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): August 6, 2009

NABI BIOPHARMACEUTICALS

(Exact Name of Registrant as specified in its charter)

Delaware (State or Other Jurisdiction of Incorporation)

000-04829

(Commission File Number)

59-1212264 (IRS Employer Identification No.)

12276 Wilkins Avenue, Rockville, Maryland 20852 (Address of principal executive offices) (Zip Code)

Registrant's telephone number, including area code: (301) 770-3099

Not Applicable

(Registrant's name or former address, if change 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 (<i>see</i> General Instruction A.2. below):				
	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 2.02. Results of Operations and Financial Condition

On August 6, 2009, Nabi Biopharmaceuticals (the "Company") issued a press release announcing its results of operations for the three and six months ended June 27, 2009. A copy of the press release announcing these results is furnished as Exhibit 99.1 to this report.

The information in this Item 2.02 and the exhibit attached hereto shall not be deemed to be "filed" for purposes of Section 18 of the Securities Exchange Act of 1934 (the "Exchange Act") or otherwise subject to the liability of that section, and it shall not be incorporated by reference into any filing under the Securities Act of 1933 or the Exchange Act, regardless of any general incorporation language in such filing.

Item 8.01. Other Events

On August 6, 2009, the Company issued a press release announcing it had signed a definitive agreement for the sale of PentaStaph (Pentavalent S. aureus Vaccine) and related assets to GlaxoSmithKline. A copy of the press release is furnished as Exhibit 99.2 to this report.

The information furnished in this Item 8.01 and the exhibit attached hereto shall not be deemed to be "filed" for purposes of Section 18 of the Securities Exchange Act of 1934 (the "Exchange Act") or otherwise subject to the liability of that section, and it shall not be incorporated by reference into any filing under the Securities Act of 1933 or the Exchange Act, regardless of any general incorporation language in such filing.

Item 9.01. Financial Statements and Exhibits

Exhibit number	Description
99.1	Earnings Press Release
99.2	PentaStaph Press Release

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.

NABI BIOPHARMACEUTICALS

By: /s/ Raafat E.F. Fahim, Ph.D.

Raafat E.F. Fahim, Ph.D.

President and Chief Executive Officer

Date: August 6, 2009



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NEWS RELEASE

FOR IMMEDIATE RELEASE

Nabi Biopharmaceuticals Announces Second Quarter 2009 Financial Results

Rockville, Maryland, August 6, 2009 – Nabi Biopharmaceuticals (NASDAQ: NABI) today announced its second quarter financial results for the three month period ended June 27, 2009. The Company reported a net loss from continuing operations of \$5.8 million, or \$0.11 per share, compared to a net loss of \$5.7 million, or \$0.11 per share, for the period ended June 28, 2008. Including results from discontinued operations, Nabi had a net loss of \$5.8 million, or \$0.11 per share, for the current period compared to a net loss of \$3.7 million, or \$0.07 per share in the second quarter of 2008. The 2008 results have been adjusted to reflect the retrospective application of FSP 14-1 accounting rules for convertible debt instruments that were adopted in the first quarter of 2009.

General and Administrative expenses were \$2.4 million for the quarter ended June 27, 2009 compared to \$2.9 million in the prior year period. The decrease reflects our ongoing efforts to reduce overall infrastructure costs as well as a reduction in stock-based compensation expense. Research and Development expenses were \$3.4 million in the second quarter of 2009 compared to \$3.3 million in 2008, reflecting increased efforts to prepare for the planned Phase III NicVAX trial including manufacturing-related activities.

For the six months ended June 27, 2009, the Company's net loss from continuing operations was \$12.9 million, or \$0.25 cents per share, compared to \$13.3 million, or \$0.25 cents per share, for the six months ended June 28, 2008. Including results from discontinued operations, the net loss was \$12.9 million, or \$0.25 per share, compared to a net loss of \$11.0 million, or \$0.21 per share, in 2008. General and Administrative expenses for the current six-month period were \$5.4 million compared to \$8.1 million in 2008 while Research and Development expense was \$7.2 million for the current six-month period compared to \$6.5 million for the comparable 2008 period. The 2008 results have been adjusted to reflect the retrospective application of FSP 14-1 accounting rules for convertible debt instruments that were adopted in the first quarter of 2009.

