ivoclar vivadent
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CE 0123

Declaration of conformity

Manufacturer Address	Ivoclar Vivadent AG Bendererstrasse 2 FL – 9494 Schaan Liechtenstein
Product	**ExciTE® F**
Type of material	Dental adhesive with fluoride release
Product category	Adhesives
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, 2014-12-01 Replaces version of: 2014-06-23
Valid until	2019-11-27

Signature		
Name Position	Dr. Thomas Hirt CTO	Dipl. Ing. Patrik Oehri Director CQM and Regulatory Sicherheitsbeauftragter (Product Safety Officer)
Date	2014-12-01	2014-12-01

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Attachment to Declaration of conformity

Product	**ExciTE® F**
Dated	2014-12-01

613724	ExciTE F VivaPen Refill 1x2 ml
626969	ExciTE F VivaPen Refill 3x2 ml
627177	ExciTE F VivaPen Intro Pack 1x2 ml
630375	ExciTE F Refill 1x5 g
630376	ExciTE F Refill 2x5 g
630377	ExciTE F Single Dose Refill 50
688370	ExciTE F-Adhese Univ Promo VivaPen
688371	ExciTE F-Adhese Univ Promo Bottle

Place, Valid from	Schaan, 2017-01-26 Replaces version of: 2014-12-01	
Signature		
Name Position	Dr. Erik Braziulis Scientific Service, R&D	Sandra Cakebread Regulatory Affairs, R&D
Date	2017-01-26	2017-01-26

This Declaration of Conformity is handled without the suffix "AN", "EN", "AL", "WW" etc after the article number. The suffix is only for in-house reference and relates to the language version (e.g. English, Spanish, French, etc) and/or electrical plug of the product. The suffix does not belong to the official article number.