



Xcelience
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Xcelodose® 600

- Stand alone analytical projects or fully integrated development support

Clinical Trial Manufacturing:

- 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 to phase IIb, including individual capsule filling of API, tableting, encapsulation, placebo matching, over-encapsulation or coating of comparators.
- Process validation and technology transfer

Xcelience has expertise and technology in direct API-to-capsule filling, utilizing the Xcelodose® 600 system for precision micro filling of API into capsules. In addition, the CFS 1200™ system allows for rapid development and manufacturing of multiple strengths of simple liquid-filled capsule products.

Xcelodose® and CFS1200™ are registered trademarks of Capsugel BVBA.

Xpert Consulting:

Xcelience has launched a product-development consulting service to help its client base in planning, executing and evaluating the development of drug products. The Xpert Consulting service uses specialized expertise within Xcelience's scientific staff and is spearheaded by Dr. Steve Bannister, Scientific Director and Principal Consultant. His scientific training and experience includes pharmaceuticals, analytical chemistry, bioanalysis and pharmacokinetics. He has managed components of product development at the laboratory level and directed integrated discovery and development functions at the executive level.

Facility Overview

Xcelience's 50,000-sq.-ft. Tampa laboratories are fully cGMP compliant and registered with the FDA and DEA.

Vital Statistics

Year Founded: 1997

Number of Employees: 75

Key Personnel: Derek Hennecke, president and chief executive officer; Randall Guthrie, vice president; Dr. Steve Bannister, scientific director; Irene LoJacono, director, project management.

Who We Are

XCELIENCE, a Tampa-based pharmaceutical contract research organization, is the premier source for unsurpassed quality in drug development. Our experienced scientists' commitment to quality in supporting drug discovery and innovative drug development solutions, has allowed us to consistently and efficiently move compounds through the research and development stages to market. Xcelience's unique corporate structure creates teams led by accomplished project managers.

Major Markets

The Pharmaceutical Industry.

Services Offered

XCELIENCE'S core services include but are not limited to:

Preformulation:

Xcelience's range of preformulation studies allows us to characterize your API at the earliest stage of drug development

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, and emulsions)
- Semi-solids (creams, ointments, gels, liquids, suspensions, and solutions)

Analytical:

- 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

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