## Medical selector guide











Tritan™ copolyester	Suggested application	Specific gravity (ASTM D792)	Tensile yield		Tensile break (ASTM D638)		Flexural modulus	Izod impact, notched, J/m	Deflection temperature, °C (°F)		Optical properties (ASTM D1003)		Chemical	Contains certified
			Elongation, %	Stress, MPa (psi)	Elongation, %	Stress, MPa (psi)	(ASTM D790), MPa (105 psi)	(ff·lbf/in.) @ 23°C (73°F) (ASTM D256)	@ 1.82 MPa (264 psi)	@ 0.455 MPa (66 psi)	Transmittance,	Haze, %	resistance	recycled content*
MX710	Y sites		6	43 (6,200)	210	53 (7,700)	1,550 (2.3)	980 (18.4)	85 (185)	99 (210)	91	<1	Best	
MX711 (mold release)	IV components	1.18												
MX711 Renew 50 (mold release)	IV components	1.18	6	43 (6,200)	210	53 (7,700)	1,550 (2.3)	980 (18.4)	85 (185)	99 (210)	90	<1	Best	<b>✓</b>
MX731 (mold release)	Medical devices	1.18	7	43 (6,200)	210	52 (7,500)	1,570 (2.3)	860 (16.1)	81 (178)	94 (201)	91	<1	Better	
MX731 Renew 50 (mold release)	Medical devices	1.18	7	43 (6,200)	210	52 (7,500)	1,575 (2.28)	860 (16.1)	81 (178)	94 (201)	91	<1	Better	<b>✓</b>
MX810	Device housings	1.17	7	44 (6,400)	140	53 (7,700)	1,580 (2.3)	650 (12.2)	92 (198)	109 (228)	92	<1	Good	
MX811 (mold release)														

 $<sup>{}^{\</sup>star}Certified\ recycled\ content\ allocated\ using\ ISCC\ mass\ balance.$ 

## **EASTMAN**

**Eastman Corporate Headquarters** P.O. Box 431

Kingsport, TN 37662-5280 U.S.A.

U.S.A. and Canada, 800-EASTMAN (800-327-8626) Other locations, +(1) 423-229-2000

eastman.com/locations

Although the information and recommendations set forth herein are presented in good faith, Eastman Chemical Company ("Eastman") and its subsidiaries make no representations or warranties as to the completeness or accuracy thereof. You must make your own determination of its suitability and completeness for your own use, for the protection of the environment, and for the health and safety of your employees and purchasers of your products. Nothing contained herein is to be construed as a recommendation to use any product, process, equipment, or formulation in conflict with any patent, and we make no representations or warranties, express or implied, that the use thereof will not infringe any patent. NO REPRESENTATIONS OR WARRANTIES, EITHER EXPRESS OR IMPLIED, OF MERCHANTABILITY, FITNESS FOR A PARTICULAR PURPOSE, OR OF ANY OTHER NATURE ARE MADE HEREUNDER WITH RESPECT TO INFORMATION OR THE PRODUCT TO WHICH INFORMATION REFERS AND NOTHING HEREIN WAIVES ANY OF THE SELLER'S CONDITIONS OF SALE.

Safety Data Sheets providing safety precautions that should be observed when handling and storing our 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.

© 2024 Eastman. Eastman brands referenced herein are trademarks of Eastman or one of its subsidiaries or are being used under license. Non-Eastman brands referenced herein are trademarks of their respective owners.