

Property comparison and ranking guide

Eastman clear medical grade plastics



Property comparison and ranking guide—Eastman clear medical grade plastics

Physical comparison	ASTM test method	DuraStar™ polymers			Eastar™ copolyester		
		MN611	MN621	MN631	MN058	MN021	MN211
Physical properties							
Clarity							
Haze (%)	D1003	0.3	0.3	<1	<1	1	0.3
Transmittance (%)	D1003	91	91	92	82	84	91
Izod impact strength, notched @ 23°C (73°F), J/m (ft-lbf/in.)	D256	80 (1.5)	370 (7.0)	80 (1.5)	51 (1.0)	40 (0.8)	101 (1.9)
Flexural modulus, MPa (10 ⁵ psi)	D790	2000 (2.9)	1900 (2.8)	1900 (2.7)	2400 (3.5)	2500 (3.6)	2100 (3.0)
Elongation @ break (%)	D638	300	310	270	90	120	110
Tensile stress @ break, MPa (psi)	D638	51 (7400)	53 (7700)	43 (6300)	24 (3500)	25 (3600)	28 (4100)
Tensile stress @ yield, MPa (psi)	D638	47 (6900)	46 (6700)	50 (7200)	58 (8400)	58 (8400)	50 (7300)
Heat deflection temperature @ 0.455 MPa (66 psi), °C (°F)	D648	74 (165)	73 (164)	70 (163)	69 (156)	69 (156)	70 (158)
Heat deflection temperature @ 1.82 MPa (264 psi), °C (°F)	D648	65 (149)	65 (149)	66 (150)	63 (145)	65 (149)	63 (145)
Specific gravity	D792	1.20	1.20	1.19	1.33	1.33	1.27
Vicat softening point, °C (°F)	D1525	—	—	86 (186)	80 (176)	—	85 (185)
Thermal glass transition temperature, T _g , °C (°F)	—	87 (189)	87 (189)	87 (189)	80 (176)	80 (176)	81 (178)
Barrier							
Oxygen	—	●	●	●	●	●	●
Water	—	●	●	●	●	●	●
Processing							
Drying temperature, °C (°F)	—	71 (160)	70 (160)	70 (160)	160 (320)	150–160 (300–320)	71 (160)
Drying time, hrs	—	3–4	3	4	4–6	4–6	4–6
Melt temperature, °C (°F)	—	232–277 (450–530)	250–290 (480–550)	230–280 (450–530)	277–293 (530–560)	275–295 (530–565)	249–271 (480–520)
Mold temperature, °C (°F)	—	16–38 (60–100)	15–30 (60–80)	15–30 (60–80)	16–32 (60–90)	10–30 (50–90)	16–38 (60–100)
Injection speeds	—	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate
Product summary							
Sterilization							
Gamma	—	●	●	●	●	●	●
EtO	—	●	●	●	●	●	●
E-beam	—	●	●	●	●	●	●
Gas plasma	—	●	●	●	●	●	●
Autoclave	—	○	○	○	○	○	○
Joining							
Solvent bonding	—	●	●	●	●	●	●
Ultrasonic bonding	—	●	●	●	●	●	●
Laser welding	—	●	●	●	●	●	●
Adhesives	—	●	●	●	●	●	●
Swaging (cold bending)	—	●	●	●	●	●	●
Radio frequency welding	—	○	○	○	○	○	○
Thermal bonding	—	○	○	○	○	○	○
Process							
Injection molded	—	●	●	●	●	●	●
Extrusion blow molded	—	○	○	○	○	○	●
Injection blow molded	—	●	●	●	○	●	●

