



Experience, flexibility and quality time after time.



Topical Dosage (Gels, Lotions, Creams and Liquids) Product Development and Clinical Manufacturing Phase I and II

SL Pharma Labs has experience in working with both large and small molecules as well as novel technologies. Clients range from large, multi-national through virtual start-up organizations. The commonality across the client base from innovator drug companies through generic organizations is a desire for efficient collaborative support that is done on time and on budget every time.

Topical dosage product development and clinical manufacturing services include:

- » pre-formulation and formulation development
- » analytical development and validation
- » microbiological development and validation
- » process development
- » lab-scale, feasibility and batch manufacture
- » container and closure compatibility and integrity testing
- » batch release and stability specification development and testing
- » packaging and labeling
- » technology transfer
- » development of technical/regulatory documentation



SL Pharma Labs is located in the First State Industrial Park campus which is approximately 35 minutes from the Philadelphia International Airport and 5 minutes from the I-95 Newport exit in Delaware.

The facility houses formulation development, chemistry, microbiology and analytical laboratories, stability chambers, QA storage and clean rooms ranging from Class 100 – Class 100,000. SL Pharma Labs can handle cytotoxic compounds and Controlled Substances Schedules II through V. Additionally, the facility operates under cGMP and maintains a clean record with the FDA since the establishment of the company in 1997.

SL Pharma Labs, Inc.
1300 First State Blvd, Suite C
Wilmington, DE 19804
T (302) 636-0202 / F (302) 636-0204
info@slpharmalabs.com
<http://www.slpharmalabs.com>