Eastman TRITAN[™] copolyester



Some dos and don'ts for injection molding

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Eastman Tritan[™] copolyester is a thermoplastic material that is ideally suited for a variety of injection molding applications because of its outstanding clarity, toughness, and chemical resistance. Some dos and don'ts that should be helpful in processing this material follow.

Drying

Do

- Keep drying system clean to prevent contamination of Eastman Tritan[™] copolyester.
- Dry thoroughly with a dehumidified drying system to prevent
- Bubbles or appearance problems on finished product.
- Degradation due to hydrolysis, which causes loss of toughness.
- Use air with a dew point of -30° to -40°C (-20° to -40°F).
- Check dehumidifying system with a dew point tester.
- Use air temperatures of 82° to 88°C (180° to 190°F).
 Air temperatures below 65°C (150°F) will substantially increase drying time. Air temperatures above 108°C (226°F) may cause pellets to stick together.
- Dry material for a minimum of 4 hours.
- Keep desiccant bed clean.
- Check heater circuits and elements.
- Control temperature precisely.
- Use air-drying volume of 1.0 cfm/lb [0.06 (m³/min)/kg] of material processed per hour.

Don't

- Attempt to decrease drying time by raising temperature above 108°C (226°F). This may cause pellets to stick together.
- Remove dried material from the drying system until just prior to processing. The moisture level increases rapidly when Eastman Tritan[™] copolyester is exposed to ambient conditions. Even in the hopper, the material will begin to absorb moisture. It is good practice to keep the material in the hopper as dry as possible.

Injection molding

Do

- Keep the material free from contamination to produce tough, clear parts.
- Purge the machine thoroughly, preferably with Eastman Tritan[™] copolyester.
- Use standard molding equipment.
- Use conventional mold-design engineering to include full-round, large runners, large gates, and generously radiused corners.
- Use slow screw speeds of between 30 and 60 rpm, or even slower if recovery is erratic.
- Use slow injection speed to reduce streaking and gate splay. If injection speed is programmable, a slow/moderate/slow profile may be used to produce the best appearance and a reasonable fill time.
- Use an actual melt temperature near the recommended 282°C (540°F) when at the recommended 5–6 minutes or less melt residence time. Cooler melt may contribute to residual stress or screw recovery issues. In cases of longer residence time (big barrel and small shot), consider and test parts from lower melt temperatures approaching 260°C (500°F).
- Plan shot size to be at least 40% and preferably 75% to 80% of machine capacity.
- Expect properties to be dependent on the interplay of temperature, time, and moisture.
- Maintain mold surface temperature around 38° to 66°C (100° to 150°F).

Don't

- Use high-back pressures except when including regrind material or color concentrates.
- Use large mold vents since excessive gas is not produced. Small, conventional vents of 0.001 to 0.0015 in. (0.025 to 0.038 mm) are adequate.
- Discard scrap material. If scrap material is initially dried and processed properly and is free of contamination, it may be blended with virgin material up to a level of 25%.
- Rely on visual inspection to judge the physical properties and quality of a molded piece. Test it under exacting conditions.



The results of insight

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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.

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