EU Declaration of Conformity as per MDR 2017/745

Manufacturer:	PRODUITS DENTAIRES SA Rue des Bosquets 18 1800 Vevey Switzerland
CHRN	CHRN-MF-20000511
SRN	CH-MF-000023147

European Authorized Representative	PD Dental EU Thonon-les-Bains, 74200 France
European Authorized Representative SRN	FR-AR-000017144

We declare under our sole responsibility that the medical device of the following Generic Device Group:

Plastic Pipettes

Reference:	*See attached table*
Risk class:	*I*
As per rule:	*5 Annex VIII, MDR 2017/745*
Intended Purpose:	*Device used for delivering small quantities of liquid into the root canal or pulp chamber during endodontic or restorative treatment*
Trade name:	*Plastic Pipettes*
Basic UDI-DI:	*764015916PIPETTESB4*

Meets all the requirements of the medical device regulation ***MDR 2017/745***

Conformity assessment procedure:	*MDR 2017/745 Article 52, Annex II and III and ODim July 2020.*	
Quality & Regulatory Affairs Engineer	Silvia Contini	Vevey, 2022-04-22
President	*Nicolas Gehrig*	Vevey, 2022-04-22

Plastic Pipettes

Reference	*Designation*
11750	PLASTIC PIPETTES 0.3mL ASSORTED (5)
11752	PLASTIC PIPETTES 1mL ASSORTED (5)
11754	PLASTIC PIPETTES 0.3mL ASSORTED (25)
11757	PLASTIC PIPETTES 1mL ASSORTED (25)