

EU Declaration of conformity



Name of the manufacturer:

**Fielmann Group AG
Fuhlsbüttler Straße 399
22309 Hamburg**

Single Registration Number (SRN):

DE-MF-000006245

We declare under our sole responsibility, pursuant to article 52(7) MDR, the conformity with Regulation (EU) 2017/745 of the European Parliament and of the Council for the following product:

Basic UDI DI according to Annex VI MDR:

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Product name:

BD 605 FA ELLA II

Risk class:

1 (according to rules set out in Annex VIII MDR)

Intended use:

**Spectacle frames to be combined with
corrective lenses to treat refractive errors**

The above mentioned product is a medical device according to article 2(1) of the regulation. It fulfils the basic safety and performance requirements according to Annex I of the regulation and complies with the state of the art and applicable harmonized standards.

Hamburg, 09.09.2025

Place, date

Valter Larentis
Qualified Person (Art. 15 MDR)

