



Name of the manufacturer:

**Fielmann Group AG
Weidestraße 118a
22083 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:

4066209metfuladu6Z

Product name:

BD 587 FA EMILIA

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, 09.01.2025

Place, date

Valter Larentis
Qualified Person (Art. 15 MDR)

fielmann
Fielmann Group AG
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