



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>		
Catalog Number/Model Number	Formulation 9554X Product Name	Device Classification
Applicable SKU: 90153FPH, 94199RR, 94200GM, 94377CQ, 94377FL, 94377GM, 94377RR 90152CQH, 90152GMH, 90152FLH, 90152RRH, 90152FPH	Blink TotalCare Solution	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 RTF9554-5020

Notified Body: 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 Certificate 2021-03-12

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

Signature

Date