ivoclar vivadent
passion vision innovation

Declaration of conformity

CE 0123

Manufacturer Address	Ivoclar Vivadent AG Bendererstrasse 2 FL – 9494 Schaan Liechtenstein
Product	**IPS Empress® Direct**
Type of material	Restorative material
Classification	Medical Device Class IIa

We herewith declare that the above mentioned products meet the provisions of the following EC Council Directives and its implementation in national law. All supporting documentations are retained under the premises of the manufacturer and the notified body.

Directives	Medical Device Directive 93/42/EEC, Annex II. 3
Standards	EN ISO 4049 - Polymer-based filling, restorative and luting materials
Notified Body Address	TÜV SÜD Product Services GmbH Ridlerstrasse 65 DE – 80339 München Deutschland
Place, Valid from	Schaan, 22 April 2009 Replaces version of: ---

Signature		
Name Position	Dr. Volker Rheinberger CTO	Dipl. Ing. Patrik Oehri Director R&D Services
Date	21 April 2009	21 April 2009