

EC Declaration of Conformity**According to Directive 93/42/EEC as amended by 2007/47/EC**

Legal Manufacturer Bausch & Lomb, Incorporated
1400 N. Goodman St.
Rochester, NY 14609 USA

European Authorized Representative(s)* Bausch & Lomb GmbH
Brunsbütteler Damm 165-173
13581 Berlin, Germany

Notified Body TUV Rheinland
LGA Products GmbH
Tillystraße 2
90431 Nürnberg
Notified Body Number: 0197

EC Certificate Number EC Cert: HD 60146676 0001

Product (s) Independent Viewing Chamber or
Independent Corneal Viewing Chamber

Product Code(s) See Attached Table on Page 2

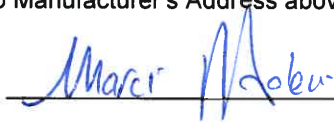
GMDN Code 45358

Classification IIa, Rule 2 according to Directive 93/42/EEC Annex IX

We hereby declare the conformity of the above mentioned products with the European Medical Device Directive 93/42/EEC as amended by 2007/47/EC Annex II, Section 3. Above product(s) is/are developed and manufactured in compliance with the MDD and the applicable European harmonized standards.

Place of Issue: Refer to Manufacturer's Address above

Signature:



Date: 13 October 2020

Name/Title/Position: Marci Halevi, Director, Regulatory Affairs

* The previous EU Authorized Rep address may appear on product manufactured prior to 29-Mar-2019.
Bausch & Lomb, Incorporated
106 London Road
Kingston-upon-Thames
KT2 6TN UK

EC DECLARATION OF CONFORMITY
According to Directive 93/42/EEC as amended by 2007/47/EC

Legal Manufacturer	Bausch & Lomb, Incorporated 1400 N. Goodman St. Rochester, NY 14609 USA
Product (s)	Independent Viewing Chamber or Independent Corneal Viewing Chamber

Item Number (SKU)	Product Name
IVC-12	Independent Corneal Viewing Chamber