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Document No.DOCCP-9EU02

DECLARATION OF CONFORMITY

Manufacturer's name	NIDEK Co., Ltd.	SRN	NA	
Manufacturer's address	34-14 Maehama, Hi	roishi-cho, Gamagori,	Aichi 443-0038,	Japan
European Representative name	NIDEK S.A.	SRN	FR-AR-00000	00341
European Representative address	Ecoparc, rue Benjan	nin Franklin, 94370 Su	cy En Brie, FRA	NCE
Identification of device	CHART PROJECTO	OR		
Model No.	CP-9			
Classification (Annex VIII, MDR)	I (Rule 13)			
Category (for RoHS)	8			
Basic UDI-DI	4987669105	G5		

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	April 3, 2020
COUNCIL of 5 April 2017 on medical devices	CE
COUNCIL DIRECTIVE 2011/65/EU concerning restriction of the use of certain hazardous substances.	April 3, 2020

Place: Aichi, Japan

Effective date: August 30, 2021

August 26, 202/

Signed by

Katsuaki Tohyama / Senior Manager

Person responsible for regulatory

compliance NIDEK Co., Ltd. Date of signature: