | IIA |
| Conformity assessment procedure performed: | Annex IX – Conformity assessment based on a quality management system (Chapter I) |
| Notified Body | BSI Group The Netherlands B.V. 2797 |
| Certificate(s) issued | EU quality management system certificate MDR 747884 R000 |
| Technical standards and/or Common Specifications applied: | EN 62366 [2015] - EN ISO 10993-1 [2020] - EN ISO 13485 [2016] - EN ISO 13544-1 [2007+A1:2009] - EN ISO 13544-2 [2002+A1:2009] - EN ISO 13544-3 [2001+A1:2009] - EN ISO 14971 [2019] - EN ISO 15223-1 [2021] - EN ISO 20417 [2021] |

With this Declaration of Conformity, issued under the sole responsibility of FIAB SpA as the Manufacturer, we hereby declare

- that the medical devices specified meet the provision of the Regulation (EU) 2017/745 for medical devices
- that the procedures of FIAB quality management system according to ISO 13485 have been followed, Certificate of Registration no.MD77846 issued by BSI
- that the products do not contain medicinal substances, elements of animal origin or their derivatives, human blood derivatives, and are latex free

Signature:

Vicchio, 03/02/2023

Alberto Calabrò
Managing Director

Declaration Code EU-00000073-101

First issued: 03/02/2023

Last revised: 03/02/2023

Cod 99500201MD4A

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Attachment of EU Declaration of Conformity – List of models

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| OS/100 - OS/100A-30 - OS/100D - OS/100N - OS/100P - OS/100PD |
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| Declaration Code | EU-00000073-101 | First issued: | 03/02/2023 |
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