

Lot Release Testing

Lancaster Laboratories offers a full range of services to support quality control and release testing to ensure the safety and efficacy of your biopharmaceutical product. Our long history of successful cGMP compliance ensures that the testing we perform for you will be acceptable to FDA, EMEA, and other regulatory authorities. Our experienced scientists can design a Lot Release Program to meet your needs as well as perform assay development and validation.

Analytical services are available to characterize your product as well as confirm its identity, purity, and activity. A full range of services is also available to ensure the absence of contaminants and adventitious agents.

Product Characterization

- Peptide Mapping
- Size Exclusion HPLC
- Carbohydrate Profiling

Activity

- In Vitro Bioassays

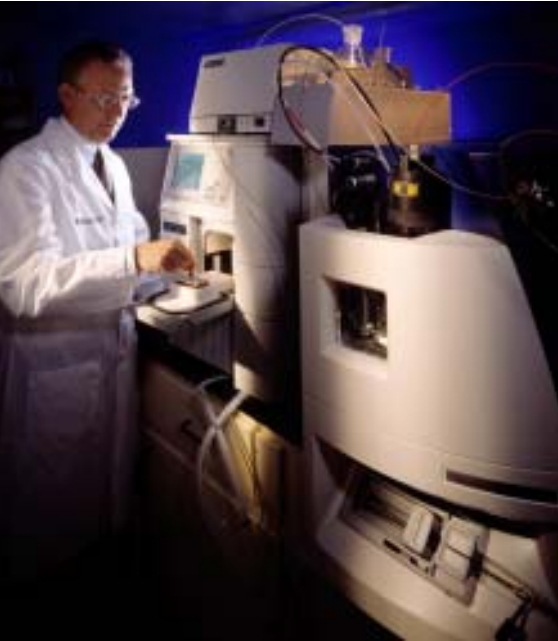
Identity

- SDS Polyacrylamide Gel Electrophoresis (SDS PAGE)
- Western Blotting
- Iso-Electric Focussing (IEF)

Contaminants

- Sterility
- Endotoxin
- Host Cell Proteins
- Host Cell Nucleic Acids
- Adventitious Agents

For more information on our Lot Release Testing or any of our biopharmaceutical capabilities please contact your project manager or Business Development at 717-656-2300.



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