

Proper sterilization of packaging made with Eastman Eastalite[™] copolyester

Medical devices must be free from bacteria to ensure patient safety and health. Sterilization, a rigorous process of eliminating these microorganisms, can subject the device—package combination to extreme temperatures, gases, radiation, and other elements that may alter the performance of certain materials Prior or esigning a medical device package, packaging engineers should first consider the method by which the packaged product will be sterilized. Engineers, therefore, need to fully understand the effects of the chosen sterilization method on packaging materials. This technical tip contains guidelines for the proper method of ethylene oxide, gamma irradiation, and electron beam (e-beam) sterilization of copolyesters. These guidelines should be followed to optimize the performance of packaging made with Eastman Eastalite[™] copolyester.

Sterilization methods

Ethylene oxide

- Bacteria are eliminated by penetration of EtO gas mixture into the sealed package.
- Efficiency is boosted by the use of moderately high temperature and humidity.
- Chamber gas can either be 100% EtO or EtO blends with other inert gasses.
- Typical cycle includes 18–24 hours preconditioning (110°F [43°C], 60% relative humidity), 8 hours total cycle (with the average gas dwell of 3–4 hours) all at 125°F (52°C), and a 24-hour aeration at 115°–120°F (46°–49°C).
- Recommended conditions for copolyesters: minimize cycle time; maximum chamber temperature of 130°F (54°C); maximum relative humidity of 50%.

Gamma irradiation

- Bacteria are eliminated by penetration of gamma irradiation from a cobalt-60 source.
- Gamma irradiation provides a high level of penetration at a low dosage rate without the generation of heat. Eastalite is highly resistant to gamma radiation.
- Typical dose is 25 to 50 kGy (2.5 to 5 M Rads). Eastalite has been sterilized up to 100 kGy without any significant physical property loss.

Electron beam radiation

- An electron beam is focused on the product. The beam scans the product to ensure uniform treatment.
- Bacteria are eliminated by penetration of high-energy electrons without heat.
- · Low penetration and high dose rate
- Typical dose is 25 to 50 kGy (2.5 to 5 M Rads). Eastalite has been sterilized up to 75 kGy without any significant physical property loss.

Table 1. Radiation device specifications		
Property	Gamma irradiation	E-beam
Energy spectrum	1.17 and 1.33 MeV	3–10 MeV
Useful power range	15–100 kW	10–200 kW
Typical dose rate	25–50 kGy	25–50 kGy

Conclusion

Gamma, e-beam, and EtO are the gentlest and most common methods for sterilizing copolyesters. However, Eastalite can be successfully and safely sterilized by hydrogen peroxide vapor, peracetic acid vapor, and plasma without any significant changes in physical properties as long as proper processing conditions are used. Proper processing conditions and controls should always be used during sterilization of any copolyester. If excessive high humidity, high temperature, high energy levels, and/or long dwell times are used, increased physical aging or other degrading issues may occur. This may cause brittleness in the package material and could lead to package integrity failure.



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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.

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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.

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