

Technical Information

ELVAX™ 3165

Ethylene Vinyl Acetate Copolymer

Description					
Product Description	ELVAX™ 3165 is an extrudable ethylene-vinyl acetate copolymer resin available in pellet form for use in conventional extrusion equipment designed to process polyethylene resins.				
Restrictions					
Material Status	Commercial: Active				
Typical Characteristics					
Composition	18% By Weight Vinyl Acetate comonomer content Thermal Stabilizer: BHT antioxidant				
Applications	This resin is designed to provide a low temperature heat seal to itself or many other materials commonly used in flexible packaging applications. The melt properties of this resin allow it to be processed on blown film equipment over a wide range of film thickness and blow-up ratios. It can also be co-extruded with a variety of other polymers. This resin is typically used as low temperature seal layer in co-extruded films.				
Typical Properties					
Physical	Nominal Values		Test Method(s)		
*Density ()	0.94 g/cm ³	ASTM D792		ISO 1183	
*Melt Flow Rate (190°C/2.16kg)	0.7 g/10 min	ASTM D1238		ISO 1133	
Thermal	Nominal Values		Test Method(s)		
*Melting Point (DSC)	89°C (192.2°F)	ASTM D3418		ISO 3146	
Freezing Point (DSC)	68°C (154.4°F)	ASTM D3418			
Vicat Softening Point ()	69°C (156.2°F)	ASTM D1525		ISO 306	
Processing Information					
Maximum Processing Temperature	235 °C (455 °F)				
General Processing Information	Resin melt temperature should be maintained in the range of 160-185°C (320-365°F) to provide a suitable viscosity and melt strength for blown film extrusion. Higher temperatures may be more appropriate for coextrusion with other grades. Selection of a specific melt temperature will depend on considerations such as desired gauge, height of tower, cooling capacity, extruder hold up time, winding conditions, and other machine variables. ELVAX TM can be used in conventional extrusion equipment designed to process polyethylene resins. However, corrosion-protected barrels, screws, adapters, and				

FDA Status Information

products.

ELVAX™ 3165 resin complies with Food and Drug Administration Regulation 21 CFR 177.1350(a)(1) - - Ethylene-vinyl acetate copolymers, subject to the limitations and requirements therein. This Regulation describes polymers that may be used in contact with food, subject to the finished food-contact article meeting the extractive limitations under the intended conditions of use, as shown in paragraph (b)(1) of the Regulation.

ethylene vinyl acetate (EVA) resins may thermally degrade and release corrosive by-



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Regulatory Information

For information on regulatory compliance outside of the U.S.A., consult your local Dow representative.

Safety & Handling

THE IMPORTANCE OF PROPER HANDLING & STORAGE:

Maintaining proper handling and storage conditions for ELVAX™ resins is very important to ensure overall product quality and keep the resin in a free-flowing state. If the ELVAX™ resin is subjected to sunlight, rain or excessive temperatures, then the resin may not process properly or achieve the desired characteristics in the final product.

It is crucial for ELVAX™ resins to be kept under proper storage and handling conditions because improper storage and handling may cause the resin to "block" (massing of pellets into large clumps that can hinder the ease of material transfer) or lose the ability to flow freely.

Please refer to the ELVAX™ Handling Guide for additional information.

For additional information on appropriate Handling & Storage of this polymeric resin, please refer to the material Safety Data Sheet.

A Product Safety Bulletin, material Safety Data Sheet, and/or more detailed information on extrusion processing and/or compounding of this polymeric resin for specific applications are available from your Dow representative.

Product Stewardship

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b. use in cardiac prosthetic devices regardless of the length of time involved ("cardiac prosthetic devices" include, but are not limited to, pacemaker leads and devices, artificial hearts, heart valves, intra-aortic balloons and control systems, and ventricular bypass-assisted devices);

c. use as a critical component in medical devices that support or sustain human life; or

d. use specifically by pregnant women or in applications designed specifically to promote or interfere with human reproduction.

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