Net cash used in operating activities from continuing operations was \$12.8 million for the first six months of 2009 compared to \$13.1 million used in the first six months of 2008. Cash, cash equivalents and marketable securities totaled \$108.7 million at June 27, 2009, excluding \$5.7 million of restricted cash related to discontinued operations. The reduction of \$21.6 million from the year-end 2008 balance of \$130.3 million includes \$12.8 million of cash used in continuing operating activities and \$13.6 million in repurchases of company stock and convertible debentures, partially offset by the release of \$4.5 million of restricted cash held in escrow.

During the second quarter, the Company repurchased \$10.4 million, par value, of its 2.875% convertible senior notes for \$10.1 million, leaving a balance of just \$6 million. The Company also repurchased 937,255 shares of its common stock at an average price of \$2.91 per share. A total of 11.1 million shares have been repurchased since the inception of the program.

On March 31, 2009 Biotest Pharmaceuticals Corporation made indemnification claims against Nabi totaling \$56 million. On June 1, 2009 Nabi announced that Biotest had withdrawn \$50.4 million of these claims, leaving a remaining claim of \$5.7 million, for which cash in that amount is being held in escrow pending resolution. We believe this remaining claim is unsubstantiated and are seeking the release of the remaining escrow funds.

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On June 3, 2009 Nabi announced that it had received scientific advice from the European Medicines Agency (EMEA) on NicVAX regarding the appropriate design of the pivotal phase III clinical studies and safety data base. The scientific advice is designed by EMEA to help improve the chances of a positive marketing approval in Europe if the clinical trial results are positive.

"The scientific advice we received from the European Medicines Agency confirms and supports our NicVAX Phase III trial design and protocol that was previously endorsed by the U.S. FDA through a Special Protocol Assessment. We now have support and guidance from the world's leading regulatory agencies," said Dr. Raafat Fahim, President and Chief Executive Officer of Nabi Biopharmaceuticals. "We are closely controlling costs, have significantly reduced our debt level and eliminated the potential risk from a large indemnification claim. Our announcement earlier today of the PentaStaph sale validates the interest in Nabi and its products. We continue to have interest from and discussions with potential strategic partners for our remaining assets as we pursue our strategic alternatives process."

Financial Results Conference Call and Webcast Information

The Company will host a live webcast at 4:30 p.m. EDT today to discuss these results.

The live webcast can be accessed at:

http://phx.corporate-ir.net/phoenix.zhtml?p=irol-eventDetails&c=100445&eventID=2332831

(Due to the length of this URL, it may be necessary to copy and paste this hyperlink into your browser. Remove the space if one exists.) or via the Nabi Biopharmaceuticals website at http://www.nabi.com.

If you do not have Internet access, the U.S./Canada call-in number is 866-804-6922 and the international call-in number is 857-350-1668. The passcode is 70728172. An audio replay will be available for U.S./Canada callers at 888-286-8010 and for international callers at 617-801-6888. The replay passcode is 25229128. This audio replay will be available through August 13, 2009. An archived version of the webcast will also be available on the company's website at http://www.nabi.com.

About Nabi Biopharmaceuticals

Nabi Biopharmaceuticals leverages its experience and knowledge in powering the immune system to develop products that target serious medical conditions in the areas of nicotine addiction and gram-positive bacterial infections. Nabi Biopharmaceuticals is currently developing NicVAX® (Nicotine Conjugate Vaccine), an innovative and proprietary investigational vaccine for treatment of nicotine addiction and prevention of smoking relapse, and PentaStaphTM (Pentavalent S. aureus Vaccine), a vaccine designed to prevent the most dangerous and prevalent strains of *S. aureus* bacterial infections. The company is headquartered in Rockville, Maryland. For additional information about Nabi Biopharmaceuticals, please visit our Web site:http://www.nabi.com.

