



CoreRx, Inc.

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ABOUT

Year Founded: 2006

Number of Employees: 7

Key Personnel: Dr. Todd R. Daviau, Director, Analytical & Preformulation; Jim Davis, Director – Analytical Laboratory; Brian McMillan, Director - Formulations; Rob Trivedi, Director – Technical Operations; Mark Licarde, Director - Manufacturing.

WHO WE ARE

CoreRx, formed in the spring of 2006, already has a proven record of developing formulations for clients throughout the country.

The management team of CoreRx was the Management team of the Pharmaceuticals Division of MDS Pharma Services prior to its divestiture in 2006. Our commitment to unsurpassed quality in drug discovery and innovative development solutions has allowed us to consistently and efficiently move compounds through the research and development stages to market.

CoreRx's unique corporate structure creates teams led by accomplished scientists. These scientific teams work intensively with each client, becoming an extension of their own organization.

SERVICES OFFERED

CoreRx's Services include but are not limited to:

Preformulation

CoreRx's range of preformulation services allow us to characterize your API at the earliest stage of drug development. Our scientists have experience in **preformulation** of new substances, **deformulation** and reformulation of new and generic pharmaceutical products.

Formulation Development

The scientific team creates options for formulation development to meet your dosage range and preferred route of delivery.

We are able to formulate the following dosage forms:

- Solids (tablet formulation, hard gelatin capsule formulation, granules and powders)
- Liquids (oral and topical liquids, suspensions, lyophilized dosage forms, parenterals and emulsions)
- Semi-solids (creams, ointments, gels, liquids, suspensions, and solutions)

Clinical Trial Manufacturing

Our scientists are some of the industry's most respected in the field of formulation development scale up.

- Manufacturing batches for stability testing, proof of concept or prototype development,
- Scaling up and production of pilot batches,
- Manufacturing of clinical trials supplies from post drug discovery through phase III,
- Process validation and technology transfer.

Analytical

At CoreRx™ our services center on small molecule analysis - including deformulation, preformulation, method development and qualification, drug substance and finished product release, and stability testing.

- FDA and ICH guideline-driven analytical capabilities
- State-of-the art facilities and analytical instrumentation provide an optimal environment for stability studies, including a full range of temperature and relative humidity storage conditions
- Stand-alone analytical projects or fully integrated development support,

Facility Overview

CoreRx's Tampa laboratories are fully cGMP compliant and registered with the FDA and DEA. We are very proud of our world-class scientists and unparalleled environment.

CORE STRENGTHS

CoreRx™ focuses on supporting and streamlining the drug development process for biotechnology and pharmaceutical innovators by offering services such as pre-formulation/formulation development, analytical development, and clinical manufacturing. We have a proven track record and extensive expertise providing comprehensive drug development services to biotechnology and pharmaceutical companies worldwide.

At CoreRx™ we maintain open communication with our clients, so that we can provide innovative solutions efficiently and on budget while maintaining our core values of quality, integrity, and efficiency.

CoreRx™ offers diversified technical resources, capacity, flexibility, and the experience to develop and manufacture virtually any clinical dosage form with strict quality compliance and value.

Our Priority is your success...