

Dow Packaging & Speciality Plastics Product Data Sheet

ELVALOY™ 742

Copolymer

Description				
Product Description	ELVALOY™ 742 is an ethylene/vinyl acetate/carbon monoxide (E/VA/CO) copolymer designed for polymer modification.			
Restrictions				
Material Status	Commercial: Active			
Other Restrictions	Dow recommends to consume the product within 18 months after date of production. See CoA (Certificate of Analysis) supplied with the resin to confirm date of manufacture. If no CoA is available, contact your Dow representative to determine the date of manufacture based on the production batch or lot number.			
Typical Characteristics				
Uses	Polymer Modifier			
Composition	By Weight			
Features	Permanent, non-migrating PVC modifier. CPVC, TPU modifier.			
Characteristics / Benefits	ELVALOY™ 742 copolymer deliver toughness and flexibility that is locked in to PVC. ELVALOY™ 742 copolymer will not migrate like liquid plasticizers.			
	Tensile StrengthElongation @ Break Duromete Hardness Molecular Weight Distribution -	1200% 55 Shore A	- ASTM D638 - ASTM D2240	
Applications	PVC Modification			
Typical Properties				
Physical	Nominal Values		Test Method(s)	
*Density ()	1.02 g/cm ³	ASTM D792		ISO 1183
*Melt Flow Rate (190°C/2.16kg)	35 g/10 min	ASTM D1238		ISO 1133
Thermal	Nominal Values		Test Method(s)	
*Melting Point (DSC)	45°C (113°F)	ASTM D3417		ISO 3146
*Glass Transition Temperature (DSC)	-32°C (-25.6°F)	ASTM D3418		
Processing Information				
*Maximum Processing Temperature	240 °C (464 °F)			

FDA Status Information

ELVALOY™ 742 resin complies with Food and Drug Administration Regulation 21 CFR 175.105 - - Adhesives. This Regulation describes adhesives that may be used as components of articles intended for use in packaging, transporting, or holding food, subject to the limitations and requirements therein.

The information and certifications provided herein are based on data we believe to be reliable, to the best of our knowledge. The information and certifications apply only to the specific material designated herein as sold by Dow and do not apply to use in any process or in combination with any other material. They are provided at the request of and without charge to our customers. Accordingly, Dow cannot guarantee or warrant such certifications or information and assumes no liability for their use.

Regulatory Information

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

Dow representative.

Safety & Handling

For 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

The Dow Chemical Company and its subsidiaries ("Dow") has a fundamental concern for all who make, distribute, and use its products, and for the environment in which we live. This concern is the basis for our Product Stewardship philosophy by which we assess the safety, health, and environmental information on our products and then take appropriate steps to protect employee and public health and our environment. The success of our Product Stewardship program rests with each and every individual involved with Dow products — from the initial concept and research, to manufacture, use, sale, disposal, and recycle of each product.

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Medical Applications Policy

NOTICE REGARDING MEDICAL APPLICATION RESTRICTIONS: Dow will not knowingly sell or sample any product or service ("Product") into any commercial or developmental application that is intended for:

- a. long-term or permanent contact with internal bodily fluids or tissues. "Long-term" is contact which exceeds 72 continuous hours (or for PELLETHANE™ Polyurethane Elastomers only, which exceeds 30 days);
- 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.

Dow requests that customers considering use of Dow products in medical applications notify Dow so that appropriate assessments may be conducted.

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Additional Information

North America Europe/Middle East Latin America

U.S. & Canada: 1-800-441-4369 **All Countries** +31-11567-2626 Argentina: +54-11-4319-0100 1-989-832-1426 +800-3694-6367 +55-11-5188-9000 Brazil: +57-1-219-6000 Mexico: +1-800-441-4369 Italy: +800-783-825 Colombia: Mexico: +52-55-5201-4700 South Africa +800-99-5078 Asia Pacific +800-7776-7776 +60-3-7958-5392

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