EU Declaration of conformity

MIOPTICS

| 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 | 8170 |
| Colors | P01,P02,P03 |
| 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, 16.04.2024

Place and Date of issue

Pavel Linder MDR responsible person

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