Fielmann

Zentrale

EU Declaration of conformity

CE

Name of the manufacturer:	**Fielmann Group AG** **Fuhlsbüttler Straße 399** **22309 Hamburg**
Single Registration Number (SRN):	**DE-MF-000006245**

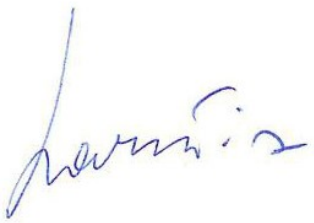
We declare under our sole responsibility, pursuant to article 52(7) MDR, the conformity with Regulation (EU) 2017/745 of the European Parliament and of the Council for the following product:

Basic UDI DI according to Annex VI MDR:	**4066209plafuladu38**
Product name:	**MC 642 FA PHILIPP**
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, 10.02.2026

Place, date



Valter Larentis
Qualified Person (Art. 15 MDR)