



Name of manufacturer **Eschenbach Optik GmbH
Fürther Straße 252, 90429 Nürnberg
Germany**

Single Registration Number (SRN) **DE-MF-000007711**

We declare under sole responsibility that the product

Name of product **spectacle frame / models see annex**

Basic UDI-DI **4064158FRA-MET-FULL-001BR, 4064158FRA-PLA-FULL-0024U,
4064158FRA-MIX-FULL-003KD, 4064158FRA-OTH-FULL-004HW,
4064158FRA-MET-HALF-005YE, 4064158FRA-PLA-HALF-006RH,
4064158FRA-MIX-HALF-00785, 4064158FRA-OTH-HALF-0086N,
4064158FRA-MET-LESS-009CK, 4064158FRA-PLA-LESS-01055,
4064158FRA-MIX-LESS-011KN, 4064158FRA-OTH-LESS-012J7,
4064158FRA-MET-OTH-013J3, 4064158FRA-PLA-OTH-014X4,
4064158FRA-MIX-OTH-015A6, 4064158FRA-OTH-OTH-016AM**

Nomenclature **GMDN Code 32816
EMDN Code Q02100201, EMDN Code Q02100202
EMDN Code Q02100203, EMDN Code Q02100299**

Risk class **Class I – non sterile, no measuring function**

Conformity assessment procedure **pursuant to article 52 (7) for medical devices**

conforms with the following regulations:

Regulation (EU) 2017/745 of the European Parliament and of the Council on medical devices

Regulation (EC) No 1907/2006 on the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH)

Complies with European standard EN ISO 12870

We operate a systematic procedure to monitor the product after placing it on the market.

Nuremberg, 05.06.2024
Place and date of issue

sgd. P. Braunhofer
**Name,
Dr. Peter Braunhofer
CEO**

sgd. A. Jahnke
**Name,
Andreas Jahnke
MDR Responsible Person**