



EU DECLARATION OF CONFORMITY		CE
REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL. of 5 April 2017 on MEDICAL DEVICES		
MANUFACTURER (Name / Brand or commercial name of the manufacturer/complete legal address)	QUICK WONDER GLASSES WENZHOU CO.,LTD WEST OF 11TH FLOOR, NO 1 HUAIJIANG ROAD, LUCHENG DISTRICT,WENZHOU, CHINA	
SRN (MANUFACTURER)	CN-MF-000012339	
AUTHORIZED EUROPEAN REPRESENTATIVE	Luxus Lubenswelt GmbH Kochstr.1,47877, Willich,Germany	
COMPETEN AUTHORITY Bezirksksregieung Dusseldorf, Dezernat 24	Cecilienallee 2, Dusseldorf,Nordrhein-Westfalen,Germany Dez24.mpg@brd.nrw.de	
SRN(AUTHORIZED EUROPEAN REPRESENTATIVE)	DE-AR-000005110	
<p>DECLARE UNDER THEIR RESPONSIBILITY AS MANUFACTURER THAT THE PRODUCTS DESCRIBES BELOW MEET THE REQUIREMENTS OF THE REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL OF 5 APRIL 2017 ON MEDICAL DEVICES</p> <p>PRODUCT DESCRIPTION</p> <p>CORRECTIVE: SPECTACLE FRAMES</p> <p>BRAND NAME:  </p> <p>INTENDED USE: An ophthalmic device made of metal, acetate, or plastic in order to include optical lenses manufactured in accordance with a standard prescription and used to correct the refractive errors of a patient's eyesight, e.g., refractive ametropia (myopia, hyperopia, and astigmatism), and possibly to protect the eyes against radiation or mechanical hazards.</p> <p>UDI-DI BASIC: 697471200KKAQ8,697471200KMQY,697471200KKCQC,697471200RSAS3,697471200RSMST,697471200RSCS7</p> <p>Classification (CLASS): class I (anexo VIII regla 1)</p> <p>Specifics of the product: OPTICAL</p> <p>The products:</p> <p>-Comply with European standards EN ISO 12870: 2016 / EN 16128 (if applicable)</p> <p>-Are in conformity with the EU REGULATION 2017-745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices;</p> <p>-Are marked "CE";</p> <p>-Are in conformity with the general obligation of security requested by 2001/95/CEE Directive and all its updates.</p>		
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Signed by:

Function: Vice president

Date and place: 18/02/2022 Wenzhou China

