Johnson&Johnson vision

DECLARATION OF CONFORMITY

Manufacturer:

AMO Ireland Block B Liffey Valley Office Campus Quarryvale, Co. Dublin Ireland SRN: IE-MF-000013704 Manufacturer Production Facility: AMO (Hangzhou) Co., Ltd. 200, 4th Avenue Hangzhou Economic & Technological Development Zone, 310018 Hangzhou, Zhejiang PEOPLE'S REPUBLIC OF CHINA

Devices Concerned		
Catalog Number/Model Number	Formulation 7723XA	Device Classification
	Product Name	
Applicable SKU: 90059GMH, 90068RHH, 90276GMH	Lens Plus Ocupure	Class IIb Rule 15

We, AMO Ireland., declare exclusively under sole responsibility that the device(s) listed above meet the provisions of Annex II of Council Directive 93/42/EEC of 14 June 1993 concerning medical devices.

Standards Applied:

Applied standards are listed in the Essential Requirements Checklist RTF7723XA-5040

Notified Body:	TÜV SÜD Product Service GmbH Ridlerstraße 65 80339 Müchen Germany	
Notified Body Identification Number:	CE0123	
EC Certificate Number:	G1 001630 0011	
Start of CE Marking	2021-03-12	
PH3074 REV. 2 Declaration of Confo	Declaration of Conformity – AMO Ireland (Formulation 7723XA)	

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(Date, Lot, or serial number):

Note: This Declaration of Conformity to European Medical Devices Directive 93/42/EEC has been re-issued and signed for administrative purposes.

We confirm that these products were listed on a valid CE Declaration of Conformity prior to the Date of Application (26 May 2021) of Regulation (EU) 2017/745 and may continue to be placed on the market in compliance with Article 120 Transitional Provisions of the Regulation.

Signature of Regulatory Representative:

Name and Title:

Nicole Kassner, Associate Director Regulatory Affairs, Johnson & Johnson Vision



Signature

Date

Signature of Quality Representative:

Name and Title:

Vincent Jordan, CQ Director EMEA for Johnson & Johnson Vision

Dublin, Ireland



Digitally signed by Vincent Jordan Adobe Reader version: 11.0.23

Signature

Date