



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 frames with sun protection / 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

Complies with European standard EN ISO 12870

Risk category **Category I**
Conformity assessment procedure **internal production control (module A) set out in annex IV**



and conforms with the following regulations:

Regulation (EU) 2016/425 on personal protective equipment

Complies with European standard EN ISO 12312-1

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

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

Nuremberg, 06.05.2024

Place and date of issue

sgd. P. Braunhofer

Name,
Dr. Peter Braunhofer
CEO

sgd. A. Jahnke

Name,
Andreas Jahnke
MDR Responsible Person