

EU Safety Data Sheet

IPS Empress CAD Multi



Date of issue / Reference

04.04.2007

lise / Verison 2

Replaces version of

07.08.2006

casa

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04.04.2007

Sheet No. 1663

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 /
Designation

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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.

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Sheet No. 1663

Page 2 of 5

6 Accidental release measures

Clean up mechanically.

7 Handling and storage

7.1 Handling

Only adequately trained personnel should handle this product.

7.2 Industrial hygiene

Usual hygienic measures for dental practice.

7.3 Storage

No specific requirements.

7.4 Place of storage

No specific requirements.

7.5 Fire- and explosion-protection

None.

8 Exposure controls / Personal protection

8.1 Technical measures

Provide adequate local ventilation.

8.2 Control of threshold limits

Producer Industry recommends an exposure limit of 10 mg/m³.

8.3 Personal protective equipment

8.3.1 Respiratory protection

Respiratory Protection. Avoid inhalation of dust while grinding.
In dusty atmospheres, use an approved dust respirator.

8.3.2 Hand protection

Not required.

8.3.3 Eye protection

Safety goggles.

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

not applicable

9.7 Viscosity

not applicable

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IPS Empress CAD Multi



Date of issue / Reference	04.04.2007	lise / Verison 2
Replaces version of	07.08.2006	casa
Date of printing	04.04.2007	Sheet No. 1663

Page 3 of 5

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.

11.2 Subacute / Chronic toxicity

No adverse effects anticipated by this route of exposure incidental to 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 Disposal considerations

Take to an approved landfill or a waste incineration plant, under conditions approved by the local authority.

13.1 EU waste key

08 02 99

14 Transport information

14.1 Transport at land

ADR

RID

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casa

Date of printing

04.04.2007

Sheet No. 1663

Page 4 of 5

		UN Number	---	Kemler Number	---
		Packing Group	---		
		Proper shipping name	---		
14.2	Transport at sea	ADNR	---	IMDG	---
		UN Number	---		
		EMS	---	MFAG	---
		Packing Group	---		
		Proper shipping name	---		
14.3	Air transport	ICAO / IATA-DGR	---		
		UN Number	---		
		Proper shipping name	---		
		Subsidiary Risk	---		
		Labels	---		
		Packing Group	---		
	Passenger airplane	Packing Instructions	---		
		max.	---		
	Cargo Airplane	Packing Instructions	---		
		max.	---		
14.4	Further information	Product is not classified as a dangerous good for transport.			

15 Regulatory information

The product is a medical device according to the EC-directive 93/42/EEC.

This product is classified as a medical device under US and Canadian regulations and has been reviewed by the US Food and Drug Administration and Health Canada.

This product does not require classification as Dangerous Goods.

15.1 UN number

15.2 National regulations

15.3 EU number

15.4 Hazard symbols

15.5 Hazard designation

15.6 Risk phrases

15.7 Safety phrases

15.8 MAK value

15.9 BVD classification (CH)

15.10 VbF (D)

15.11 Further information

None.

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Replaces version of	07.08.2006	casa
Date of printing	04.04.2007	Sheet No. 1663

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',
© ASSiST Applied Software Solutions in Science and Technology AG, Weiherweg 3, CH-4104 Oberwil, Switzerland

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