Admix Pharma Laboratories, LLC.

CONTRACT TESTING AND PRODUCT DEVELOPMENT LABORATORY SERVICES



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With Admix Pharma Laboratories, LLC

you will enjoy a host of attractive benefits that can effectively extend your in-house pharmaceutical products development and manufacturing capabilities for long and short term projects. You will quickly realize how efficiently and effectively you can relieve the internal workload of your laboratory and pharmaceutical products development department. In addition, you have our commitment to meet all agreed-upon delivery dates.

All the Services You're Looking For

You'll find that Admix Pharma Laboratories,

LLC offers a full range of technologically advanced services. You can choose the specific services that offer the most appropriate solutions to all your needs.

Make the Most of Your Time & Resources

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Ask Yourself 5 Simple Questions

- 1. Has your company identified a number of products you would like to bring to market?
- 2. Does your company require additional Product Development capacity?
- 3. Does your company require additional Analytical Services capacity?
- 4. Does your company require additional Manufacturing capacity?
- 5. Does your company require additional Regulatory capacity?
 - If your answer to **2 or more** of these questions is YES, then your company needs the help of **Admix Pharma Laboratories, LLC**., one of the most knowledgeable, experienced, and capable companies in the field of pharmaceutical products contract development and manufacturing.



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Our Services

The Scope of Our Services Includes

Analytical Site

- Active Pharmaceutical Ingredients
 Testing
- Inactive Pharmaceutical Ingredients/Excipients Testing
- Liquid, Solid, and Semi-Solid Dosage Forms Products Testing
- Stability Testing
- Analytical Method Development and Validation
- Cleaning Validation Testing



Product Development Site

- Liquid, Solid and Semi-Solid Dosage
 Forms Product Development
- Active Pharmaceutical Ingredients
 Sourcing
- Inactive Pharmaceutical Ingredients/Excipients Sourcing
- Product Marketability Evaluation (will it be financially effective to bring a proposed product to market).
- Pilot Batches Manufacturing and Stability Evaluation
- Stability Indicating Analytical Testing Method Development and Validation
- Submission (ANDA)/Stability Batch Manufacturing, Testing, and Stability Evaluation
- Identification/Selection/Evaluation of an Appropriate CRO for Performance of Bioequivalence Studies
- Bioequivalence Studies Performance
 Monitoring
- Bioequivalence Studies Reports
 Evaluation
- Compiling ANDA Submissions
- Electronic Submission of ANDA (e-CTD format)
- Pre-marketing Process Validation
- Commercial Scale Manufacturing