Documents provided by us based on your request, present the sole property of Bausch Health/BAUSCH+LOMB, or the respective manufacturer and can be used solely for the designated purposes in accordance with by the Regulation

(EU) 2017 A 1 5 Che Europart at liament and of the Council of 5 April 2017 approximation and a samended.

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EU Declaration of Conformity

In accordance with Annex IV of Regulation (EU) 2017/745 of the European Parliament and of the Council

Technical Documentation Number	745-002
Legel Manufacturar	Bausch & Lomb Incorporated 1400 North Goodman St. Rochester, NY 14609 USA SRN: US-MF-000001029
European Authorized Representative	Bausch & Lomb Incorporated Cork Road Industrial Estate Waterford, X91 V383, Ireland SRN: IE-AR-000000094
Notified Body	Nalional Standards Authority of Ireland (NSAI) 1 Switt Square Northwood, Santry Dublin 9, Ireland Nobilied Body Number: 0050
Producta	Bausch + Lomb ULTRA® ONE DAY (kalillicon A) Copiaci Lenses
Product Code(s)	KAPBSDD
Baeic UDI-DI	0310119B0020YV
Globaj Medical Devices Nomenclatura Code	47841 – Solt corrective contact lons, dally- disposable
EMDN Code	Q021004010101
Classification	Class IIa (Annex VIII, Rule 5)
Conformity Assessment Probedure	Regulation (EU) 2017/745 Annex IX Conformity essessment based on a quality management system and on assessment of technical documentation
Intended Purppse	Bausch + Lomb kalifilicon A Contact Lens is Indicated for the deily wear correction of retractive ametropia (myopia and hyperopia) in aphakic and/or non-aphakic persons with non- diseased eyes that exhibit refractive astigmatism of 2,00 (diopters or less, that does not interfere with visual acuity. The lens is to be prescribed for single-use disposable wear and is to be discarded efter each removal.

Documents provided by us based on your request, present the sole property of Bausch Health/BAUSCH+LOMB, or the respective manufacturer and can be used solely for the designated purposes in accordance with by the Regulation (EU) 2017 245 S the European Parliament and of the Council of 5 April 2017 on madical devices as amended.

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Hochesler, NY 14609 USA 565 338,8000

This EU Declaration of Conformity is issued under the sole responsibility of the Legal Manufacturer.

We, Bausch & Lomb Incorporated, hereby declare the above listed Medical Devices comply with Medical Device Regulation (EU) 2017/745. Above products are developed and manufactured in compliance with the EU MDR and the applicable European harmonized standards and Common Specifications as applicable.

This declaration is made on the basia of EU Quality System Cartificate Number MD19.1854 and EU MDR EC Cartificate Number 745.002, issued by the Notified Body stated above, in accordance with Annex (X, Chapters) and ii) of Medical Device Regulation (EU) 2017/745 and the products above comply with Annex L.

Place of Issue; Refer to Legal Menulecturer's Address above.

Signature:

Name/Title/Position:

Date: