



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:

**4066209mixfuladuBH**

Product name:

**LN 062 FA LINDA**

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, 14.05.2024

Place, date

Valter Larentis  
Qualified Person (Art. 15 MDR)

**fielmann**  
Fielmann Group AG  
Weidestraße 118a  
22083 Hamburg