KENMARK EYEWEAR

Declaration of Conformity

MANUFACTURER		
Name of Company	**Address**	**SRN**
Kenmark Eyewear	1800 Research Dr Louisville, KY 40299 U.S.A.	US-MF-000040231

AUTHORIZED REPRESENTATIVE		
Name of Company	**Address**	**SRN**
MedEnvoy Global B.V.	Prinses Margrietplantsoen 33 Suite 123 2595 AM The Hague The Netherlands Phone: +31 70 3262148 Email: EU-AR@medenvoyglobal.com	NL-AR-000024028

EMDN code	Description
Q02100203	Metal and plastic spectacle frames without lenses

GMDN code	Description
35065	Prescription spectacles

PRODUCT IDENTIFICATION	
Basic-UDI-DI	**Name of Device(s) / Model Name**
0715317900311TC	Plastic / Acetate frames
0715317900319TU	Metal / other material frames

Intended Purpose
Frames and non-corrective lenses intended for cosmetic enhancements per the user preference.

RISK CLASS FOR MEDICAL DEVICES	
Device Classification	
Risk Classification	MDR Class I
Risk Classification Rule	Rule 1
Product List	See related product list, PL-001 (Product List Spectacle Frames, Rev. A)

KENMARK EYEWEAR

This Declaration of Conformity is issued under the sole responsibility of Kenmark Eyewear. We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices.

COMPANY REPRESENTATIVE:

NAME: Susan Parker

TITLE: VP of Supply Chain

PLACE: Louisville, KY USA

DATE: August 26, 2024

SIGNATURE: DocuSigned by: Susan Parker D111300A71CF4CC...

Revision	Description of Change
1.0	Initial release under EU 2017/745.
1.1	Correct order postal code MedEnvoy Global B.V.