

Eastman TRITAN™

copolyester

Enhanced capability injection molding material

Eastman Tritan™ copolyester provides customers a clear polymer product with a balance of performance and processability. This balance helps Tritan create exciting opportunities to differentiate your products through lasting aesthetic appeal and value. Some advantages include excellent toughness, chemical resistance, higher temperature resistance, low and stable shrinkage rates. Mold release and nonmold release versions are available as well as colors and UV additives.

- Excellent chemical resistance
- Excellent toughness
- Ease of processing
 - Easy-dry resin
 - Improved cycle time
 - Same mold shrinkage value as polycarbonate
- Good optical properties
- · Higher temperature resistance
- Performs well in secondary operations

Basic properties		Eastman Tritan™ copolyester	
Property	ASTM method	MX710 and MX711	MX810 and MX811
Specific gravity	D792	1.18	1.17
Mold shrinkage, in./in. or mm/mm	D955	0.005-0.007	0.005-0.007
Tensile stress @ yield, psi (MPa)	D638	6,200 (43)	6,400 (44)
Tensile stress @ break, psi (MPa)	D638	7,700 (53)	7,700 (53)
Elongation @ tensile break, %	D638	210	140
Flexural modulus, psi (MPa)	D790	225,000 (1,550)	228,000 (1,585)
Rockwell hardness, R scale	D785	112	115
Notched Izod impact strength @ 23°C (73°F), ft-lb/in. (J/m)	D256	18.4 (980)	12.2 (650)
Notched Izod impact strength @ -40°C (-40°F), ft-lb/in. (J/m)	D256	2.1 (110)	2.4 (126)
Deflection temperature @ 66 psi (0.455 MPa), °C (°F)	D648	99 (210)	109 (228)
Deflection temperature @ 264 psi (1.82 MPa), °C (°F)	D648	85 (185)	92 (198)

Suggested drying and processing conditions

Drying conditions		
Drying temperature, °C (°F)	88 (190)	
Drying time, hr	4	
Dryer air dew point, °C (°F)	< -29 (< -20)	
Machine conditions		
Injection speed	slow	
Screw speed, rpm	30-60	
Pack and hold pressure, MPa	35–50	
Cushion, in.	0.2-0.4	
Back pressure, MPa	10–15	

Processing temperatures		
Zones, °C (°F)		
Rear	Set barrel temperatures to reach target melt temperature, up to 10°–20°C (20°–40°F) below targe depending on shear heating.	
Center		
Front		
Nozzle, °C (°F)	282 (540)	
Hot runners, °C (°F)	282 (540)	
Melt temperature, °C (°F)	282 ± 10 (540 ± 20)	
Mold temperature, °C (°F)	38°-66°C (100°-150°F)	



The results of insight

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Safety Data Sheets providing safety precautions that should be observed when handling and storing Eastman products are available online or by request. You should obtain and review the available material safety information before handling any of these products. If any materials mentioned are not Eastman products, appropriate industrial hygiene and other safety precautions recommended by their manufacturers should be observed.

It is the responsibility of the medical device manufacturer ("Manufacturer") to determine the suitability of all component parts and raw materials, including any Eastman product, used in its final product to ensure safety and compliance with requirements of the United States Food and Drug Administration (FDA) or other international regulatory agencies.

Eastman products have not been designed for nor are they promoted for end uses that would be categorized either by the United States FDA or by the International Standards Organization (ISO) as implant devices. Eastman products are not intended for use in the following applications: (1) in any bodily implant applications for greater than 30 days, based on FDA-Modified ISO-10993, Part 1, "Biological Evaluation of Medical Devices" tests (including any cosmetic, reconstructive, or reproductive implant applications); (2) in any cardiac prosthetic device application, regardless of the length of time involved, including, without limitation, pacemaker leads and devices, artificial hearts, heart valves, intra-aortic balloons and control systems, and ventricular bypass assisted devices; or (3) as any critical component in any medical device that supports or sustains human life.

For manufacturers of medical devices, biological evaluation of medical devices is performed to determine the potential toxicity resulting from contact of the component materials of the device with the body. The ranges of tests under FDA-Modified ISO-10993, Part 1, "Biological Evaluation of Medical Devices" include cytotoxicity, sensitization, irritation or intracutaneous reactivity, systemic toxicity (acute), subchronic toxicity (subacute), implantation, and hemocompatibility. For Eastman products offered for the medical market, limited testing information is available on request. The Manufacturer of the medical device is responsible for the biological evaluation of the finished medical device.

The suitability of an Eastman product in a given end-use environment is dependent on various conditions including, without limitation, chemical compatibility, temperature, part design, sterilization method, residual stresses, and external loads. It is the responsibility of the Manufacturer to evaluate its final product under actual end-use requirements and to adequately advise and warn purchasers and users thereof.

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