

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	252.860
Product (s)	Soft Corrective Contact Lens, Daily-Disposable (nesofilcon A)
GMDN Code	47841
Classification	Class IIa, Rule 5, 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: April 30, 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.

Bausch & Lomb, Incorporated
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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 North Goodman Street Rochester, NY 14609 U.S.A.
Product (s)	Soft Corrective Contact Lens, Daily-Disposable (nesofilcon A)

Product Name
BAUSCH + LOMB Biotrue ONEday (nesofilcon A) Contact Lenses
BAUSCH + LOMB Biotrue ONEday (nesofilcon A) For Presbyopia Contact Lenses
BAUSCH + LOMB Biotrue ONEday (nesofilcon A) For Astigmatism Contact Lenses
See Attachment 1 for Private Label Names