ivoclar vivadent®

Declaration of conformity

CE 0123

Manufacturer Address	Ivoclar Vivadent AG Bendererstrasse 2 FL – 9494 Schaan Liechtenstein
Product	**IPS e.max® Press**
Type of material	Ceramic system
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	ISO 6872 Dental ceramic ISO 9693 Metal-ceramic dental restorative systems
Notified Body Address	TÜV Product Services Ridlerstrasse 65 DE – 80339 München Deutschland
Place, Valid from	Schaan, 08 May 2007 Replaces version of: 03 October 2005

Signature		
Name Position	Dr. Volker Rheinberger CTO	Dipl. Ing. Patrik Oehri Director R&D Services
Date	07 May 2007	07 May 2007