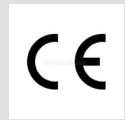


EU DECLARATION OF CONFORMITY

REGULATION (EU) 2017/745 ACCORDING TO ANNEX IV OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL.
of 5 April 2017 on MEDICAL DEVICES,



MANUFACTURER	IFLEX EYEWEAR COMPANY LIMITED Rm 904 Hilder Centre, 2 Sung Ping Street Hung Hom, Kowloon, Hong Kong
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SRN	HK-MF-000052203
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AUTHORIZED EUROPEAN REPRESENTATIVE	ICARE EUROPE SARL CHEMIN DU MOULIN CARRON, 38, ECULLY, France
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SRN	FR-AR-000010357
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THE MANUFACTURER DECLARES UNDER THEIR RESPONSIBILITY THAT THE PRODUCTS DESCRIBED BELOW MEET THE REQUIREMENTS OF THE REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL OF 5 APRIL 2017 ON MEDICAL DEVICES,

PRODUCT DESCRIPTION
FRAMES FOR OPHTHALMIC GLASSES
BRANDS
REFORM EYEWEAR

In all their variants (colours and size)

INTENDED USE:

The frame is an essential accessory so that the ophthalmic lenses can be carried and compensate for a visual defect.

Classification	CLASS I, Rule 1, Annex VIII, Chapter III, Regulation (EU) 2017/745
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PRODUCT FAMILY	BASIC UDI-DI
PLASTIC FULL RIM	489715328FFPRZ
METAL FULL RIM	489715328FFMRT
MIXED FULL RIM	489715328FFXSH
PLASTIC HALF RIM	489715328FHMRZ
METAL HALF RIM	489715328FHPS7
MIXED HALF RIM	489715328FHXP
<ul style="list-style-type: none"> Conformity assessment procedure Name and ID of the notified body Certificate CE number 	Annex II, Annex III, and Annex IV N.A N.A
<i>The manufacturing quality Management System follows the standards</i>	EN ISO 9001:2015, ISO 13485:2016

Applied standard & common Specification: EN ISO 14971 / EN ISO 15223-1/ EN 1041 / EN ISO 10993-1 +AC:2010 / EN 62366 / EN ISO 12870:2016/

Signed by: Cedric Bimar

Function: Managing Director

Date: 09/03/2026

Place of Issue and SRN: Hung Hom, Kowloon, Hong Kong - HK-MF-000052203

Signature: