Fielmann

Zentrale

CE

Name of the manufacturer:	**Fielmann 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:	**4066209plafuladu38**
Product name:	**BD 548 FLEX FA**
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 acording 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, 04.07.2023

Place, date

Valter Larentis
Qualified Person (Art. 15 MDR)

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Fielmann AG
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