



## EU Declaration of conformity

# MI.OPTICS

Name of manufacturer **MI.OPTICS s.r.o.**  
**Jindřichova 308**  
**344 01 Domažlice**  
**Czech Republic**

Single Registration Number (SRN) **CZ-MF-000012845**

We declare under sole responsibility that the product:

Name of product **Spectacle frames FFX**

Trade Name **CUBISTA**

Model **8335**

Colors **T01,T02,T03,T04,T05,T06**

Basic UDI-DI **8596510FFX002E**

Nomenclature **GMDN Code 32816, EMDN Code Q02100203**

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 of the European Parliament and of the Council (EU) 2017/745 on medical devices
- Complies with European standard EN ISO 12870
- 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.

Domažlice,  
06.12.2022

Place and Date of issue

Pavel Linder  
MDR responsible person

Jiri Kaiser  
CEO MI.OPTICS s.r.o.