^aNot medical grade ^bFilm properties ^cNo break ^dWith special additives

○ = Poor ● = Fair ● = Average ● = Good ● = Excellent

Injection molded properties											
		Eastman Tritan™ copolyester			Eastman Provista™ copolymer	Tenite™ cellulose			Ecdel™ elastomers		
MN006	MB002	MX711	MX731	MX811	MP002	350A-14 ^a	360A-7	360A-16	9965	9966	9967
<1.0	1.3	<1	<1	<1	1.3	<8.5	<8.5	<8.5	1 ^b	1 ^b	1 ^b
92	91	90	91	92	91	>90	>90	>90	93 ^b	93 ^b	94 ^b
NB ^c	NB ^c	980 (18.4)	860 (16.1)	650 (12.2)	NB ^c	416 (7.8)	203 (3.8)	>533 (>10)	NB ^c	NB ^c	NB ^c
1800 (2.6)	1900 (2.7)	1550 (2.25)	1575 (2.28)	1585 (2.28)	1900 (2.7)	1517 (2.2)	1862 (2.7)	1241 (1.8)	150 (2.2)	150 (2.2)	150 (2.2)
330	300	210	210	140	300	40	50	45	300	400	400
54 (7800)	48 (7000)	53 (7700)	52 (7500)	53 (7700)	48 (7000)	37 (5300)	41 (5900)	30 (4400)	20 (2900)	22 (3200)	23 (3300)
44 (6300)	47 (6900)	43 (6200)	43 (6200)	44 (6400)	47 (6900)	32 (4600)	41 (6000)	27 (3900)	14 (2030)	14 (2030)	13 (1900)
73 (163)	73 (163)	99 (210)	94 (201)	109 (228)	73 (163)	84 (183)	92 (198)	80 (176)	58 (136)	58 (136)	58 (136)
64 (147)	63 (145)	85 (185)	81 (178)	92 (198)	63 (145)	76 (169)	82 (180)	72 (162)	46 (115)	44 (111)	42 (108)
1.23	1.25	1.18	1.18	1.17	1.25	1.20	1.21	1.19	1.13	1.13	1.13
88(190)	85 (185)	—	—	—	85 (185)	100 (212)	107 (225)	92 (198)	170 (338)	170 (338)	170 (338)
85 (185)	85 (185)	110 (212)	110 (212)	120 (248)	85 (185)	110 (230)	118 (244)	97 (207)	-3 (27)	-3 (27)	-3 (27)
●	●	●	●	●	●	○	○	○	●	●	●
●	●	●	●	●	●	○	○	○	○	○	○
71 (160)	70 (160)	88 (190)	88 (190)	88 (190)	70 (160)	70 (160)	70 (160)	70 (160)	65 (150)	65 (150)	65 (150)
6	4-6	4-6	4-6	4-6	4-6	4	4	4	4	4	4
249-271 (480-520)	220-235 (430-450)	260-282 (500-540)	260-282 (500-540)	260-282 (500-540)	210-225 (415-440)	200-225 (390-430)	200-225 (390-430)	200-225 (390-430)	225-260 (435-500)	225-260 (435-500)	225-260 (435-500)
16-38 (60-100)	5-32 (40-90)	38-66 (100-150)	38-66 (100-150)	38-66 (100-150)	15-40 (60-100)	55-60 (130-140)	55-60 (130-140)	55-60 (130-140)	50-80 (120-180)	50-80 (120-180)	50-80 (120-180)
slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate	slow to moderate
●	●	●	●	●	●	●	●	●	●	●	●
●	●	●	●	●	●	●	●	●	●	●	●
●	●	●	●	●	●	●	●	●	●	●	●
●	●	●	●	●	●	●	●	●	●	●	●
○	○	○	○	○	○	○	○	○	●	●	●
●	●	●	●	●	●	●	●	●	●	●	●
●	●	●	●	●	●	●	●	●	○	○	○
●	●	●	●	●	●	●	●	●	○ ^d	○ ^d	○ ^d
●	●	●	●	●	●	●	●	●	●	●	●
○	○	○	○	○	○	○	○	○	●	●	●
○	○	○	○	○	○	○	○	○	●	●	●
●	●	●	●	●	●	●	●	●	●	●	●
○	●	○	○	●	●	○	○	○	●	●	●
○	●	●	●	●	●	○	○	○	●	●	●

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Material 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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