



Clinical Data to Present at Lazard Life Sciences Conference

NEWTON, Mass., Nov 27, 2006 (BUSINESS WIRE) -- Clinical Data Inc. (NASDAQ: CLDA) today announced that it will participate in the Lazard Capital Markets 3rd Annual Life Sciences Conference in New York City on Wednesday, November 29, 2006.

Drew Fromkin, President and CEO, will speak to the attendees of the conference at 8:00 a.m. EST. A live audio webcast of the presentation will be available to all interested parties at Clinical Data's Investor Relations website at <http://www.clda.com>. An archived replay of the webcast will also remain accessible for 90 days through February 27, 2007, within the "Events and Presentations" area of the "Investors" section of the Clinical Data website.

About Clinical Data, Inc.

Clinical Data, Inc., a worldwide leader in providing comprehensive molecular and pharmacogenomics services as well as genetic tests to improve patient care, is organized under three worldwide divisions segmented by service offerings and varying client constituents: PGxHealth(TM); Cogenics(TM); and Vital Diagnostics(TM). The Pharmacogenomics and Molecular Services (TM) division, Cogenics, consolidates operations of Genaissance Pharmaceuticals, Inc., Lark Technologies, Inc., Icoria, Inc., and Genome Express SA to provide a comprehensive range of molecular biology and pharmacogenomics services to pharmaceutical, biotech, academic, agricultural, and government clients. These services are offered in both research and regulated environments and have applications across the lifecycle of pharmaceutical product development including pharmacovigilance requirements post-launch. PGxHealth builds upon existing assets and know-how acquired from Genaissance Pharmaceuticals, Icoria, and Genome Express in the areas of genomics-based, genetic tests and therapeutic efficacy and safety biomarker development for drug utilization. PGxHealth develops, validates and commercializes novel Therapeutic Diagnostics(TM), in some instances in combination with new and existing therapeutics, to improve patient care. In addition, PGxHealth has a therapeutic drug candidate, vilazodone, currently in late stage clinical trials for the treatment of depression. Vital Diagnostics serves the clinical laboratory in the traditional in-vitro diagnostics market worldwide. With a focus on the physician's office, hospital and small-to-medium sized laboratory segments and customers in approximately 100 countries, Vital Diagnostics has achieved a leading market share for instruments and reagents sold into moderately complex physicians' office laboratories within the United States. Clinical Data is headquartered in Newton, Mass. with operations in Texas, Connecticut, North Carolina, Rhode Island, and California as well as internationally in the UK, France, the Netherlands, and Italy.

SAFE HARBOR STATEMENT UNDER THE PRIVATE SECURITIES LITIGATION REFORM ACT OF 1995

This press release contains certain forward-looking information about Clinical Data that is intended to be covered by the safe harbor for "forward-looking statements" provided by the Private Securities Litigation Reform Act of 1995, as amended. Forward-looking statements are statements that are not historical facts. Words such as "expect(s)", "feel(s)", "believe(s)", "will", "may", "anticipate(s)" and similar expressions are intended to identify forward-looking statements. These statements include, but are not limited to statements regarding: our ability to successfully integrate the operations, business, technology and intellectual property obtained in our acquisitions; our ability to obtain regulatory approval for, and successfully introduce our new products; our ability to expand our long-term business opportunities; our ability to maintain normal terms with our customers and partners; financial projections and estimates and their underlying assumptions; and statements regarding future performance. All of such statements are subject to certain risks and uncertainties, many of which are difficult to predict and generally beyond the control of the Company, that could cause actual results to differ materially from those expressed in, or implied or projected by, the forward-looking information and statements. These risks and uncertainties include, but are not limited to: whether vilazodone will advance further in the clinical trials process and whether and when, if at all, vilazodone will receive final approval from the U.S. Food and Drug Administration and equivalent foreign regulatory agencies and for which indications; whether vilazodone will be successfully marketed if approved; the extent to which genetic markers (haplotypes) are predictive of clinical outcomes and drug efficacy and safety; our ability to achieve the expected synergies and operating efficiencies from all of our acquisitions; the strength of our intellectual property rights; competition from pharmaceutical, biotechnology and diagnostics companies; the development of and our ability to take advantage of the market for pharmacogenetic and biomarker products and services; general economic downturns; and other risks contained in our various SEC reports and filings, including but not limited to our Quarterly Report on Form 10-Q for the quarter ended September 30, 2006, and our subsequent Current Reports on Form 8-K filed with the Securities and Exchange Commission. Our audience is cautioned not to place undue reliance on these forward-looking statements that speak only as of the date hereof, and we do not undertake any obligation to revise and disseminate forward-looking statements to reflect events or circumstances after the date hereof, or to reflect the occurrence of or non-occurrence of any events.

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