






DECLARATION OF CONFORMITY

Manufacturer's name NIDEK Co. Ltd.
 Manufacturer's address 34-14 Maehama, Hiroishi-cho, Gamagori, Aichi 443-0038, Japan
NIDEK s.a.
 European Representative Europarc, 13 rue Auguste Perret, 94042 Créteil, France
 Identification of device AUTO LENSMETER
 Model No. LM-7/LM-7P
 Classification(Annex IX, MDD) I
 Category (Annex I, RoHS) 8
 Classification(2014/53/EU, RE) 1

We herewith declare under our sole responsibility that the above mentioned products meet the provisions of the following EC Council Directives and Standards. All supporting documentation is retained under the premises of the manufacturer. The supporting technical documentation is also retained at NIDEK s.a., Europarc, 13 rue Auguste Perret, 94042 Créteil, France.

General applicable directives	Date CE Mark was affixed
COUNCIL DIRECTIVE 93/42/EEC of concerning medical devices	August 4, 2017 
COUNCIL DIRECTIVE 2011/65/EU concerning restriction of the use of certain hazardous substances	August 4, 2017 
Directive 2014/53/EU of the European Parliament and of the council on the market of radio equipment	August 4, 2017 

Place: Aichi, Japan
 Signed by

Effective date: August 4, 2017


 Hiroyuki Torii
 Director
 Top Management
 NIDEK Co., Ltd.

Date of signature: August 3, 2017