DECLARATION OF CONFORMITY TO COUNCIL DIRECTIVE 93/42/EEC CONCERNING MEDICAL DEVICES

MANUFACTURER:	Lianyungang TianNuo Optical Instrument Co.,Ltd No.3 JinQiao Road, Dapu Industrial park,222000 Lianyungang, Jiangsu, PEOPLE'S REPUBLIC OF CHINA
MEDICAL DEVICE:	Trial lens set
SPECIFICATION:	232
CLASSIFICATION - ANNEX IX:	Class I With Measuring, rule 1
CONFORMITY ASSESSMENT ROUTE:	MDD 93/42/EEC Annex V

WE, THE MANUFACTURER, HERE WITH DECLARE THAT THE STATED MEDICAL DEVICE MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC CONCERNING MEDICAL DEVICES;
ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER, AND THE MANUFACTURER IS EXCLUSIVELY RESPONSIBLE FOR THE DECLARATION OF CONFORMITY.

NOTIFIED BODY:	**TÜV SÜD PRODUCT SERVICE GMBH RIDLERSTR. 65 - 80339 M NCHEN, GERMANY**
IDENTIFICATION NUMBER	CE **0123**
(EC) CERTIFICATE(S):	**G2M 069 181 0007 REV.01**
EXP. DATE:	**<2024-03-03>**
EC REP ***EUROPEAN REPRESENTATIVE:***	**Shanghai International Holding Corp. GmbH (Europe) Eiffestrasse,80, D-20537, Hamburg, Germany Tel: 0086-021-65951371，0049-40-2513175，Fax: 0049-40-255726**

This declaration of conformity is only valid before 21st, March, 2020.

START OF CE-MARKING: **<2009-03-04>**

PLACE, DATE OF DECLARATION: Lianyungang, Jiangsu, China ***, 2019-03-18***

SIGNATURE:

NAME: MR. QIN LIANGPING
POSITION:
GENERAL MANAGER