BAUSCH + LOMB

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EC Declaration of Conformity

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

Legal Manufacturer	Bausch & Lomb Incorporated 1400 North Goodman Street Rochester, NY 14609 U.S.A.
European Authorized Representative(s)*	Bausch & Lomb Incorporated Cork Road Industrial Estate Waterford, X91 V383, Ireland
Notified Body	National Standards Authority of Ireland (NSAI) 1 Swift Square Northwood, Santry Dublin 9, Ireland Notified Body number: 0050
EC Certificate Number	252.124
Product (s)	Multipurpose Contact Lens Care Solutions and Lens Care Kits
GMDN Code	45870, 45088
Classification	Class IIb, Rule 15, 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 Legal Manufacturer's Address above

Signature: Nancy A. Fehrman Date: Sept. 2, 2020

Name/Title/Position: Nancy Fehrman, Senior Manager, Regulatory Affairs

*The previous EU Authorized Rep address may appear on product manufactured prior to 29- March-2019.
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