BIOENVISION INC
Form DEFA14A
October 05, 2007
UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

SCHEDULE 14A

Proxy Statement Pursuant to Section 14(a) of
the Securities Exchange Act of 1934 (Amendment No.)

Filed by the Registrant x
Filed by a Party other than the Registrant o
Check the appropriate box:

o	Preliminary Proxy Statement
o	**Confidential, for Use of the Commission Only (as permitted by Rule 14a-6(e)(2))**
o	Definitive Proxy Statement
x	Definitive Additional Materials
o	Soliciting Material Pursuant to §240.14a-12

BIOENVISION, INC.
(Name of Registrant as Specified In Its Charter)

(Name of Person(s) Filing Proxy Statement, if other than the Registrant)

Payment of Filing Fee (Check the appropriate box):

x No fee required.

o Fee computed on table below per Exchange Act Rules 14a-6(i)(1) and 0-11.

(1) Title of each class of securities to which transaction applies:

(2) Aggregate number of securities to which transaction applies:

(3) Per unit price or other underlying value of transaction computed pursuant to Exchange Act Rule 0-11 (set forth the amount on which the filing fee is calculated and state how it was determined):

(4) Proposed maximum aggregate value of transaction:

(5) Total fee paid:

o Fee paid previously with preliminary materials.

o Check box if any part of the fee is offset as provided by Exchange Act Rule 0-11(a)(2) and identify the filing for which the offsetting fee was paid previously. Identify the previous filing by registration statement number, or the Form or Schedule and the date of its filing.

(1) Amount Previously Paid:

(2) Form, Schedule or Registration Statement No.:

(3) Filing Party:

(4) Date Filed:

This filing consists of a press release issued by Bioenvision, Inc.

ADDITIONAL INFORMATION AND WHERE TO FIND IT

In connection with the proposed acquisition of Bioenvision, Inc. (Bioenvision) by Genzyme Corporation (Genzyme) and the required approval of the transaction by Bioenvision s stockholders, Bioenvision filed a definitive proxy statement and other relevant documents concerning the transaction with the Securities and Exchange Commission (SEC) on September 7, 2007. Stockholders of Bioenvision are urged to read the definitive proxy statement and any other relevant documents because they contain important information. Investors and security holders can obtain free copies of the definitive proxy statement and other relevant documents when they become available by contacting Bioenvision Investor Relations at (212) 750-6700 ext. 160. In addition, documents filed with the SEC by both Genzyme and Bioenvision are available free of charge at the SEC s web site at http://www.sec.gov.

Information regarding the identity of the persons who may, under SEC rules, be deemed to be participants in the solicitation of stockholders of Bioenvision in connection with the transaction, and their interests in the solicitation, is set forth in the proxy materials filed by Bioenvision with the SEC.

FORWARD-LOOKING STATEMENTS

Certain statements contained in the press release are forward-looking statements, including express or implied statements regarding the future approval by Bioenvision s stockholders of the pending agreement and plan of merger with Genzyme and regarding Bioenvision obtaining regulatory approval of its products. Because these statements are subject to risks and uncertainties, actual results may differ materially from those expressed or implied by such forward-looking statements. Specifically, factors that could cause actual results to differ materially from those expressed or implied by such forward-looking statements include, but are not limited to: risks associated with whether the merger of Wichita Bio Corporation with and into Bioenvision will be approved by the stockholders of Bioenvision; risks associated with the uncertainty as to whether such merger will in fact occur, risks associated with disruptions from the proposed merger transaction which may harm relationships with customers, employees, suppliers and partners; risks associated with the outcome of litigation and regulatory proceedings to which we are currently a party and may become a party in the future; risks associated with preclinical and clinical developments in the biopharmaceutical industry in general and in Bioenvision s compounds under development in particular; the potential failure of Bioenvision s compounds under development to prove safe and effective for treatment of disease; uncertainties inherent in the early stage of Bioenvision s compounds under development; failure to successfully implement or complete clinical trials; failure to receive marketing clearance from regulatory agencies for our compounds under development; acquisitions, divestitures, mergers, licenses or strategic initiatives that change Bioenvision s business, structure or projections; the development of competing products; uncertainties related to Bioenvision s dependence on third parties and partners; and those risks described in Bioenvision s filings with the SEC. Bioenvision assumes no obligation to update any forward-looking statements as a result of new information or future events or developments, except as required by law and the statements contained in the press release are current as of the date hereof only.

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For Immediate Release

Bioenvision Adjourns Special Meeting of Stockholders
- Inspector of Elections Tabulating Final Vote, Meeting to Reconvene October 10 -

New York, NY - (October 5, 2007) - Bioenvision, Inc. (NASDAQ: BIVN) announced today that at its reconvened special meeting of stockholders, a vote was held regarding the approval of the Agreement and Plan of Merger, dated May 29, 2007, between Bioenvision and Genzyme Corporation, as amended by Amendment No. 1 thereto, dated August 8, 2007 (the Merger Agreement). After the vote was taken, the special meeting of stockholders was adjourned so that the inspector of elections could have additional time to tabulate the final vote with respect to approval of the Merger Agreement. The special meeting of stockholders is scheduled to reconvene on Wednesday, October 10 at 11 a.m. local time at the offices of Goodwin Procter LLP, at 599 Lexington Avenue in New York.

Additional Information and Where to Find It
In connection with the proposed transaction, Bioenvision has filed a definitive proxy statement, a proxy supplement and other materials with the Securities and Exchange Commission (the SEC). We urge investor to read the proxy materials carefully, as they contain important information about Bioenvision and the proposed Merger Agreement. Investors can obtain free copies of the definitive proxy statement as well as other filed documents containing information about Bioenvision at http://www.sec.gov, the SEC s free internet site. These filings are also accessible in the Investors section of the company s website at http://www.bioenvision.com.

About Bioenvision, Inc.

Bioenvision s primary focus is the acquisition, development, and marketing of compounds and technologies for the treatment of cancer. Bioenvision s product pipeline is focused on: Evoltra® (clofarabine) and Modrenal®. For more information on Bioenvision please visit our website at www.bioenvision.com.

Certain statements contained in this press release are forward-looking statements, including express or implied statements regarding the future approval by Bioenvision s stockholders of the pending agreement and plan of merger with Genzyme. Because these statements are subject to risks and uncertainties, actual results may differ materially from those expressed or implied by such forward-looking statements. Specifically, factors that could cause actual results to differ materially from those expressed or implied by such forward-looking statements include, but are not limited to: risks associated with whether the merger of Wichita Bio Corporation with and into Bioenvision will be approved by the stockholders of Bioenvision; risks associated with the uncertainty as to whether such merger will in fact occur, risks associated with disruptions from the proposed merger transaction which may harm relationships with customers, employees, suppliers and partners; risks associated with the outcome of litigation and regulatory proceedings to which we are currently a party and may become a party in the future; risks associated with preclinical and clinical developments in the biopharmaceutical industry in general and in Bioenvision s compounds under development in particular; the potential failure of Bioenvision s compounds under development to prove safe and effective for treatment of disease; uncertainties inherent in the early stage of Bioenvision s compounds under development; failure to successfully implement or complete clinical trials; failure to receive marketing clearance from regulatory agencies for our compounds under development; acquisitions, divestitures, mergers, licenses or strategic initiatives that change Bioenvision s business, structure or projections; the development of competing products; uncertainties related to Bioenvision s dependence on third parties and partners; and those risks described in Bioenvision s filings with the SEC. Bioenvision assumes no obligation to update any forward-looking statements as a result of new information or future events or developments, except as required by law and the statements contained in this press release are current as of the date of this release only.

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