DECLARATION OF CONFORMITY TO COUNCIL DIRECTIVE 93/42/EEC OF 14 JUNE 1993 CONCERNING MEDICAL DEVICES

MANUFACTURER:	**Ray Vision International Corporation** Rm. 103 Building C,11 Dewai Street, Xicheng District, Beijing 100088, China
MEDICAL DEVICE:	Digital Visual Acuity Chart Model: Elite
CLASSIFICATION - ANNEX IX:	CLASS I MEASURING, RULE 1
CONFORMITY ASSESSMENT ROUTE:	ANNEX VII+V

WE, THE MANUFACTURER, HEREWITH DECLARE THAT THE STATED MEDICAL DEVICES MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC OF 14 JUNE 1993 CONCERNING MEDICAL DEVICES;
INCLUDING, AT 21 MARCH 2010, THE AMENDMENTS BY COUNCIL DIRECTIVE 2007/47/EEC.
ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

STANDARDS APPLIED: EMC DIRECTIVE: 2004/108/EC ANNEX II, LOW VOLTAGE DIRECTIVE: 2006/95/EC

NOTIFIED BODY:	TUV SUD PRODUCT SERVICE GMBH RIDLERSTR 65,0-80339 MUNCHEN, GERMANY
IDENTIFICATION NUMBER	CE 0123
(EC) CERTIFICATE(S):	G2M090268272911
EC REP EUROPEAN REPRESENTATIVE:	Medizintechnik De candia, Henschelring 13, D - 85551 Kirchheim b. München, Germany
START OF CE-MARKING:	2009-03-27
PLACE, DATE OF DECLARATION:	Beijing, China, 2012-08-07
SIGNATURE:	Ligo Wang Product Certification Manager