EU DECLARATION OF CONFORMITY

This declaration of conformity is issued under the sole responsibility of the manufacturer MEDOP.

PPE Identification	Type	Eye protection –Prescription spectacles			
	References	DANUBIO (DANU MARR L/ DANU MARR M / DANU MARR XL)			
	Batch				
Manufacturer	MEDOP Bruno Mauricio Zabala street, 16-4º izq – 48003 Bilbao – Vizcaya - Spain www.medop.es				
Object of the declaration	Spectacles for industrial use fixed with prescription lenses (glass and CR39)				
Relevant Union harmonisation legislation	Regulation (EU) 2016/425 Regulation (EU) 2017/475 RD 1591/2009				
Relevant harmonised standards	EN 166:2001				
PPE category	II				
Medical Device Class	I				
Notified Body that performed the EU type-examination (Module B)	SGS Fimko OY, (No 0598) Takomotie 8, Fl-00380 Helsinki, Finland				
Certificate reference	FI21/968043				
Conformity Assessment Procedure	Not Apply X	Module C2 ☐	Module D ☐		

Bilbao, on 25th May 2021

MEDOP, S.A. APTDO 1660 - 48080 BILBAO

Inés Gómez-Rubiera

Legal Representative