Material Safety Data Sheet

Telio Stains



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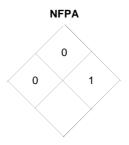
Company Ivoclar Vivadent AG, Bendererstrasse 2, FL - 9494 Schaan

Fürstentum Liechtenstein

1 Commercial product name and supplier

1.1 Commercial product name / Designation

Telio Stains



HMIS						
ПІЛІЭ						
Н	0					
F	0					
R	1					

1.2 Application / Use Light-curing characterization material

1.3 Producer Ivoclar Vivadent AG, Bendererstrasse 2, 9494 Schaan

Principality of Liechtenstein (FL)

1.4 Supplier Ivoclar Vivadent, Inc.

175 Pineview Drive, Amherst NY 14228, USA

2785 Skymark Ave., Unit 1 Mississagua, ON L4W4Y3, Canada MSDS prepared by Anderjeet Gulati. Tel. No. 716 691-0010

1.5 24 Hour Emergency Assistance Emergency-Call USA- Infotrac: 1-800-535-5053

Emergency-Call Canada - Canutec: 1-613-996-6666

General MSDS Assistance US: 1-800-533-6825 Canada: 1-800-263-8182

2 Composition

2.1 Chemical characterization Paste of dimethacrylates, inorganic fillers, initiators, stabilizers and

pigments

2.2 Hazardous components

CAS No. 1565-94-2 < 38 % Bis-GMA

Xi: Irritant. R36: Irritating to eyes. R38: Irritating to skin.

CAS No. 109-16-0 < 18 % Triethylene glycoldimethacrylate

Xi: Irritant. R43: May cause sensitisation by skin contact. R36/37/38: Irritating to

eyes, respiratory system and skin.

CAS No. 72869-86-4 < 33 % Urethane dimethacrylate

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

sensitisation by skin contact.

2.3 Further information None.

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

May cause sensitization by skin contact.

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4	First aid measures	
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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.

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

Test method:

5.3 Flash point

5.4 Ignition temperature

not determined

5.5 Explosion limits Lower:

Upper:

not applicable

5.6 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

8.1 Exposure controls Good general ventilation should be sufficient.

8.2 Exposure limit values None established.

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8.3 Occupational exposure controls

8.3.1 Respiratory protection Not required.

8.3.2 Hand protection Gloves.

Commercial medical gloves do not provide protection against the

sensitizing effect of methacrylates.

Check penetration time for the specific gloves used with the glove

manufacturer.

8.3.3 Eye protection Safety goggles.

8.3.4 Other None.

8.4 Environmental exposure controls Not relevant.

9 Physical and chemical properties

9.1 Appearance Paste

9.2 Colour various

9.3 Odour practically odourless

9.4 Change of physical state Test method:

9.5 Density

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 Further information

Part. coeff. n-octanol/water

Evaporat. rate

None.

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Marine pollutant

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14.3	Air transport	ICAO / IATA-	DGR			
r	•	UN Number				
		Proper shippin	g name			
		Subsidiary Ris	k			
		Labels				
		Packing Group)			
	Passenger airplane	Packing Instru	ctions			
		max.				
	Cargo Airplane	Packing Instru	ctions			
		max.				
14.4	Further information					
		Product is not classified as a dangerous good for transport.				ransport.
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.				
15.1	National regulations					
15.2	NFPA Storage					
15.3	Further information	None.				
16	Other information	No other infor	mation			
as de	above mentioned data correspond to our scription of the products in regard to new antees on properties.		_	_		

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