ivoclar vivadent®
passion vision innovation

CE 0123

Declaration of conformity

Manufacturer Address	Ivoclar Vivadent AG Bendererstrasse 2 9494 Schaan Liechtenstein
Product	**SR Vivodent® S PE / SR Orthotyp® S PE** **SR Vivodent® S DCL / SR Orthotyp® S DCL** **SR Ortholingual® S DCL**
Type of material	Resin Teeth
Product category	Resin Teeth
Classification	Medical Device Class IIa

We hereby declare under our exclusive responsibility that the above mentioned products meet the provisions of the following EC Council Directives and its implementation in national law. All supporting documentation is retained on the premises of the manufacturer and the notified body.

Directives	Medical Device Directive 93/42/EEC, Annex II. 3
Notified Body Address	TÜV SÜD Product Service GmbH Ridlerstrasse 65 80339 München Deutschland
Place, Valid from	Schaan, 2018-12-11 Replaces version of: 2017-03-06
Valid until	2023-12-10

Signature		
Name Position	Dr. Thomas Hirt CTO	Dipl. Ing. Patrik Oehri Director CQM and Regulatory Sicherheitsbeauftragter (Product Safety Officer)
Date	2018-12-11	2018-12-11