

DECLARATION OF CONFORMITY

We Luxottica Group SpA, with registered office in Piazzale Cadorna 3 – 20123 Milan (MI)-Italy, declare that the models and the relative spare parts, of which we are the manufacturer:

BRAND	TRADEMARK
A0	ALAIN MIKLI
AN	ARNETTE
AR	GIORGIO ARMANI
AX	ARMANI EXCHANGE
BV	BVLGARI
BE	BURBERRY
CH	CHANEL
DK/DY	DONNA KARAN / DONNA KARAN NEW YORK
DG/DD	DOLCE & GABBANA / D&G
EA	EMPORIO ARMANI
HC	COACH
LU	LUXOTTICA
MK	MICHAEL KORS
MU	MIU MIU
MT	MOSLEY TRIBES
OO / OX/ OY/ OJ	OAKLEY
OV	OLIVER PEOPLES
PO	PERSOL
PH	POLO RALPH LAUREN
PM	PAUL SMITH
PR/PS	PRADA / PRADA LINEA ROSSA
RA/RL/PL	RALPH LAUREN
RB/RJ/RX/RY	RAY-BAN
SF	SFEROFLEX
SH	S+ARCK EYES
SM	STELLA MCCARTNEY
TF	TIFFANY & CO.
TY	TORY BURCH
VA	VALENTINO
VE / VR	VERSACE / VERSUS
VO	VOGUE

- all ophthalmic frames comply with Annex VII of the Medical Device Directive 93/42/EEC amended by the Directive 2007/47/EC according to the current reference standard EN ISO 12870 ophthalmic optic spectacle frames general requirement and test methods, risk classification: class I according to the Annex IX;
- all sunglasses are in conformity with the EU Regulation of the European Parliament and of the Council 2016/425 and the harmonized standard EN ISO 12312-1:2013/A1:2015



DIRECTIVES

EU Regulation of the European Parliament and of the Council 2016/425

"Personal protective equipment"

EEC Directive 94/27 CEE "Nickel Directive"

EEC Directive 95/2001 CEE "General Product Safety"

EEC Directive 93/42 CEE "Medical Devices"

CE Regulation 1907/2006 "REACH"

US FDA Reg. 21C.F.R 801.410 "Drop Ball Test" for Sunglasses and sun lenses

<u>US FDA Reg.21 C.F.R. subchapter H</u> – medical devices, part 801 – labelling

US Treasury Decision 74-38 (January 22, 1974; 39 F.R. 2470)

US Consumer Product Safety Act

US Consumer Product Safety Improvement Act of 2008

California Proposition 65 "The Safe Drinking Water and Toxic Enforcement Act"

STANDARDS

EN ISO 12312-1:2013/A1:2015 "Eye and face protection-Sunglasses and related eyewear.

EN ISO 12311:2013 "Personal Protective Equipment – Test methods for sunglasses and related eyewear"

ISO 12870:2016 "Ophthalmic optics - Spectacle frames - Requirements and test methods"

EN 12472:2005+A1:2009 "Method for the simulation of wear and corrosion for the detection of nickel release from coated items"

EN 16128:2015 "Reference method for the testing of spectacle frames and sunglasses for nickel release"

ISO 8624:2011 "Measuring system for spectacles frames"

ANSI Z80.3:2018 "American National Standard. Non prescription Sunglasses and Fashion Eyewear - Requirements".

AS/NZS 1067.1:2016 "Eye and face protection - Sunglasses and fashion spectacles" QB2457:1999, "Sunglasses"

<u>GB10810.3:2006</u> "Spectacle lenses and related eye wear – Part3: Transmittance specifications and test methods"

NOTES

The authorization and registration Number by the Health Ministry, as Medical Devices producer of the First category is:

473617 (frames without lenses - others)

473615 (metal and plastic frames without lenses)

473613 (plastic frames without lenses)

473553 (metal frames without lenses)

The F.D.A. Registration Number is: 9614827; FDA listing number A974287 (frames), A974283 (sunglasses), B018782 (lenses)

This conformity declaration is issued under exclusive responsibility of the Manufacturer. Done at Agordo, on 30 July 2019

Luxottica Group S.p.A.

Carlo Salomoni

Group Customer Service Director

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Iscr. Registro Imprese di Milano n. 00891030272