

Are you interested in first-hand experience in development and manufacturing of pharmaceutical and therapeutic products? Are you motivated, enjoy problem solving, and have great attention to detail?

In this role, you'll be an integral part of a biotech startup and be able to experience what it takes to bring products to market. Working in Quality Assurance, you'll have responsibility to work with personnel across all departments and help develop efficient and effective processes and systems. Your work will directly contribute to cGMP and Quality System Compliance in our FDA-regulated company. You'll have an opportunity to contribute to a fast-growing environment and learn from an experienced team.

The ideal candidate is pursuing a BS or MS in Biomedical Engineering, Biology, or similar degree. Some experience in an FDA-regulated industry is desired but not required. Basic familiarity with Quality and Regulatory requirements is desired.

Captozyme is a leader in health care using novel enzymes and live biotherapeutics. Whether creating our own pipeline of products or working with partners to advance the discovery of new opportunities, we are committed to advancing gut health research and making a lasting difference in the industry. Learn more at [www.Captozyme.com](http://www.Captozyme.com).

Interested candidates should send their resume to [jobs@captozyme.com](mailto:jobs@captozyme.com).

Captozyme Inc.  
1622 NW 55<sup>th</sup> Place  
Gainesville, FL 32653  
[www.captozyme.com](http://www.captozyme.com)