



Discovery Labs Provides Status of Previously Submitted Response to FDA Approvable Letter for Surfaxin[®]

Warrington, PA, August 15, 2005 — **Discovery Laboratories, Inc. (Nasdaq: DSCO)**, announces that it has received a preliminary communication from the U.S. Food and Drug Administration (FDA) indicating that the Company's previously submitted Response Letter is not considered complete at this time. The FDA notified Discovery that it will provide, later this week, its official written communication regarding select matters for which the FDA may be requesting additional information. The Company previously submitted its Response Letter to the FDA's Approvable Letter for Surfaxin[®] (lucinactant) for the prevention of Respiratory Distress Syndrome (RDS) in premature infants on July 29, 2005.

Robert J. Capetola, Ph.D., President and CEO of Discovery, commented, "The response we recently submitted to the FDA's Approvable Letter for Surfaxin was encyclopedic and comprehensive. Based upon the thoroughness of our Response Letter, we are optimistic that the FDA's issues will center on a few select items that are capable of being addressed in an as timely and efficient manner as possible. Of course, we will have greater clarity on these matters once we have received and fully assessed the FDA's official communication which we expect later this week. At that time, we anticipate holding a conference call to communicate more detailed information with our shareholders. We believe that our relationship with the FDA has always been collaborative and very productive. We anticipate prompt discussions with the FDA and moving forward to finalize a complete response as quickly as possible."

DISCLOSURE NOTICE: The information in this press release includes certain "forward-looking" statements relating to the timing of FDA matters relating to Discovery's NDA for Surfaxin for RDS, which assume that all of the conditions in the Approvable Letter are timely satisfied. Any statements contained in this press release that do not describe historical facts are forward-looking statements that involve risks and uncertainties that could cause actual results to differ materially from those discussed in such forward-looking statements. Such risks and uncertainties include the following: uncertainties relating to Discovery Labs' ability to successfully address the comments and concerns of the FDA contained in the approvable letter for Surfaxin, including uncertainties relating to the sufficiency of the additional details to be submitted to the FDA by Discovery Labs; the timing of FDA actions regarding Surfaxin, including the timing of its review of the supporting data or the additional details submitted in relation thereto; the outcome of FDA actions in response to the submissions, including the possibility that the FDA will not consider the submissions complete or that the FDA will require additional information in support of the submissions; the possibility that even if the response is deemed complete by the FDA, the FDA could respond to these submissions by issuing an additional approvable letter with additional conditions for approval or the FDA could issue a not approvable letter; the ability to resolve final labeling for Surfaxin with the FDA; uncertainties regarding market acceptance of Surfaxin; uncertainties relating to third-party reimbursements; uncertainties relating to Discovery Labs' ability to continue to operate at commercial scale in compliance with FDA regulations and other applicable manufacturing requirements; uncertainties relating to patents and proprietary rights; issues previously raised by earlier FDA inspections of the Totowa, NJ facility of Laureate Pharma, Inc. (Laureate), Discovery's contract manufacturer for Surfaxin, and other risks identified in Discovery Labs' Securities and Exchange Commission filings. Many factors could cause the resolution of those

or other issues to differ materially from Discovery's forward-looking statements, including that the timing, scope and duration of a resolution of the issues related to the Approvable Letter will depend on the ability of Discovery (and Laureate) to assure the FDA of the quality and reliability of its basic quality controls, process assurances and documentation requirements that support the commercial production manufacturing process under applicable cGMPs. The reader of this release should understand that the failure to reach resolution of any issues could result in delays in ultimate approval of Discovery's potential products beyond the first quarter of 2006. Discovery Labs further cautions readers not to place undue reliance on any forward-looking statements which speak only as of the date they are made. Discovery Labs disclaims any obligation to publicly update or revise any such statements to reflect any change in expectations or in events, conditions or circumstances on which any such statements may be based, or that may affect the likelihood that actual results will differ from those set forth in the forward-looking statements.

About Discovery Labs

Discovery Laboratories, Inc. is a biotechnology company developing its proprietary surfactant technology as Surfactant Replacement Therapies (SRT) for respiratory diseases. Surfactants are produced naturally in the lungs and are essential for breathing. Discovery's technology produces a precision-engineered surfactant that is designed to closely mimic the essential properties of natural human lung surfactant. Discovery believes that through its technology, pulmonary surfactants have the potential, for the first time, to be developed into a series of respiratory therapies for patients in the neonatal intensive care unit, critical care unit and other hospital settings, where there are few or no approved therapies available.

Discovery has received an Approvable Letter from the FDA for Surfaxin, the Company's lead product, for the prevention of Respiratory Distress Syndrome (RDS) in premature infants, and has filed a Marketing Authorization Application with the EMEA for clearance to market Surfaxin in Europe. Discovery is also conducting various clinical programs to address Acute Respiratory Distress Syndrome (ARDS) in adults, Bronchopulmonary Dysplasia (BPD) in premature infants, Neonatal Respiratory Failures in premature infants, severe asthma in adults, and Meconium Aspiration Syndrome (MAS) in full-term infants.

For more information, please visit our corporate website at www.Discoverylabs.com.

To the extent that statements in this press release are not strictly historical, including statements as to business strategy, outlook, objectives, future milestones, plans, intentions, goals, future financial conditions, future collaboration agreements, the success of Discovery's product development, events conditioned on stockholder or other approval, or otherwise as to future events, all such statements are forward-looking, and are made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. These forward-looking statements are subject to certain risks and uncertainties that could cause actual results to differ materially from the statements made. Among the factors which could affect Discovery's actual results and could cause results to differ from those contained in these forward-looking statements are the risk that financial conditions may change, risks relating to the progress of Discovery's research and development, the risk that Discovery will not be able to raise additional capital or enter into additional collaboration agreements (including strategic alliances for our aerosol and Surfactant Replacement Therapies), risk that Discovery will not be able to develop a successful sales and marketing organization in a timely manner, if at all, risk that Discovery's internal sales and marketing organization will not succeed in developing market

awareness of Discovery's products, risk that Discovery's internal sales and marketing organization will not be able to attract or maintain qualified personnel, risk of delay in the FDA's or other health regulatory authorities' approval of any applications filed by Discovery, risks that any such regulatory authority will not approve the marketing and sale of a drug product even after acceptance of an application filed by Discovery for any such drug product, risks relating to the ability of Discovery's third party contract manufacturers to provide Discovery with adequate supplies of drug substance and drug products for completion of any of Discovery's clinical studies, other risks relating to the lack of adequate supplies of drug substance and drug product for completion of any of Discovery's clinical studies, and risks relating to the development of competing therapies and/or technologies by other companies. Companies in the pharmaceutical and biotechnology industries have suffered significant setbacks in advanced clinical trials, even after obtaining promising earlier trial results. Data obtained from tests are susceptible to varying interpretations, which may delay, limit or prevent regulatory approval. Those associated risks and others are further described in Discovery's filings with the Securities and Exchange Commission including the most recent reports on Forms 10-K, 10-Q and 8-K, and any amendments thereto.

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