



Contact: Tricia J. Richardson
Novavax, Inc.
240-268-2031

Novavax CEO to Present at Phacilitate Vaccine Forum Barcelona 2009

ROCKVILLE, MD (June 18, 2009) – /PRNewswire-FirstCall/ - Novavax, Inc. (NASDAQ: NVAX) a clinical-stage vaccine company, announced today that Dr. Rahul Singhvi, President and Chief Executive Officer, will be presenting at the Phacilitate Vaccine Forum Barcelona 2009 conference on June 22 and June 23, 2009 at The Fira Palace, Barcelona, Spain. Dr. Singhvi is participating in the Panel Session ‘*How are VLP based vaccines helping to shape the manufacturing processes of the future?*’ and will also be speaking to the Case Study: *Highlighting the clinical milestones to date: What progress is being made with respect to immune response in healthy adults?* Presentations will be available on www.novavax.com after the event.

ABOUT NOVAVAX

Novavax, Inc. (Nasdaq: NVAX) is a clinical stage biotechnology company, creating novel vaccines, including H1N1, to address a broad range of infectious diseases worldwide using advanced proprietary virus-like particle (VLP) technology. The Company produces these VLP based, potent, recombinant vaccines utilizing new and efficient manufacturing approaches. Additional information about Novavax is available at www.novavax.com and in the Company’s various filings with the Securities and Exchange Commission.

Forward-Looking Statement

Statements herein relating to future development results and performance, conditions or strategies and other matters, including expectations regarding product and clinical developments, are forward-looking statements within the meaning of the Private Securities Litigation Reform Act. Novavax cautions that these forward-looking statements are subject to numerous assumptions, risks and uncertainties, which change over time. Factors that may cause actual results to differ materially from the results discussed in the forward-looking statements or historical experience include risks relating to the early stage of Novavax’s product candidates under development; current results may not be predictive of future pandemic results, results of our seasonal influenza vaccine or any other vaccine that we may develop; further testing is required before regulatory approval can be applied for and the FDA may not approve a vaccine even if further trial results are similar to those disclosed previously by the company; uncertainties relating to clinical trials, including possible delays in initiating or completing the trials and safety and efficacy results; dependence on the efforts of third parties; competition for clinical resources and patient enrollment from drug candidates in development by other companies with greater resources and visibility; and risks that we may lack the financial resources and access to capital to fund our operations including further clinical trials. Further

information on the factors and risks that could affect Novavax's business, financial conditions and results of operations, is contained in Novavax's filings with the U.S. Securities and Exchange Commission, which are available at <http://www.sec.gov>. These forward-looking statements speak only as of the date of this press release, and Novavax assumes no duty to update forward-looking statements.

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