

HEMISPHERX BIOPHARMA INC
Form DEFA14A
September 19, 2011

SCHEDULE 14A
(Rule 14a-101)
INFORMATION REQUIRED IN PROXY STATEMENT
SCHEDULE 14A INFORMATION
Proxy Statement Pursuant to Section 14(a)
of the Securities Exchange Act of 1934

Filed by the Registrant

Filed by a Party other than the Registrant

Check the appropriate box:

- Preliminary Proxy Statement
- Confidential, for Use of the Commission Only (as Permitted by Rule 14a-6(e)(2))
- Definitive Proxy Statement
- Definitive Additional Materials
- Solicitation Material Pursuant to Rule 14a-11(c) or rule 14a-12

Hemispherx Biopharma, 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):

- No fee required.
- Fee computed on table below per Exchange Act Rules 14a-6(i)(1) and 0-11.
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| 1) | Title of each class of securities to which transaction applies: |
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| 3) | Per unit price or other underlying value of transaction computed pursuant to Exchange Act Rule 0-11: |
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- Fee paid previously with preliminary materials.

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(1) Amount Previously Paid:

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

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(4)

Date Filed:

Company/Investor Contact:
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Hemispherx Biopharma's 2011 Stockholder Annual Meeting Set

Philadelphia, PA – September 19, 2011: Hemispherx Biopharma, Inc. (NYSE Amex: HEB) (the “Company”) will hold its Annual Meeting of stockholders on Thursday, October 13, 2011, at 10 a.m. EDT, in Philadelphia, PA. The meeting will be held at the Embassy Suites Hotel, 1776 Benjamin Franklin Parkway, Philadelphia, PA, 19103.

The record date for determining the stockholders entitled to notice of, and to vote at, the Annual Meeting has been set as the close of business on August 18, 2011.

Proposals for action by stockholders include: the election of five Directors; the ratification of McGladrey & Pullen, LLP to audit the financial statements of Hemispherx; the approval, by non-binding vote, of executive compensation; the amendment of the Company's Certificate of Incorporation to increase the number of authorized shares of Common Stock but with restrictions on the use of those shares when such use is for fund-raising purposes; and the transaction of such other matters as may properly come before the meeting or any adjournment thereof.

The mailing of the proxy material commenced on September 16, 2011. The Company has posted copies of the proxy statement, the annual report for the fiscal year ended 2010 and the quarterly report for the quarter ended June 30, 2011 on its website at <http://www.hemispherx.net/content/investor/annualMeeting.asp>.

The Company emphasizes the importance of your vote!

If you are a U.S. resident and received a proxy card containing your Control Number, please take a moment to vote your shares via Internet, phone, or mail. If you are a non-U.S. stockholder, you are encouraged to contact your custodian bank/broker at your earliest convenience.

Important Information for Foreign Stockholders

Hemispherx urges all stockholders who owned shares on August 18, 2011, the Record Date for the Meeting, to vote. A significant number of stockholders are domiciled in Europe and are less readily accessible for notification purposes.

European banks and brokerage houses do not necessarily forward the Proxy materials to stockholders. Accordingly, if you are a European stockholder, you most likely will need to contact your bank or brokerage house directly in order to exercise your right to vote. As we are a Delaware corporation, there is no need for your bank or brokerage house to block your shares. Banks and brokerage houses simply need to certify the number of shares owned by their clients on August 18, 2011 and cast votes by October 11, 2011 (7 pm US Eastern Daylight Time).

We have posted a copy of the proxy statement and related documents on our website at <http://www.hemispherx.net/content/investor/annualMeeting.asp>. For non-U.S. stockholders, we also have posted blank proxy cards in English, French, German and Dutch which they may use to instruct their bank or brokerage house to vote on their behalf.

If you need assistance in voting your shares, Hemispherx suggests that you contact Morrow & Co., LLC, its proxy solicitation agent. Morrow & Co. can be reached in the U.S. at 203-658-9400 or in London at +44-207-222-4645. Stockholders may also contact Dianne Will, Investor Relations for Hemispherx, collect at 518-398-6222 or via e-mail at ir@hemispherx.net.

About Hemispherx Biopharma

Hemispherx Biopharma, Inc. is an advanced specialty pharmaceutical company engaged in the manufacture and clinical development of new drug entities for treatment of seriously debilitating disorders. Hemispherx's flagship products include Alferon N Injection® (FDA approved for a category of sexually transmitted diseases) and the experimental therapeutics Ampligen® and Alferon® LDO. Ampligen® is an experimental RNA nucleic acid being developed for globally important debilitating diseases and disorders of the immune system. Hemispherx's platform technology includes components for potential treatment of various severely debilitating and life threatening diseases. Hemispherx has patents comprising its core intellectual property estate and a fully commercialized product (Alferon N Injection®). The Company wholly owns and exclusively operates a GMP certified manufacturing facility in the United States for commercial products. For more information please visit www.hemispherx.net.

Information contained in this news release, other than historical information, should be considered forward-looking and is subject to various risk factors and uncertainties. For instance, the strategies and operations of Hemispherx involve risk of competition, changing market conditions, change in laws and regulations affecting these industries and numerous other factors discussed in this release and in the Company's filings with the Securities and Exchange Commission. Any specifically referenced investigational drugs and associated technologies of the Company (including Ampligen® and Alferon® LDO) are experimental in nature and as such are not designated safe and effective by a regulatory authority for general use and are legally available only through clinical trials with the referenced disorders. The forward-looking statements represent the Company's judgment as of the date of this release. The Company disclaims, however, any intent or obligation to update these forward-looking statements. Clinical trials for other potential indications of the approved biologic Alferon N Injection® do not imply that the product will ever be specifically approved commercially for these other treatment indications. The planning, completion, results or submission of clinical trials do not imply that any study product will ever be approved commercially for the studied or other treatment indications.
