



BioLink Life Sciences, Inc.

Enhancing the POWER of Drugs & Diagnostics

250 Quade Drive, Cary, NC 27513

New BioLink Capabilities Accelerate FDA Approvals for Clients

Cary, N.C. – June 9, 2009 – BioLink Life Sciences has added cGMP-compliant analytical testing to its portfolio of services that enable their clients to accelerate progress toward FDA approvals. BioLink provides both analytical and synthetic support to biomedical, pharmaceutical, and life sciences companies in the U.S. and abroad. The Company focuses its expertise on solving problems in a cost- and time-effective manner and links its clients to other experts with GMP manufacturing, preclinical, and regulatory expertise. This “value plus” combination provides a quicker path to FDA approval.

GMP refers to the Good Manufacturing Practice regulations enforced by the US Food and Drug Administration. These regulations, which have the force of law, require that manufacturers, processors, and packagers of drugs, medical devices, foods, and supplements take proactive steps to ensure that their products are safe, pure, and effective. “By significantly upgrading our equipment and IT systems, we have ensured compliance with current GMP,” BioLink’s President Deanna Nelson noted. “The enhancement allows us to better meet client needs in 2009 and beyond.”

BioLink Life Sciences, Inc. is a privately owned biotechnology and life sciences company with headquarters and laboratories located in Cary, NC. Since the Company’s founding in 2001, BioLink has provided contract research and development support to other pharmaceutical and biotech companies, who are developing new approaches for the diagnosis and treatment of a variety of diseases.

To learn more about BioLink’s products, contact BioLink at (919) 678-9478, or visit the company’s website at www.biolinkonline.com.

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