

**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION**

Washington, D.C. 20549

**FORM 8-K**

**Current Report  
Pursuant to Section 13 or 15(d) of the  
Securities Exchange Act of 1934**

Date of Report (Date of earliest event reported): **June 18, 2012**

**BIOSANTE PHARMACEUTICALS, INC.**

(Exact name of registrant as specified in its charter)

**Delaware**  
(State or other jurisdiction  
of incorporation)

**001-31812**  
(Commission File Number)

**58-2301143**  
(I.R.S. Employer  
Identification Number)

**111 Barclay Boulevard**  
**Lincolnshire, Illinois**  
(Address of principal executive offices)

**60069**  
(Zip Code)

Registrant's telephone number, including area code: **(847) 478-0500**

**N/A**  
(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

**Item 8.01. Other Events.**

On June 18, 2012, BioSante Pharmaceuticals, Inc. ("BioSante") received notice from The NASDAQ Stock Market ("NASDAQ") indicating that BioSante regained compliance with the minimum bid price requirement for continued inclusion of its common stock on The NASDAQ Global Market under NASDAQ Listing Rule 5450(a)(1).

**Item 9.01 Financial Statements and Exhibits.**

(d) *Exhibits.*

<u>Exhibit No.</u>	<u>Description</u>
99.1	News Release dated June 19, 2012 (filed herewith)

**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 hereunto duly authorized.

**BIOSANTE PHARMACEUTICALS, INC.**

By: /s/ Phillip B. Donenberg  
Phillip B. Donenberg  
Senior Vice President of Finance,  
Chief Financial Officer and Secretary

Dated: June 19, 2012

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**BIOSANTE PHARMACEUTICALS, INC.**

**FORM 8-K**

**Exhibit Index**

<b>Exhibit No.</b>	<b>Description</b>	<b>Method of Filing</b>
99.1	News Release dated June 19, 2012 (filed herewith)	Filed herewith

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**BioSante Pharmaceuticals, Inc.**  
 111 Barclay Boulevard  
 Lincolnshire, Illinois 60069  
[www.biosantepharma.com](http://www.biosantepharma.com)

**FOR IMMEDIATE RELEASE**

**NASDAQ: BPAX**

**BioSante Pharmaceuticals Regains Compliance with  
 NASDAQ Minimum Bid Price Listing Requirement**

LINCOLNSHIRE, Illinois (June 19, 2012) — BioSante Pharmaceuticals, Inc. (NASDAQ: BPAX) announced today that it has received notice from The NASDAQ Stock Market indicating that BioSante has regained compliance with the minimum bid price requirement for continued inclusion of its common stock on The NASDAQ Global Market.

As previously announced by BioSante on June 11, 2012 the Company plans to initiate two new LibiGel Phase III efficacy trials. This decision is based on an extensive analysis of previous efficacy data, consultation with key opinion leaders (KOLs) in female sexual dysfunction, testosterone therapy and placebo effects, as well as a meeting with the U.S. Food and Drug Administration (FDA). BioSante also intends to continue the on-going LibiGel Phase III cardiovascular and breast cancer safety study as per protocol. LibiGel (testosterone gel) is in development for the treatment of female sexual dysfunction (FSD), specifically, hypoactive sexual desire disorder (HSDD) in menopausal women.

**About BioSante Pharmaceuticals, Inc.**

BioSante is a specialty pharmaceutical company focused on developing products for female sexual health and oncology. BioSante's products include LibiGel® (transdermal testosterone gel) for the treatment of female sexual dysfunction (FSD), specifically hypoactive sexual desire disorder (HSDD), which is in Phase III clinical development. BioSante also is developing a portfolio of cancer vaccines, with 17 Phase I and Phase II clinical trials currently on-going. Four of these vaccines have been granted Orphan Drug designation by the U.S. Food and Drug Administration (FDA). BioSante's other products include an FDA-approved testosterone gel for male hypogonadism, which is licensed to Teva Pharmaceuticals USA, Inc., and the Pill-Plus™, an oral contraceptive in Phase II clinical development by Pantarhei Bioscience B.V. BioSante's first FDA-approved product, Elestrin™ (estradiol gel) indicated for the treatment of hot flashes associated with menopause, is marketed in the U.S. by Jazz Pharmaceuticals, BioSante's licensee. Additional information is available online at: [www.biosantepharma.com](http://www.biosantepharma.com).

**Forward-Looking Statements**

*To the extent any statements made in this release deal with information that is not historical, these are forward-looking statements under the Private Securities Litigation Reform Act of 1995. Such statements include, but are not limited to, statements about the continued listing of BioSante's common stock on The NASDAQ Global Market and other statements identified by words such as "plans," "intends," "may," "could," other words of similar meaning, derivations of such words and the use of future dates. Forward-looking statements by their nature address matters that are, to different degrees, uncertain. Uncertainties and risks may cause BioSante's actual results to be materially different than those expressed in or implied by BioSante's forward-looking statements. For BioSante, particular uncertainties and risks include, among others, risks and uncertainties related to the market price of BioSante's securities and its continued listing on The NASDAQ Global Market, uncertainties regarding clinical testing, the difficulty of developing pharmaceutical products, obtaining regulatory and other approvals and achieving market acceptance; the marketing and other success of BioSante's licensees or sublicensees and BioSante's future revenues, if any, from its licensees and sublicensees; uncertainties relating to the future and costs of BioSante's product development programs and BioSante's need for and ability to obtain additional financing if needed. More detailed information on these and additional factors that could affect BioSante's actual results are described in BioSante's filings with the Securities and Exchange Commission, including its most recent annual report on Form 10-K and subsequent quarterly report on Form 10-Q. All forward-looking statements in this release speak only as of the date of this release and are based on BioSante's current beliefs and expectations. BioSante undertakes no obligation to update or revise any forward-looking statement, whether as a result of new information, future events or otherwise.*

For more information, please contact:

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