

## **Soligenix to Present at the 20th Annual GCFF Investor Conference in Vancouver, British Columbia, Canada**

### **Review of Two Phase 3 Clinical Programs with Near-Term Readouts**

PRINCETON, N.J., Oct. 3, 2019 /PRNewswire/ -- Soligenix, Inc. (Nasdaq: SNGX) (Soligenix or the Company), a late-stage biopharmaceutical company focused on developing and commercializing products to treat rare diseases where there is an unmet medical need, announced today that its Chief Scientific Officer, Oreola Donini, PhD, will deliver a corporate presentation to review the Company's two pivotal phase 3 clinical programs in cutaneous T-cell lymphoma and oral mucositis in head and neck cancer at the Global Chinese Financial Forum (GCFF) Conference on Saturday, October 5, 2019 at 11:55 am Pacific Time. The event will take place at Vancouver Executive Hotel, Vancouver, BC, Canada.

To access the Soligenix corporate presentation, please visit [here](#).

### **About the GCFF and the 2019 GCFF Conference**

The GCFF is the most prominent series of bi-lingual financial functions in both North America and China. Established in 2000, GCFF's mandate is to provide a world-class platform connecting both the China and North American financial markets, bringing together companies, financial institutions and investors who are interested in exploring new investment opportunities.

With an extensive network covering both the North American and Chinese markets, GCFF is supported by a strong foundation of financial resources. This enables GCFF to organize resourceful financial events that facilitate business growth and networks among financial institutions, public companies, private companies, and investors of all levels.

The 2019 (20<sup>th</sup> Annual) GCFF Conference brings together investment opportunities in the small-cap growth market in North America to meet with the largest and fastest growing Chinese investor community in Canada. The Annual Main Event will be a great platform for the Chinese investors to learn the latest market trends and share market insights.

For more information about the conference, please refer to the [conference website](#).

### **About Soligenix, Inc.**

Soligenix is a late-stage biopharmaceutical company focused on developing and commercializing products to treat rare diseases where there is an unmet medical need. Our Specialized BioTherapeutics business segment is developing SGX301 as a novel photodynamic therapy utilizing safe visible light for the treatment of cutaneous T-cell lymphoma, our first-in-class innate defense regulator (IDR) technology, dusquetide (SGX942) for the treatment of oral mucositis in head and neck cancer, and proprietary formulations of oral beclomethasone 17,21-dipropionate (BDP) for the prevention/treatment of gastrointestinal (GI) disorders characterized by severe inflammation including pediatric Crohn's disease (SGX203) and acute radiation enteritis (SGX201).

Our Public Health Solutions business segment includes active development programs for RiVax<sup>®</sup>, our ricin toxin vaccine candidate and SGX943, our therapeutic candidate for antibiotic resistant and emerging infectious disease. The development of our vaccine programs incorporates the use of our proprietary heat stabilization platform technology, known as ThermoVax<sup>®</sup>. To date, this business segment has been supported with government grant and contract funding from the National Institute of Allergy and Infectious Diseases (NIAID), the Biomedical Advanced Research and Development Authority (BARDA), and the Defense Threat Reduction Agency (DTRA).

For further information regarding Soligenix, Inc., please visit the Company's website at [www.soligenix.com](http://www.soligenix.com).

This press release may contain forward-looking statements that reflect Soligenix, Inc.'s current expectations about its future results, performance, prospects and opportunities, including but not limited to, potential market sizes, patient populations and clinical trial enrollment. Statements that are not historical facts, such as "anticipates," "estimates," "believes," "hopes," "intends," "plans," "expects," "goal," "may," "suggest," "will," "potential," or similar expressions, are forward-looking statements. These statements are subject to a number of risks, uncertainties and other factors that could cause actual events or results in future periods to differ materially from what is expressed in, or implied by, these statements. Soligenix cannot assure you that it will be able to successfully develop, achieve regulatory approval for or commercialize products based on its technologies, particularly in light of the significant uncertainty inherent in developing therapeutics and vaccines

against bioterror threats, conducting preclinical and clinical trials of therapeutics and vaccines, obtaining regulatory approvals and manufacturing therapeutics and vaccines, that product development and commercialization efforts will not be reduced or discontinued due to difficulties or delays in clinical trials or due to lack of progress or positive results from research and development efforts, that it will be able to successfully obtain any further funding to support product development and commercialization efforts, including grants and awards, maintain its existing grants which are subject to performance requirements, enter into any biodefense procurement contracts with the U.S. Government or other countries, that it will be able to compete with larger and better financed competitors in the biotechnology industry, that changes in health care practice, third party reimbursement limitations and Federal and/or state health care reform initiatives will not negatively affect its business, or that the U.S. Congress may not pass any legislation that would provide additional funding for the Project BioShield program. In addition, there can be no assurance as to timing or success of the Phase 3 clinical trial of SGX942 (dusquetide) as a treatment for oral mucositis in patients with head and neck cancer receiving chemoradiation therapy or the Phase 3 clinical trial of SGX301 (synthetic hypericin) for the treatment of cutaneous T-cell lymphoma. There also can be no assurance as to timing or success of the preclinical/clinical trials of RiVax<sup>®</sup>, that RiVax<sup>®</sup> will be approved for the PRV program or the amount for which a PRV for RiVax<sup>®</sup> can be sold. These and other risk factors are described from time to time in filings with the Securities and Exchange Commission, including, but not limited to, Soligenix's reports on Forms 10-Q and 10-K. Unless required by law, Soligenix assumes no obligation to update or revise any forward-looking statements as a result of new information or future events.

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