



## 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

<i>Devices Concerned</i>		
<b>Catalog Number/Model Number</b>	<b>Formulation 7723XA Product Name</b>	<b>Device Classification</b>
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  
München Germany

Notified Body Identification Number: CE0123

EC Certificate Number: G1 001630 0011

Start of CE Marking 2021-03-12

(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

**Nicole Kassner**

Digitally signed by Nicole Kassner  
DN: cn=US, o=JNJ, ou=Subscribers, cn=Nicole Kassner,  
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Signature

Date

Signature of Quality Representative:

Name and Title:

Vincent Jordan, CQ Director EMEA for Johnson & Johnson Vision

Dublin, Ireland

**Vincent Jordan**

Digitally signed by Vincent Jordan  
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email=vjordan2@its.jnj.com, c=IE  
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Date