EU Safety Data Sheet		SR Ivocap Clear Poly	ivoclar				
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Company		Ivoclar Vivadent AG, Bendererstrasse 2, FL - 9494 Schaan Fürstentum Liechtenstein					
1	Commercial product name and sup	plier					
1.1	Commercial product name / Designation	SR Ivocap Clear Polymer					
1.2	Application / Use	Denture base 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	Dust conception					
		Dust generation					
3	Composition						
3.1	Chemical characterization	> 98 % Polymethylmethacrylate (C	CAS-No. 9011-1	4-7)			
3.2	Hazardous components						
	CAS No. 94-36-0	< 1.5 % Benzoylperoxide EINECS/ELINCS No.: 202-327-6 Xi: Irritant. E: Explosive. R3: Extreme risk other sources of ignition. R7: May cause fi sensitisation by skin contact.					
3.3	Further information	None.					
4	First aid measures						
4.1	Eye contact	Flush with plenty of water. Consul	t a physician if i	rritation persists.			
4.2	Skin contact	Wash thoroughly with soap and water.					
4.3	Ingestion	Give large amounts of water.					
4.4	Inhalation	Take into fresh air.					
4.5	Further information	If respiratory irritation is experienced, call a physician.					

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5	Fire-fighting measures						
5.1	Suitable extinguishing media	Extinguishing powder. Water fog. Carbon dioxide.					
5.2	Extinguishing media to avoid	Do not use di	rect water stream.				
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	Avoid dust build-up. Only adequately trained personnel should handle this product.					
7.2	Industrial hygiene	Avoid breathing dust. Usual hygienic measures for dental practice.					
7.3	Storage	Store at 12-28 °C / 54-82 °F.					
7.4	Place of storage	Avoid exposure to direct sunlight.					
7.5	Fire- and explosion-protection	Dust in air may be explosive.					
8	Exposure controls / Personal protect	tion					
8.1	Exposure controls	Provide adequ	uate local ventilation.				
8.2	Exposure limit values	Producer Industry recommends an exposure limit of 10 mg/m3.					
8.3	Occupational exposure controls						
8.3.1	Respiratory protection	Avoid breathing dust. In dusty atmospheres, use an approved dust respirator.					
8.3.2	Hand protection	Gloves.					
8.3.3	Eye protection	Safety goggles.					
8.3.4	Other	Use only with adequate ventilation.					
8.4	Environmental exposure controls	ntrols					

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9	Physical and chemical properties				
9.1	Appearance	powder			
9.2	Colour	white			
9.3	Odour	low			
9.4	Change of physical state				Test method:
9.5	Density	1.2 g/cm ³ (20°C))		
9.6	Vapour pressure	C ()			
		not applicable			
9.7	Viscosity				
		not applicable			
9.8	Solubility				
	Solubility in water				
		non soluble			
9.9	pH				
		Not applicable.			
9.10	Flash point	>250 °C			
9.11	Ignition temperature	>400 °C			
9.12	Explosion limits	Lower:			
		Upper: not determined			
0.13	Further information	not determined			
9.13	Part. coeff. n-octanol/water				
	Evaporat. rate				
	-	None.			
10	Stability and reactivity				
		> 250 °C			
10.1	Thermal decomposition	> 250 °C			
10.2	Hazardous decomposition products	Product may de	accordance to inst compose at elevate irritating (Methylm	ed temperatures	s generating vapours
10.3	Conditions / materials to avoid	none known			
10.4	Further information	None.			

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11	Toxicological information						
11.1	Acute toxicity		thacrylate: The oral ide: The oral LD50				
11.2	Subacute / Chronic toxicity	No adverse effects anticipated by this route of exposure incidental to proper industrial handling. In rare cases can cause an allergic reaction.					
11.3	Further information	Dust may irritate eyes. Cured material: Biological evaluation according to OECD principles and EEC directives no cytotoxic nor allergic potential, no skin irritation.					
12	Ecological information						
12.1	Ecotoxicity	No data availa	ıble.				
12.2	Mobility	No data availa	ıble.				
12.3	Persistence and degradability	No data availa	ıble.				
12.4	Bioaccumulative potential	No data availa	ıble.				
12.5	Further information	No ecological problems to be anticipated if properly handled and used. nearly insoluble					
13	Disposal considerations	Burn in an adequate incinerator under carefully controlled conditions in accordance with local and national regulations.					
13.1	EU waste key	20 01 39					
14	Transport information						
14.1	Transport at land	ADR		RID			
		UN Number Packing Grou Proper shippir		Kemler Numbe	er		

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14.2	Transport at sea	ADNR			IMDG	
		UN Number EMS Packing Group Proper shippin Marine polluta	ig name		MFAG	
14.3	Air transport	ICAO / IATA- UN Number Proper shippin Subsidiary Ris Labels Packing Group	ig name k	 		
	Passenger airplane	Packing Instru	ctions			
	Cargo Airplane	max. Packing Instru max.	ctions			
14.4	Further information	Product is not	classified as	s a dang	erous 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 regulations and has been reviewed by the US Food and drug Administration. This product does not require classification as Dangerous Goods.				
15.1	UN number					
15.2	National regulations					
15.3 15.4 15.5	EINECS/ELINCS number Hazard symbols Hazard designation					
15.6 15.7	Risk phrases Safety phrases					
15.8	AGW value	ml/m³ (ppn	1)			
15.9	BVD classification (CH)					
	VbF (D) 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)