Versaflex[™] CL2250

Thermoplastic Elastomer **Avient Corporation**

Technical Data

Product Description

Versaflex[™] CL2250 is an easy processing compound designed for use in injection molding applications where FDA compliance, clarity and enhanced heat resistance are required.

- Excellent Clarity
- Overmold Adhesion to Polypropylene
- Superior Colorability
- Very Good Heat and Boil Resistance

General Material Status · Commercial: Active Technical Datasheet Literature¹ Avient Corporation Search for UL Yellow Card Versaflex™ • · Africa & Middle East · Latin America Availability Asia Pacific North America Good Colorability · Good Processability · High Clarity Features · Good Moldability · Good Processing Stability · Medical/Healthcare · Pacifiers · Transparent or Translucent Uses Applications · Personal Care Parts Overmolding • ISO 10993-5 • FDA 21 CFR 177.1210² Agency Ratings • ISO 10993-4 USP Class VI³ **RoHS** Compliance · RoHS Compliant Appearance · Clear/Transparent · Pellets Forms · Injection Molding **Processing Method**

Physical	Nominal Value Unit	Test Method
Density / Specific Gravity	0.888 g/cm ³	ASTM D792
Melt Mass-Flow Rate (MFR) (190°C/2.16 kg)	13 g/10 min	ASTM D1238
Molding Shrinkage - Flow	0.80 to 1.2 %	ASTM D955
Mechanical	Nominal Value Unit	Test Method
Flexural Modulus	13.9 MPa	ASTM D790
Elastomers	Nominal Value Unit	Test Method
Tensile Stress ^{5, 6}		ASTM D412
100% Strain, 23°C	1.52 MPa	
300% Strain, 23°C	2.32 MPa	
Tensile Strength ^{5, 6} (Break, 23°C)	5.70 MPa	ASTM D412
Tensile Elongation ^{5, 6} (Break, 23°C)	770 %	ASTM D412
Tear Strength	24.5 kN/m	ASTM D624
Compression Set (23°C, 22 hr)	20 %	ASTM D395B
Hardness	Nominal Value Unit	Test Method
Durometer Hardness (Shore A, 10 sec)	50	ASTM D2240
Fill Analysis	Nominal Value Unit	Test Method
Apparent Viscosity (200°C, 11200 sec^-1)	11.9 Pa·s	ASTM D3835



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PROSPECT

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Nominal Value Unit	
20 %	
171 to 188 °C	
193 to 221 °C	
193 to 227 °C	
210 to 227 °C	
210 to 221 °C	
13 to 38 °C	
0.00 to 0.552 MPa	
25 to 75 rpm	
	20 % 171 to 188 °C 193 to 221 °C 193 to 227 °C 210 to 227 °C 210 to 221 °C 13 to 38 °C 0.00 to 0.552 MPa

Color concentrates based on polypropylene (PP), ethylene vinyl acetate (EVA), or low density polyethylene (LDPE) are most suitable for coloring Versaflex[™] CL2250. Improved color dispersion can be achieved by using higher melt flow concentrates (with a melt flow from 25-40 g/10 min). Typical loadings for color concentrates are 1% to 5% by weight. Liquid color can be used, but mineral oil based carriers may have a significant effect on the final hardness value. Concentrates based on PVC should not be used. A high color match consistency can be obtained by the use of precolored compounds available from GLS. The final determination of color concentrate suitability should be determined by customer trials.

Purge thoroughly before and after use of this product with a low flow (0.5 - 2.5 MFR) polyethylene (PE) or polypropylene (PP).

Regrind levels up to 20% can be used with Versaflex[™] CL2250 with minimal property loss, provided that the regrind is free of contamination. To minimize losses during molding, the melt temperature should remain as low as possible. The final determination of regrind effectiveness should be determined by the customer.

Versaflex[™] CL2250 has excellent melt stability. Maximum residence times may vary, depending on the size of the barrel. Generally, the barrel should be emptied if it is idle for periods of 8 - 10 minutes or longer.

Drying is not Required

Injection Speed: 0.5 to 2 in/sec 1st Stage - Boost Pressure: 100 to 800 psi 2nd Stage - Hold Pressure: 30% of Boost Hold Time (Thick Part): 4 to 10 sec Hold Time (Thin Part): 1 to 3 sec

Notes

¹ These links provide you with access to supplier literature. We work hard to keep them up to date; however you may find the most current literature from the supplier.

² Please contact GLS Thermoplastic Elastomers for a copy of the FDA compliance letter.

- ³ Please contact PolyOne GLS Thermoplastic Elastomers for a complete copy of the GLS Healthcare Policy.
- 1. The Customer must notify GLS of any FDA Class I and/or European Union Class I medical devices for each specific product and application. 2. The Customer shall not knowingly manufacture, use, sell or otherwise supply, directly or indirectly products or compounds made from GLS
- products in any of the following without prior written approval by GLS for each specific product or application: a. Cosmetics

b. Drugs and other Pharmaceuticals

- c. Temporary or permanent implantation in the human body, regardless of the intended duration of implantation
- d. Class II and Class III Medical Devices as defined in 21 CFR 860.3 ("Medical Devices")
- e. Class IIa, IIb and III as defined in Directive 93/42/EEC

⁴ Typical properties: these are not to be construed as specifications.

⁵ Die C

⁶ 2 hr



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