LUXOTTICA

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*
KL / KX	*KILLER LOOP*
LC / LU / LX	*LUXOTTICA*
MK	*MICHAEL KORS*
MU	*MIU MIU*
MT	*MOSLEY TRIBES*
OO / OX	*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*
SH	*S+ARCK EYES*
SM	*STELLA MCCARTNEY*
TF	*TIFFANY & CO.*
TY	*TORY BURCH*
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 disposition of the EEC statements n. 89/686/EEC according to the current reference standard EN ISO 12312-1

- This conformity declaration is issued under exclusive responsibility of the Manufacturer.

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Codice Fiscale 00891030272 - Partita IVA 10182640150
Capitale Sociale € 28.653.640,38 i.v.
Iscr. R.E.A. Milano n. 1348098
Iscr. Registro Imprese di Milano n. 00891030272

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DIRECTIVES

EEC Directive 89/686 CEE "Personal eye-protection Directive"
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
US FDA Reg.21 C.F.R. subchapter H – 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 "Eye and face protection-Sunglasses and related eyewear.
ISO 12870:2012 "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 16128:2011 "Reference test method for release of nickel from those parts of spectacle frames and sunglasses intended to come into close and prolonged contact with the skin"
ISO 8624:2011 "Measuring system for spectacles frames"
ANSI Z80.3:2010 "American National Standard. Nonprescription Sunglasses and Fashion Eyewear - Requirements".
AS/NZS 1067:2003 "Australian/New Zealand Standard TM. Sunglasses and fashion spectacles"
QB2457:1999, GB10810.1 "Sunglasses","Spectacle Lenses"
GB13511 "Assembled spectacles"
CNS 15067:2012 "Sunglasses and sunglare filters for general use and filter for direct observation of the sun"

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 (lenses)

Done at Agordo, on 26 Feb 2016

Luxottica Group S.p.A.
Group Quality Director
Carlo Salomoni

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