

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) September 20, 2012

NAVIDEA BIOPHARMACEUTICALS, INC.

(Exact name of registrant as specified in its charter)

<u>Delaware</u> (State or other jurisdiction of incorporation)	<u>001-35076</u> (Commission File Number)	<u>31-1080091</u> (IRS Employer Identification No.)
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<u>425 Metro Place North, Suite 450, Dublin, Ohio</u> (Address of principal executive offices)	<u>43017</u> (Zip Code)
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Registrant's telephone number, including area code (614) 793-7500

(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 (see 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 8.01. Other Events.**

On September 20, 2012, Navidea Biopharmaceuticals, Inc. issued a press release announcing that it has received a Small Business Innovation Research (SBIR) grant from the National Cancer Institute (NCI), National Institutes of Health (NIH), to fund the development of the Company's radio-immuno-guided surgery monoclonal antibody targeting agent for use in detecting metastatic sites in colorectal cancer. The SBIR grant has the potential for grant money up to a total of \$1.5M over three years if fully funded. The first-year Phase I funding of \$315,000, which has already been approved, is expected to enable Navidea to complete preclinical bridging activities using a RIGS tumor-antigen-targeted monoclonal antibody and prepare a standardization clinical trial protocol. Second and third year (Phase II) funding of up to \$1.2M is contingent upon meeting certain Phase I success criteria, including Institutional Review Board (IRB) approval of the clinical trial protocol.

A copy of the complete text of the Company's September 20, 2012, press releases is attached as Exhibits 99.1 to this Current Report on Form 8-K and is incorporated herein by reference.

Statements contained or incorporated by reference in this Current Report on Form 8-K which relate to other than strictly historical facts, such as statements about the Company's plans and strategies, expectations for future financial performance, new and existing products and technologies, anticipated clinical and regulatory pathways and markets for the Company's products, are forward-looking statements. The words "believe," "expect," "anticipate," "estimate," "project," and similar expressions identify forward-looking statements that speak only as of the date hereof. Investors are cautioned that such statements involve risks and uncertainties that could cause actual results to differ materially from historical or anticipated results due to many factors including, but not limited to, the Company's continuing operating losses, uncertainty of market acceptance of its products, reliance on third party manufacturers, accumulated deficit, future capital needs, uncertainty of capital funding, dependence on limited product line and distribution channels, competition, limited marketing and manufacturing experience, risks of development of new products, regulatory risks and other risks detailed in the Company's most recent Annual Report on Form 10-K and other Securities and Exchange Commission filings. The Company undertakes no obligation to publicly update or revise any forward-looking statements.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits.

*Exhibit  
Number*

*Exhibit Description*

99.1	Navidea Biopharmaceuticals, Inc. press release, dated September 20, 2012, entitled "Navidea Awarded SBIR Grant from the NIH for Development of Radio-Immuno-Guided Surgery (RIGS) Agent Aimed at Detecting Metastatic Cancer."
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**SIGNATURES**

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.

Navidea Biopharmaceuticals, Inc.

Date: September 20, 2012

By: /s/ Brent L. Larson  
Brent L. Larson, Senior Vice President and  
Chief Financial Officer



Press Release

## **Navidea Awarded SBIR Grant from the NIH for Development of Radio-Immuno-Guided Surgery (RIGS) Agent Aimed at Detecting Metastatic Cancer**

DUBLIN OHIO, September 20, 2012 – Navidea Biopharmaceuticals, Inc. (NYSE MKT: NAVB), a biopharmaceutical company focused on precision diagnostic radiopharmaceuticals, today announced the receipt of a Small Business Innovation Research (SBIR) grant from the National Cancer Institute (NCI), National Institutes of Health (NIH), to fund the development of the Company's radio-immuno-guided surgery monoclonal antibody targeting agent for use in detecting metastatic sites in colorectal cancer. The SBIR grant has the potential for grant money up to a total of \$1.5M over three years if fully funded. The first-year Phase I funding of \$315,000, which has already been approved, is expected to enable Navidea to complete preclinical bridging activities using a RIGS tumor-antigen-targeted monoclonal antibody and prepare a standardization clinical trial protocol. Second and third year (Phase II) funding of up to \$1.2M is contingent upon meeting certain Phase I success criteria, including Institutional Review Board (IRB) approval of the clinical trial protocol.

