

## Medical Device Packaging Validation

Lansmont's Testing Services Group will recommend appropriate packaging validation procedure(s), which will allow for standardized assessment of a package's ability to withstand the hazards of the distribution environment, including handling, storage, aging, and shelf life. Upon completion of environmental and distribution testing the products ability to function properly and maintain sterility will be evaluated to ensure the packaging system performs as required.



Lansmont will utilize state of the art testing, data acquisition, and environmental simulation equipment to assess the performance of the packaging and its microbial barrier seal. Based upon the results, Lansmont will generate a detailed report that will immediately be made available via electronic media. In addition, guaranteed turnaround ensures that customers will be supplied with the desired results within the shortest possible time frame.



Lansmont's internet based **eTest** will allow customers the option to participate in the actual testing and evaluation via secure, live connection to the appropriate Testing Services Center.

### Lansmont will:

- Provide introductory customer screening
- Help define products to be tested
- Provide pre-test product/package assessment
- Provide environmental screening
- Provide packaging testing
- Provide post-test product/package assessment
- Detailed technical test report with supporting data, digital photography, video as required, concluding with possible recommendations



Testing Services Group – 3 Locations to serve you.  
Lansing, MI • Sunnyvale, CA • Huntington Beach, CA

**1-800-LANSMONT • [www.lansmont.com](http://www.lansmont.com)**

ISTA Certified Laboratories • U.S. D.O.T. Third Party Hazardous Packaging Certification Agencies • Member: AAR, AIAG, ASTM, IoPP, SAE, NIPHLE