

# fielmann

Zentrale



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:

**4066209plafuladu38**

Product name:

**EMT 008 RN**

Risk class:

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

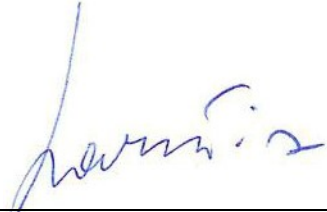
Intended use:

**Spectacles intended for near-vision and reading use only,  
having or incorporating a pair of non-prescription single-  
vision lenses of equal positive spherical power**

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, 16.02.2024

Place, date

  
Valter Larentis  
Qualified Person (Art. 15 MDR)

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