Forward-Looking Statements

Statements in this release that are not strictly historical are forward-looking statements and include statements about products in development, results and analyses of clinical trials and studies, research and development expenses, cash expenditures, licensure applications and approvals, and alliances and partnerships, among other matters. You can identify these forward-looking statements because they involve our expectations, intentions, beliefs, plans, projections, anticipations, or other characterizations of future events or circumstances. These forward-looking statements are not guarantees of future performance and are subject to risks and uncertainties that may cause actual results to differ materially from those in the forward-looking statements as a result of any number of factors. These factors include, but are not limited to, risks relating to our ability to: successfully close the sale of PentaStaph and complete the PentaStaph sale milestones; partner with third parties to fund, develop, manufacture and/or commercialize our products in development; defend against indemnification claims by Biotest; initiate and conduct clinical trials and studies; raise sufficient new capital resources to fully develop and commercialize our products in development; attract, retain and motivate key employees; collect further milestone and royalty payments under the PhosLo Agreement; obtain regulatory approval for our products in the U.S. or other markets; successfully contract with third party manufacturers for the manufacture and supply of NicVAX and PentaStaph; and comply with reporting and payment obligations under government rebate and pricing programs. Some of these factors are more fully discussed, as are other factors, in our Annual Report on Form 10-K for the fiscal year ended December 27, 2008 and our Quarterly Report on Form 10-Q for the period ended March 28, 2009 filed with the Securities and Exchange Commission.

Nabi Biopharmaceuticals CONDENSED CONSOLIDATED BALANCE SHEETS (Unaudited) (In thousands)

	June 27, 	December 27, 2008 (as adjusted)
ASSETS		
Current assets:		
Cash and cash equivalents	\$ 107,658	\$ 106,438
Marketable securities	1,034	23,900
Prepaid expenses and other current assets	2,121	1,430
Assets of discontinued operations (including restricted cash)	5,888	10,409
Total current assets	116,701	142,177
Property and equipment, net	1,083	1,315
Other assets	383	730
Total assets	\$ 118,167	\$ 144,222
LIABILITIES AND STOCKHOLDERS' EQUITY		
Current liabilities:		
Accounts payable	\$ 1,333	\$ 1,226
Accrued expenses and other current liabilities	1,555	3,030
Current liabilities of discontinued operations	3,350	3,381
Total current liabilities	6,238	7,637
2.875% convertible senior notes, net	5,775	15,202
Total liabilities	12,013	22,839
Commitments and contingencies		
Stockholders' equity:		
Convertible preferred stock	_	_
Common stock	6,271	6,239
Capital in excess of par value	363,759	363,001
Treasury stock	(45,321)	(42,187)
Other comprehensive income (loss)	(2)	60
Accumulated deficit	(218,553)	(205,730)
Total stockholders' equity	106,154	121,383
Total liabilities and stockholders' equity	\$ 118,167	\$ 144,222

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Nabi Biopharmaceuticals CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS (Unaudited)

(In thousands, except share and per share data)

	For the Three Months Ended		For the Six Months Ended	
	June 27, 2009	June 28, 2008 (as adjusted)	June 27, 2009	June 28, 2008 (as adjusted)
Operating expenses:				
General and administrative expenses	\$ 2,355	\$ 2,927	\$ 5,445	\$ 8,060
Research and development expenses	3,440	3,344	7,206	6,549
Operating loss	(5,795)	(6,271)	(12,651)	(14,609)
Interest income	83	1,217	270	3,255
Interest expense	(136)	(1,118)	(497)	(2,670)
Other income (expense), net	40	(853)	24	(722)
Loss from continuing operations before income taxes	(5,808)	(7,025)	(12,854)	(14,746)
Benefit from income taxes	_	1,300	_	1,495
Loss from continuing operations	(5,808)	(5,725)	(12,854)	(13,251)
Discontinued operations:				
Income from discontinued operations, net of tax provision	_	1,996	_	2,295
Income from discontinued operations		1,996		2,295
Net loss	\$ (5,808)	\$ (3,729)	\$(12,854)	\$ (10,956)
Basic and diluted (loss) income per share:				
Continuing operations	\$ (0.11)	\$ (0.11)	\$ (0.25)	\$ (0.25)
Discontinued operations	0.00	0.04	0.00	0.04
Basic and diluted (loss) income per share	\$ (0.11)	\$ (0.07)	\$ (0.25)	\$ (0.21)
Basic and diluted weighted average shares outstanding	50,974	51,498	51,094	52,235

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Nabi Biopharmaceuticals CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS (Unaudited) (In thousands)

