



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 | Prague |
| Model | 8452 |
| Colors | 001, 002, 003, 004 |
| 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,
24.04.2024

Place and Date of issue

Pavel Linder
MDR responsible person

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