



# DECLARATION OF CONFORMITY

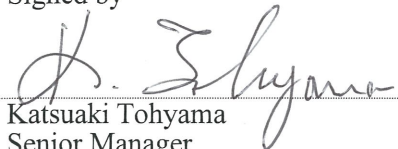
Manufacturer's name NIDEK Co., Ltd. SRN N/A  
 Manufacturer's address 34-14 Maehama, Hiroishi-cho, Gamagori, Aichi 443-0038, Japan  
 European Representative name NIDEK S.A. SRN FR-AR-000000341  
 European Representative address Ecoparc, rue Benjamin Franklin, 94370 Sucy En Brie, FRANCE  
 Identification of device AUTO LENS METER  
 Model No. LM-1800P / LM-1800PD  
 Classification (Annex VIII, MDR) I (Rule 13)  
 Category (for RoHS) 8  
 Basic UDI-DI 4987669106G7

We herewith declare under our sole responsibility that the above mentioned products meet the provisions of the following EC Council Directive(s)/Regulation. All supporting documentation is retained under the premises of the manufacturer. The supporting technical documentation is also retained at NIDEK S.A., Ecoparc, rue Benjamin Franklin, 94370 Sucy En Brie, FRANCE.

General applicable directive(s)/regulation	Date CE Marking was affixed
REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices	February 20, 2020 
COUNCIL DIRECTIVE 2011/65/EU concerning restriction of the use of certain hazardous substances.	April 21, 2014 

Place: Aichi, Japan

Signed by



Katsuaki Tohyama  
 Senior Manager  
 Person responsible for regulatory compliance  
 NIDEK Co., Ltd.

Effective date : September 30, 2021

Date of signature : August 31, 2021