
FORM 6-K

SECURITIES AND EXCHANGE COMMISSION
Washington D.C. 20549

REPORT OF FOREIGN ISSUER
PURSUANT TO RULE 13a-16 OR 15d-16 OF THE
SECURITIES EXCHANGE ACT OF 1934

For the month of September, 2009

Commission File Number: 001-32371

SINOVAC BIOTECH LTD.

39 Shangdi Xi Road
Haidian District
Beijing 100085, People's Republic of China
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F.

Form 20-F Form 40-F

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)
(1): _____

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)
(7): _____

Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also thereby furnishing the information
to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes No

If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b):

82- N/A

SINOVAC BIOTECH LTD.

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SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

SINOVAC BIOTECH LTD.

By: /s/ Weidong Yin

Name: Weidong Yin

Title: Chief Executive Officer, President

Date: September 2, 2009

Sinovac Obtains Production License for H1N1 Vaccine

BEIJING, Sept. 3 /PRNewswire-Asia/ -- Sinovac Biotech Ltd. (NYSE: SVA), a leading provider of biopharmaceutical products in China, announced today that the State Food and Drug Administration (SFDA) has approved the registration application for PANFLU.1, Sinovac's H1N1 vaccine, and has issued Sinovac a production license for this vaccine.

Following top-line results which showed that PANFLU.1, Sinovac's H1N1 vaccine, has a good safety and immunogenicity profile, the Company announced the findings of an experts' evaluation conference organized by the SFDA on August 30 and 31. The experts unanimously agreed that Sinovac's H1N1 vaccine is suitable for all people from three to 60 years old on a single shot vaccination schedule.

Mr. Weidong Yin, Chairman, President and CEO of Sinovac, commented, "The approval of PANFLU.1, Sinovac's H1N1 vaccine, is a significant milestone in the campaign for the prevention and control of the H1N1 virus. With the support of the Ministry of Health, State SFDA, Chinese Center for Disease Control and Prevention (China CDC), Sinovac was able to successfully and rapidly complete the clinical trials and registration process for the H1N1 vaccine. By leveraging our expertise in R&D, production and commercialization of human vaccines, we continue to execute our mission to provide top-quality vaccines to eliminate human diseases."

About Sinovac

Sinovac Biotech Ltd. is a China-based biopharmaceutical company that focuses on the research, development, manufacture and commercialization of vaccines that protect against human infectious diseases. Sinovac's vaccine products include Healive(R) (hepatitis A), Bilive(R) (combined hepatitis A and B), and Anflu(R) (influenza). Panflu(TM), Sinovac's pandemic influenza vaccine (H5N1), has already been approved for government stockpiling. Sinovac is developing vaccines for enterovirus 71, universal pandemic influenza, Japanese encephalitis, and human rabies. Its wholly owned subsidiary, Tangshan Yian, is conducting field trials for independently developed inactivated animal rabies vaccines.

Safe Harbor Statement

This announcement contains forward-looking statements. These statements are made under the "safe harbor" provisions of the U.S. Private Securities Litigation Reform Act of 1995. These forward-looking statements can be identified by words or phrases such as "will," "expects," "anticipates," "future," "intends," "plans," "believes," "estimates" and similar statements. Among other things, the business outlook and quotations from management in this press release contain forward-looking statements. Statements that are not historical facts, including statements about Sinovac's beliefs and expectations, are forward-looking statements. Forward-looking statements involve inherent risks and uncertainties. A number of important factors could cause actual results to differ materially from those contained in any forward-looking statement. Sinovac does not undertake any obligation to update any forward-looking statement, except as required under applicable law.

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