

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

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 60105024 0001

Product (s) OcuCoat Viscoelastic

Product Code(s) See Attached Table on Page 2

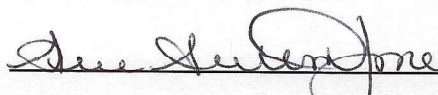
GMDN Code 35907

Classification IIb, Rule 6, 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: 10/23/2018

Name/Title/Position: Sue Sutton-Jones, Associate Director Regulatory Affairs,
Surgical Equipment & Devices

* 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

Manufacturer	Bausch & Lomb, Incorporated 1400 N. Goodman St. Rochester, NY 14609 USA
Product (s)	OcuCoat Viscoelastic

Item Number (SKU)	Product Name
CC050S	OcuCoat 1 mL single pack
CC065S	OcuCoat 1 mL six pack
CC100SL	OcuCoat 2 mL single pack