**IPS Empress CAD** 



Date of issue / Reference 04.04.2007 lise / Version 2
Replaces version of 07.08.2006 casa

Date of printing 04.04.2007 **Sheet No. 1662** Page 1 of 5

Company Ivoclar Vivadent AG, Bendererstrasse 2, FL - 9494 Schaan Principality of Liechtenstein

### 1 Commercial product name and supplier

1.1	Commercial product name /	IPS Empress CAD
	Designation	•

1.2 Application / Use Ceramic

1.3 Producer Ivoclar Vivadent AG, Bendererstrasse 2, FL - 9494 Schaan

Fürstentum Liechtenstein

1.4 Supplier

1.5 TOX emergency number Emergency-Call: +423 / 235 35 35 or 373 40 40

Ivoclar Vivadent AG, FL-9494 Schaan, Liechtenstein

### 2 Composition

2.1 Chemical characterization Ceramic ingots made of:

> 98 % SiO2, BaO, Al2O3, CaO, CeO2, Na2O, K2O, B2O3

< 2 % TiO2 and pigments

2.2 Hazardous components None.

2.3 Further information None.

### **3 Hazards identification** Grinding dust (see 8.3.1).

### 4 First aid measures In case of contact with grinding dust

4.1 Eye contact Flush eyes with plenty of water; mechanical effects only.

4.2 Skin contact No specific requirements.

4.3 Ingestion No specific requirements.

4.4 Inhalation Take into fresh air.

4.5 Further information None.

### 5 Fire-fighting measures

5.1 Suitable extinguishing media No specific requirements.

not combustible

5.2 Extinguishing media to avoid None.

5.3 Further information None.

9.7 Viscosity

IPS Empress CAD



Date of issue / Reference 04.04.2007 lise / Version 2

Replaces version of 07.08.2006 casa

Date of printing		04.04.2007	Sheet No. 1662	Page 2 of 5		
6	Accidental release measures	Clean up mec	hanically.			
7	Handling and storage					
7.1	Handling	Only adequate	ely trained personnel should	handle this product.		
7.2	Industrial hygiene	Usual hygieni	Usual hygienic measures for dental practice.			
7.3	Storage	No specific re	No specific requirements.			
7.4	Place of storage	No specific re	No specific requirements.			
7.5	Fire- and explosion-protection	None.				
8	Exposure controls / Personal prot	tection				
8.1	Technical measures	Provide adequ	ate local ventilation.			
8.2	Control of threshold limits	Producer Indu	stry recommends an exposur	re limit of 10 mg/m3.		
8.3	Personal protective equipment					
8.3.1	Respiratory protection		rotection. Avoid inhalation o spheres, use an approved dus			
8.3.2	Hand protection	Not required.				
8.3.3	Eye protection	Safety goggle	S.			
8.3.4	Other	None.				
9	Physical and chemical properties					
9.1	Appearance	Ingot				
9.2	Colour	off-white				
9.3	Odour	odourless				
9.4	Change of physical state			Test method:		
9.5	Density	not known				
9.6	Vapour pressure	HOL KHOWH				

not applicable

not applicable

**IPS Empress CAD** 



Date of issue / Reference 04.04.2007 lise / Version 2 07.08.2006 Replaces version of casa Date of printing 04.04.2007 Page 3 of 5 **Sheet No. 1662** 9.8 Solubility Solubility in water non soluble 9.9 pH not applicable 9.10 Flash point not applicable 9.11 Ignition temperature not applicable 9.12 Explosion limits Lower: Upper: not applicable 9.13 Further information None. 10 Stability and reactivity 10.1 Thermal decomposition None. 10.2 Hazardous decomposition products None. 10.3 Hazardous reactions None. 10.4 Further information None. 11 **Toxicological information** 11.1 Acute toxicity This product is not hazardous according to EEC criteria. No adverse effects anticipated by this route of exposure incidental to 11.2 Subacute / Chronic toxicity proper industrial handling. 11.3 Further information None. 12 **Ecological information** No ecological problems to be anticipated if properly handled and used. non soluble 13 Take to an approved landfill or a waste incineration plant, under **Disposal considerations** conditions approved by the local authority. 13.1 EU waste key 08 02 99

#### 14 **Transport information**

**RID** 14.1 Transport at land **ADR** 

15.9 BVD classification (CH)

15.11 Further information

15.10 VbF (D)

IPS Empress CAD



Date of issue / Reference 04.04.2007 lise / Version 2
Replaces version of 07.08.2006 casa

Date of printing 04.04.2007 **Sheet No. 1662** 

Page 4 of 5

		UN Number		Kemler Number	
		Packing Group			
		Proper shipping name	e		
14.2	Transport at sea	ADNR		IMDG	
		UN Number			
		EMS		MFAG	
		Packing Group			
		Proper shipping name	e		
14.3	Air transport	ICAO / IATA-DGR			
		UN Number			
		Proper shipping name	e		
		Subsidiary Risk			
		Labels			
		Packing Group			
	Passenger airplane	Packing Instructions			
		max.			
	Cargo Airplane	Packing Instructions			
		max.			
14.4	Further information	max. Product is not classifi		gerous good for tra	nsport.
14.4	Further information  Regulatory information		ed as a dan		
		Product is not classifi The product is a med 93/42/EEC. This product is classi	ed as a dan	according to the EC	C-directive US and
		Product is not classification.  The product is a med 93/42/EEC.  This product is classication.  Canadian regulations.	ed as a dan ical device fied as a me	according to the EC edical device under sen reviewed by the	C-directive US and
		Product is not classifi The product is a med 93/42/EEC. This product is classi	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
		The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
15	Regulatory information	The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
<b>15</b> 15.1	Regulatory information  UN number	The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
15.1 15.2	Regulatory information  UN number  National regulations	The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
15.1 15.2 15.3	Regulatory information  UN number  National regulations  EU number	The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
15.1 15.2 15.3 15.4	Regulatory information  UN number National regulations  EU number Hazard symbols	The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
15.1 15.2 15.3	Regulatory information  UN number  National regulations  EU number	The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
15.1 15.2 15.3 15.4	Regulatory information  UN number National regulations  EU number Hazard symbols	The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and
15.1 15.2 15.3 15.4 15.5	Regulatory information  UN number National regulations  EU number Hazard symbols Hazard designation	The product is a med 93/42/EEC. This product is classi Canadian regulations Drug Administration	ed as a dan ical device fied as a me and has be and Health	according to the EC edical device under een reviewed by the a Canada.	C-directive US and US Food and

None.

**IPS Empress CAD** 



Date of issue / Reference 04.04.2007 lise / Version 2 07.08.2006 Replaces version of

casa

04.04.2007 Date of printing **Sheet No. 1662** Page 5 of 5

16 Other information Version: 2

Changes: 1/6/7/8

The above mentioned data correspond to our present state of knowledge and experience. The safety data sheet serves as description of the products in regard to necessary safety measures. The indications do not have the meaning of guarantees on properties.

This safety data sheet has been generated with the safety database 'ChemManager', 
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