

# Product Release Testing /Retain Program Administration

Lancaster Laboratories offers comprehensive analytical support for release testing. From immediate release solid dosage formulations to controlled release solid dosage formulations, from liquids to suspensions, from aerosol products to transdermal products, we have the experience and capability to provide you with what Lancaster Laboratories has earned a reputation for: quality analytical data delivered on time.

Our multi-shift, experienced staff has performed release testing of in-process materials as well as finished products. Whether you need results in several days or overnight, we have the size and capacity to meet your most demanding turnaround time requirements. We have the flexibility within our organization to meet the ever-changing demands of production schedules and timelines.

Besides having the analytical testing capabilities to perform release testing, Lancaster Laboratories can take over administration of your product retain program. Upon arrival at Lancaster Laboratories, GMP reserve samples are documented as to date and time of receipt and taken directly to our reserve sample storage area. We take inventory, compare the inventory with protocols and then log these samples into our reserve sample tracking database.

When samples are received from a third party manufacturer, Lancaster Laboratories can also perform a GMP quality check. This check can include, but is not limited to, label quality, lot number, print quality, container quality and container closure. The inspection can be tailored to meet your in-house standard.

Reserve samples can be scheduled for physical observation analysis per your requirements. We generate "pull reports" from our scheduling database for each workday, pull the samples as scheduled and log the samples due for testing into our laboratory information management system (LIMS). At this point, we send you an acknowledgment, letting you know that the appropriate samples have been pulled and are scheduled for the required testing. All steps of the storage and testing process are tightly controlled and accurately documented.

We welcome client visits and audits to verify our strict compliance with current GMP requirements and to demonstrate the experience of our scientists. We're equipped with state-of-the-art instrumentation, and all analytical operations are based on written, detailed SOP's and stringent quality assurance practices.

For more information on our release testing capabilities or any of our pharmaceutical capabilities, please contact your project manager or Pharmaceutical Business Development at 717-656-2300.



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