"We are pleased that NIH/NCI has recognized the potential value for Navidea's RIGS agent to significantly benefit surgeons in effectively locating cancerous tissues during surgery and enabling their more thorough surgical removal for better patient outcomes. We appreciate NCI's support as we complete these important preclinical studies and prepare for re-initiation of human clinical trials," said Mark Pykett, V.M.D., Ph.D., President and CEO of Navidea. "The development of our RIGS monoclonal antibody agent represents another clinical advancement of Navidea's pipeline of innovative precision diagnostic products, including Lymphoseek<sup>®</sup> and NAV4694, which target devastating oncologic and neurologic diseases."

### **About the RIGS Monoclonal Antibody Targeting Agent**

The RIGS targeting agent is an investigational, tumor-specific, radio-labeled monoclonal antibody. It may be used during surgery to identify cancerous tissue undetectable by traditional diagnostic and intraoperative techniques. The RIGS monoclonal antibody agent may enable more effective surgeries caused by metastasized colorectal cancers leading to improved patient survival.

Before surgery, a cancer patient is injected with the RIGS targeting agent which circulates throughout the patient's body and binds specifically to cancer cell antigens or receptors. Concentrations of the RIGS targeting agent within affected tissue are then detected using a gamma probe and direct the surgeon to targeted tissue for removal.

### **About Colorectal Cancer and Metastatic Disease**

With nearly 140,000 new diagnoses and 50,000 deaths each year in the U.S., adenocarcinomas of the colon and rectum are a common and deadly form of cancer. When patients are first diagnosed with these diseases, they undergo surgeries to remove their tumors. Colon and rectal (colorectal) cancers, like nearly all forms of cancer, are insidious in that they can spread or metastasize from their primary sites of origin to multiple locations throughout the body. Patients who eventually succumb to colorectal cancer are killed by the uncontrolled growth of these metastatic tumors. It is well established that for colorectal cancers the most common site to which the cancer will spread first is the liver. Frequently, it is still possible to cure or dramatically extend the life of a patient with liver metastases using surgery if their cancer has not yet spread beyond the liver. About one third of colorectal cancer patients undergo surgeries to remove liver metastases.

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**NAVIDEA BIOPHARMACEUTICALS**  
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**About Navidea Biopharmaceuticals, Inc.**

Navidea Biopharmaceuticals, Inc. (NYSE MKT: NAVB) is a biopharmaceutical company focused on the development and commercialization of precision diagnostics and radiopharmaceutical agents. Navidea is actively developing four radiopharmaceutical agent platforms – Lymphoseek<sup>®</sup>, NAV4694, CFT and RIGScan<sup>™</sup> – to help identify the sites and pathways of undetected disease and enable better diagnostic accuracy, clinical decision-making and, ultimately, patient care. Navidea’s strategy is to deliver superior growth and shareholder return by bringing to market novel radiopharmaceutical agents and advancing the Company’s pipeline through selective acquisitions, global partnering and commercialization efforts. For more information, please visit [www.navidea.com](http://www.navidea.com).

*The Private Securities Litigation Reform Act of 1995 (the Act) provides a safe harbor for forward-looking statements made by or on behalf of the Company. Statements in this news release, which relate to other than strictly historical facts, such as statements about the Company’s plans and strategies, expectations for future financial performance, new and existing products and technologies, anticipated clinical and regulatory pathways, and markets for the Company’s products are forward-looking statements within the meaning of the Act. The words “believe,” “expect,” “anticipate,” “estimate,” “project,” and similar expressions identify forward-looking statements that speak only as of the date hereof. Investors are cautioned that such statements involve risks and uncertainties that could cause actual results to differ materially from historical or anticipated results due to many factors including, but not limited to, the Company’s continuing operating losses, uncertainty of market acceptance of its products, reliance on third party manufacturers, accumulated deficit, future capital needs, uncertainty of capital funding, dependence on limited product line and distribution channels, competition, limited marketing and manufacturing experience, risks of development of new products, regulatory risks and other risks detailed in the Company’s most recent Annual Report on Form 10-K and other Securities and Exchange Commission filings. The Company undertakes no obligation to publicly update or revise any forward-looking statements.*

**Contact:**

Navidea Biopharmaceuticals, Inc. – Brent Larson, Sr. VP & CFO – (614) 822-2330

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