Telio Lab LC Transpa Incisal and Base



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Company Ivoclar Vivadent AG, Bendererstrasse 2, FL - 9494 Schaan

20.07.2010

Fürstentum Liechtenstein

Telio Lab LC

1 Commercial product name and supplier

1.1 Commercial product name /

Transpa Incisal and Base

1.2 Application / Use

Designation

Light-curing characterization material

1.3 Producer

Ivoclar Vivadent AG, Bendererstrasse 2, FL - 9494 Schaan

Fürstentum Liechtenstein msds@ivoclarvivadent.com

Supplier

1.4 TOX emergency number

Official Emergency-Call: +423 / 235 33 13

Ivoclar Vivadent AG, FL-9494 Schaan, Liechtenstein

2 Hazards identification

Uncured material: Direct contact can cause eye and skin irritation.

May cause sensitization by skin contact.

3 Composition

3.1 Chemical characterization

Paste of dimethacrylates, copolymer, silicondioxide, catalysts,

stabilizers and pigments

3.2 Hazardous components

CAS No. 72869-86-4 < 30 % Urethane dimethacrylate

EINECS/ELINCS No.: 276-957-5

Xi: Irritant. R36: Irritating to eyes. R38: Irritating to skin. R43: May cause

sensitisation by skin contact.

CAS No. 6701-13-9 < 10 % Decamethylendimethacrylate

EINECS/ELINCS No.: 229-745-1

Xi: Irritant. R43: May cause sensitisation by skin contact.

3.3 Further information None.

4 First aid measures

4.1 Eye contact Flush with plenty of water. Consult a physician if irritation persists.

4.2 Skin contact Wash thoroughly with water.

4.3 Ingestion No hazards anticipated from swallowing small amounts incidentally

to normal handling.

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4.4	Inhalation	Remove to fresh air.		
4.5	Further information	None.		
5	Fire-fighting measures			
5.1	Suitable extinguishing media	Water fog, carbon dioxide, foam, dry chemicals.		
5.2	Extinguishing media to avoid	None known		
5.3	Further information	None.		
6	Accidental release measures	Clean up mechanically. Dispose of according to local and national regulations.		
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	Store at 2-28 °C / 36-82 °F		
7.4	Place of storage	Avoid exposure to light.		
7.5	Fire- and explosion-protection	Not required.		
8	Exposure controls / Personal protection			
Q 1	Exposure controls	Good general ventilation should be sufficient		

8.1 Exposure controls Good general ventilation should be sufficient.

8.2 Exposure limit values None established.

8.3 Occupational exposure controls

8.3.1 Respiratory protection Not required.

8.3.2 Hand protection

Check penetration time for the specific gloves used with the glove

manufacturer.

Commercial medical gloves do not provide protection against the

sensitizing effect of methacrylates.

8.3.3 Eye protection Safety goggles.

8.3.4 Other None.

Environmental exposure controls Not relevant. 8.4

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9	Physical and chemical properties		
9.1	Appearance	Paste	
9.2	Colour	off-white to cream	
9.3	Odour	practically odourless	
9.4	Change of physical state	Test method:	
9.5	Density		
7.5	Delisity	Not determined.	
9.6	Vapour pressure		
		not applicable	
9.7	Viscosity		
		not determined	
9.8	Solubility		
	Solubility in water	< 0.1 %	
9.9	pH		
	•	Not determined.	
9.10 Flash point			
		not applicable	
9.11	1 Ignition temperature		
		not determined	
9.12	Explosion limits	Lower: Upper:	
		not applicable	
9.13	Further information		
	Part. coeff. n-octanol/water		
	Evaporat. rate		
		None.	
10	Stability and reactivity		
10.1	Thermal decomposition	None, if used in accordance to instructions.	
10.2	Hazardous decomposition products	None under normal conditions of storage and use.	
10.3	Conditions / materials to avoid	None.	
10.4	Further information	Avoid exposure of product to light.	

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11	Toxicological information			
11.1	Acute toxicity	Oral LD50 for rats: > 5000 mg/kg		
11.2	Subacute / Chronic toxicity	Uncured material: prolonged or frequently repeated skin contact may cause allergic skin reactions.		
11.3	Further information	No hazards anticipated from swallowing small amounts incidentally to normal handling.		
12	Ecological information			
12.1	Ecotoxicity	No data available.		
12.2	Mobility	No data available.		
12.3	Persistence and degradability	No data available.		
12.4	Bioaccumulative potential	No data available.		
12.5	Further information	nearly insoluble No ecological problems to be anticipated if properly handled and used.		
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	20 01 39		
14	Transport information			
14.1	Transport at land	ADR RID		
		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 Marine pollutant		

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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 does not require classification according to the criteria	

of the EC.

15.1 UN number

15.2 National regulations

15.3 EINECS/ELINCS number

15.4 Hazard symbols Xi

15.5 Hazard designation

15.6 Risk phrases R: 43-36/38

43 May cause sensitisation by skin contact.

36/38 Irritating to eyes and skin.

5.7 Safety phrases S: 24-28-37

Avoid contact with skin.

28 After contact with skin, wash immediately with plenty

of ... (to be specified by manufacturer).

Wear suitable gloves.

15.8 AGW value

15.9 BVD classification (CH)

15.10 VbF (D)

15.11 Further information None.

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16 Other information

No other information.

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.

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Regulation (EC) No 1907/2006 (REACH)