

Florida Biologix®
13702 Innovation Drive
Alachua, FL 32615
Tel: (386) 418-8199 x203 | (877) 765-7676
Fax: (386) 462-7835
Web: www.floridabiologix.ufl.com
E-mail: caytug@floridabiologix.ufl.com



Vital Statistics

Year Founded: 2006

Number of Employees: 29

Key Personnel: Director, Richard O. Snyder, Ph.D.; Associate Director, Joyce D. Francis, Ph.D.; Scientific Coordinator, Robert K. Zwerner, Ph.D.; Director of Marketing & Sales, Chris Aytug, MBA; Director of Operations, Linna Lim, M.S.

Who We Are

Florida Biologix® is a Phase I/II contract development, manufacturing & testing organization that offers a wide range of biopharmaceutical services. Our main building is a 23,000-square foot, state-of-the-art, validated, multi-product manufacturing and testing facility; and the adjacent building houses our 5,000 square foot, fully-equipped, process development laboratories. Florida Biologix can provide a services package starting from process design and development through full-scale cGMP compliant manufacturing, purification and aseptic filling or we can work with you to develop customized programs based on your specific needs. Our excellent in-house quality department provides a full range of analytical services, facility and systems monitoring, product review and disposition and regulatory support including CMC and DMF preparation.



Florida Biologix is committed to quality, which ensures that your end product is optimal and the process exceeds expectations. Our collaborative approach ensures that the client is a partner and well informed throughout the project. Our team is flexible and experienced, which ensures that technical or business issues are handled smoothly if they arise. This allows us to meet your outsourcing needs quickly and cost-effectively; accomplishing your goal of moving biologics from the bench to the clinic. Our primary focus is to develop, produce, test and/or fill cGMP compliant biopharmaceutical clinical trial material on time and at a reasonable price.

Major Markets

Florida Biologix® serves the biotechnology industry, medical device industry and biomedical research institutions.

Services Offered

- Process and/or Product Development**
 Production batch record development, cell line engineering, clone isolation and expansion, cell culture development, purification and analytical assay development, troubleshooting, product formulation and stability studies
- cGMP Bulk Substance Production in Mammalian Cell Culture Systems**
 Recombinant proteins, enzymes, antibodies, vaccines
 Upstream cell culture and downstream purification and processing
 Master and working cell bank production
 Quality control assay validations
- Sterile Fill and Finish**
 Formulation and aseptic fill and finish into a variety of final containers, including glass vials, bags, syringes, bottles, cryovials and conical tubes via automated, semi-automated or manual processes. Stability studies and distribution to clinical sites.
- Additional Services**
 GMP storage, Quality Control Services, Quality Assurance Services, Long Term Stability Studies, Technology Assessment and Licensing Consultation, Assistance in Outsourcing Preclinical and Clinical Studies, Production of Non-cGMP Material for Research Purposes, Kit Assembly, Project Management, Regulatory Agency Audit Support

