

EU Declaration of Conformity

Manufacturers Name: COMO

Manufacturers Address: 2 Route de Lyon – Molinges

39360 CHASSAL MOLINGES

Authorized Representative Name: M. DELACROIX Jean-Luc

Authorized Representative Address : 2 Route de Lyon – Molinges

39360 CHASSAL MOLINGES

Basic UDI-DI:

For full rim metal model : : 358012METALCERCLED5

For full rim acetate models : 358012ACETATECERCLEG2

For full rim combo models : 358012COMBOCERCLE9N

For Nylon metal model : 358012METALNYLON3D

For Nylon acetate model : 358012ACETATENYLONVP

For Nylon combo model : 358012COMBONYLONWZ

Name of the Device (s): Eyeware Metal and Plastic Frames

Risk Class and Classification Rule: Class I Rule 1

Conformity assessment route: COMO uses the following procedures for the CE-labeling of their products according the Regulation MDR 2017/745: EC conformity declaration according to Annex IV and Annex VIII

This declaration of conformity is issued under the sole responsibility of COMO We hereby declare that he medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. All supporting documentation is retained at the premises of the manufacturer.

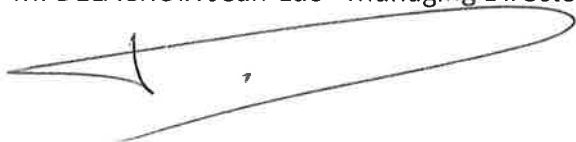
Place and date of issue:

27 June 2023

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Signature

M. DELACROIX Jean-Luc - Managing Director



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