	For the Six Months Ended	
	June 27, 2009	June 28, 2008 (as adjusted)
Cash flow from operating activities:		
Loss from continuing operations	\$ (12,854)	\$ (13,251)
Adjustments to reconcile loss from continuing operations to net cash used in operating activities from continuing operations:		
Depreciation and amortization	255	291
Non-cash intra-period tax allocation	_	(1,495)
Accretion of discount on convertible senior notes	306	1,555
Non-cash compensation	904	1,943
Other	6	945
Changes in assets and liabilities:		
Prepaid expenses and other assets	(426)	747
Accounts payable, accrued expenses and other	(1,035)	(3,822)
Total adjustments	10	164
Net cash used in operating activities from continuing operations	(12,844)	(13,087)
Net cash provided by (used in) operating activities from discontinued operations	4,488	(517)
Net cash used in operating activities	(8,356)	(13,604)
Cash flow from investing activities:		
Proceeds from sales and maturities of marketable securities, net	22,836	1,600
Capital expenditures	_	(20)
Other	_	91
Net cash provided by investing activities from continuing operations	22,836	1,671
Net cash provided by investing activities from discontinued operations	_	2,500
Net cash provided by investing activities	22,836	4,171
Cash flow from financing activities:		
Proceeds from issuances of common stock for employee benefit plans	297	54
Purchase of common stock for treasury	(3,466)	(18,658)
Other financing activities	(10,091)	(28,996)
Net cash used in financing activities from continuing operations	(13,260)	(47,600)
Net cash used in financing activities from discontinued operations	_	(23)
Net cash used in financing activities	(13,260)	(47,623)
Net increase (decrease) in cash and cash equivalents	1,220	(57,056)
Cash and cash equivalents at beginning of period		217,606
Cash and cash equivalents at end of period	\$107,658	\$ 160,550



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FOR IMMEDIATE RELEASE

Nabi Biopharmaceuticals Announces Sale of PentaStaph™ to GlaxoSmithKline PLC \$46 million Total Transaction Value

Rockville, Maryland, August 6, 2009 − Nabi Biopharmaceuticals (NASDAQ: NABI) announced that it has signed a definitive agreement for the sale of PentaStaph[™] (Pentavalent *S. aureus* Vaccine) and related assets to GlaxoSmithKline (GSK) for a total consideration of up to \$46 million. Under the terms of the agreement, Nabi will receive an initial cash payment of \$20 million when the transaction closes plus an additional \$26 million contingent upon four milestone accomplishments. The sale is subject to satisfaction of certain customary closing conditions by the buyer and seller prior to closing. Nabi anticipates closing the sale before year-end 2009 and completing the milestones within the next 18 months.

Key opinion leaders believe that successful development of a vaccine against *S. aureus* is critical in the fight against this multidrug-resistant bacterium. MRSA, or Methicillin-resistant *S. aureus*, is an increasingly prevalent strain of the bacterium. MRSA is especially troublesome in hospital-associated nosocomial infections where patients with open wounds, invasive devices and those undergoing surgery are at a greater risk of infection. MRSA is also occurring in otherwise healthy people who have not been recently hospitalized and usually manifests as skin infections through cuts, pimples and boils.

"We are very pleased that GSK is acquiring the PentaStaph program and undertaking its further development and commercialization in this area of high unmet medical need. This transaction is the next step in the execution of our strategic alternatives process and we are pleased to be able to generate near-term value from this program for our shareholders," said Dr. Raafat Fahim, President and Chief Executive Officer of Nabi Biopharmaceuticals. "We believe in the potential of PentaStaph and are confident that GSK will continue to develop this vaccine into a successful weapon against the growing threat from Staphylococcus infection."

About PentaStaph

PentaStaphTM (Pentavalent *S. aureus* Vaccine) is Nabi's five-component vaccine candidate against *S. aureus* infections. Three of the antigen components induce the production of antibodies that target *S. aureus* polysaccharides (Types 5 & 8 capsular polysaccharides and Type 336 cell-wall polysaccharide) which enhance the immune systems' ability to clear the bacteria. Types 5 & 8 capsular polysaccharides are expressed in approximately 80 percent of *S. aureus* strains, including many of the known MRSA (methicillin-resistant *S. aureus*) strains. Type 336 polysaccharide accounts for approximately 20% of *S. aureus* infections that do not form a polysaccharide capsule in the human bloodstream. Two additional antigen components induce the production of antibodies that target two of the most virulent toxins produced by the bacteria which can significantly debilitate the human immune system: Panton-Valentine Leukocidin found predominantly in community-acquired MRSA, and alpha toxin, produced by almost all *S. aureus* isolates. This multi-target approach which enhances the immune system's ability to eliminate a broad spectrum of *S. aureus* strains and neutralizes the bacterial defenses of the most virulent strains has been demonstrated in pre-clinical animal models to provide significant efficacy against the bacteria which is hoped would translate into human efficacy.

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