



Tegoglas® OL80

Cold-end coating of container and table glassware

Tegoglas[®] OL80 is a cold-end coating product based on active fatty acids for coating of hollow glassware. It is used in conjunction with special vaporizing equipment. When using Tegoglas[®] OL80, the formation of cracked residues in the vaporizing equipment is minimized.

PRODUCT DATA

Basis	Mixture of fatty acids		
Effective material content	100%		
Specific weight	0.89 g/cm³ (20°C)		
Melting point	Below 20°C		
Form	Liquid (20°C)		
Colour	Yellowish		
Odour	Mild		

PROPERTIES

- 🗕 Tegoglas® OL80 is non-toxic.
- Tegoglas® OL80 influences the contents of the glassware neither in taste nor in smell provided that it is applied correctly.
- Surfaces coated with Tegoglas® OL80 show a high surface gloss and a good surface smoothness when applied properly.

PRODUCT SAFETY

The raw materials used for the manufacture of Tegoglas® are printed in the European Positive list of additives authorized for plastic materials and articles intended to come into contact with foodstuffs.

APPLICATION

Tegoglas[®] should be applied according to the instructions of the vaporizing equipment manufacturer.

APPLICATION CONCENTRATION

Tegoglas® is used undiluted.

CONSUMPTION

About 1 – 2kg of Tegoglas[®] OL80 is used per day per line, depending on the line operating conditions.

STORAGE

Tegoglas[®] OL80 must be stored in closed containers at room temperature (about 20°C). If it is stored below 20°C, a partial crystallization may occur. This has no impact on the quality of the product if the crystallized parts are melted again prior to use by gently warming up the product. If stored at room temperature in an unopened container, the shelf life is 3 years.

PACKAGING

Packaging	Height	Depth	Wid t h	Weight	Weight
	mm	mm	mm	gross kg	net kg
60 l non-returnable PE-Drums	635	390	340	53	50



SAFETY

Please refer to our health and safety data sheet for information on toxicity and ecology.

The information and advice contained in this data sheet is to the best of our knowledge and is provided without obligation.

We are not liable for any wrong advice or any advice we may have failed to give, unless otherwise agreed by the parties.

This data sheet becomes invalid as soon as a new edition has been published.

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It is the sole responsibility of the manufacturer of the medical device to determine the suitability (including biocompatibility) of all raw materials, products and components, including any medical grade Arkema products, in order to ensure that the final end-use product is safe for its end use; performs or functions as intended; and complies with all applicable legal and regulatory requirements (FDA or other national drug agencies) It is the sole responsibility of the manufacturer of the medical device to conduct all necessary tests and inspections and to evaluate the medical device under actual end-use requirements and to adequately advise and warn purchasers, users, and/or learned intermediaries (such as physicians) of pertinent risks and fulfill any postmarket surveillance obligations.

Any decision regarding the appropriateness of a particular Arkema material in a particular medical device should be based on the judgment of the manufacturer, seller, the competent authority, and the treating physician.



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