# 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): March 31, 2009

# AVI BioPharma, Inc.

(Exact name of Company as specified in its charter)

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

**0-22613** (Commission File No.)

93-0797222 (I.R.S. Employer Identification No.)

4575 SW Research Way, Suite 200 Corvallis, OR 97333 (Address of principal executive offices)

(541) 753-3635

Registrant's telephone number, including area code

### Not Applicable

(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))
- o Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

# Item 5.02 Departure of Directors or Certain Officers; Election of Directors; Appointment of Certain Officers; Compensatory Arrangements of Certain Officers

On March 31, 2009, John W. Fara, Ph.D. finalized his decision to not stand for re-election at the end of his term as a director of AVI BioPharma, Inc. ("AVI" or the "Company"). Dr. Fara's decision is based solely on personal reasons, and was not the result of any disagreement with the Company on any matter relating to the Company's operations, policies, or practices.

Also, on March 31, 2009, the Board of Directors of the Company appointed Christopher S. Henney, Ph.D., D. Sc. and M. Kathleen Behrens, Ph.D. as members of the Company's Board of Directors and approved an increase in the number of directors from seven (7) to nine (9).

Dr. Behrens' appointment is being made pursuant to an agreement regarding Board of Director representation between AVI and Eastbourne Capital Management, L.L.C, a 5% or greater shareholder of the Company. The terms of this agreement were disclosed on the Company's Current Report on Form 8-K filed on January 30, 2009.

On March 31, 2009, the Company issued a press release announcing the foregoing changes. A copy of the Company's press release dated March 31, 2009 is attached as Exhibit 99.1 to this Form 8-K and is incorporated herein by reference.

#### Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

The following exhibits are filed herewith:

99.1 Press Release dated March 31, 2009.

## SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) of the Securities Exchange Act of 1934, as amended, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized, in the City of Portland, State of Oregon, on April 1, 2009.	
AVI Bio	oPharma, Inc.
By: <u>/s/</u>	Leslie Hudson, Ph.D.
	slie Hudson, Ph.D. esident and Chief Executive Officer
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Description

Exhibit

Exhibit 99.1

Press Release dated March 31, 2009.

AVI Press and Investor Contact:
Julie Rathbun
Investor Relations
(541) 224-2575
Investorrelations@avibio.com

### AVI BioPharma Appoints Drs. Christopher S. Henney and M. Kathleen Behrens to Its Board of Directors

**PORTLAND, OR** — **March 31, 2009** — AVI BioPharma, Inc. (NASDAQ: AVII), a developer of RNA-based drugs, today announced the appointment of Christopher S. Henney, Ph.D., D. Sc. and M. Kathleen Behrens, Ph.D. to the Company's Board of Directors effective today. The Company also announced that Company Director John Fara, Ph.D. has decided that he will not stand for re-election to the Board at the Company's upcoming annual meeting.

Dr. Henney has been one of the leaders and founders of the U.S. biotechnology industry since 1980. He co-founded three publicly held U.S. biotechnology companies: Immunex Corporation in 1980, later acquired by Amgen; ICOS Corporation in 1990, later acquired by Eli Lilly and Company; and Dendreon Corporation in 1997. At each company, Dr. Henney was a member of the Board of Directors and held a variety of executive management positions. He was CEO and Chairman of Dendreon until 2005. Dr. Henney, a former academic immunologist, held the first Chair in Basic Immunology at the Fred Hutchinson Cancer Center in Seattle. He also held faculty positions at Johns Hopkins University and the University of Washington in the field of immunology and microbiology. Dr. Henney received his Ph.D. in experimental pathology and a D.Sc. for contributions to the field of immunology from the University of Birmingham. Henney is currently chairman of Oncothyreon [ONTY] and Vice-Chair of Cyclacel [CYCC] both NASDAQ listed companies. He advises several early stage companies in the Northwest and in Australia.

Dr. Behrens' career spans the financial services and biotechnology sectors, as well as healthcare policy. Dr. Behrens served as a member of the President's Council of Advisors on Science and Technology (PCAST) from 2001 to early 2009 and she was Chair of PCAST's Subcommittee on Personalized Medicine. She has served as a public-market biotechnology securities analyst as well as venture capitalist focusing on healthcare, technology and related investments. She was instrumental in the founding of several biotechnology companies including Protein Design Labs, Inc. and COR Therapeutics, Inc., and more recently was a Director of Abgenix, Inc., which was acquired by Amgen in 2006. She worked for Robertson Stephens & Co. from 1983 through 1996, serving as a general partner and managing director. Dr. Behrens continued in her capacity as a General Partner for selected venture funds for RS Investments from 1996 through today. From 1997 to 2005, she was a director of the Board on Science, Technology and Economic Policy (STEP) for the National Research Council, and from 1993 to 2000 she was a director, President, Chair and Past Chair of the National Venture Capital Association. Dr. Behrens holds a Ph.D. in Microbiology from the University of California, Davis.

"We are very pleased to have experienced industry leaders of the stsature of Kathy and Chris join our Board of Directors. Their insight and guidance will be invaluable to AVI as we continue to

advance our promising RNA-based drugs," said Leslie Hudson, Ph.D., President and Chief Executive Officer of AVI BioPharma. "The whole Board of AVI is sincerely grateful to John Fara for his long and dedicated service during the Company's transformation from an antisense pioneer to a leading RNA therapeutics company."

### **About AVI BioPharma**

AVI BioPharma is focused on the discovery and development of RNA—based drugs utilizing proprietary derivatives of its antisense chemistry (morpholino-modified phosphorodiamidate oligomers or PMOs) that can be applied to a wide range of diseases and genetic disorders through several distinct mechanisms of action. Unlike other RNA therapeutic approaches, AVI's antisense technology has been used to directly target both messenger RNA (mRNA) and its precursor (pre-mRNA), allowing for both up- and down-regulation of targeted genes and proteins. AVI's RNA—based drug programs are being evaluated for the treatment of Duchenne muscular dystrophy as well as for the treatment of cardiovascular restenosis through our partner Global Therapeutics, a Cook Group Company. AVI's antiviral programs have demonstrated promising outcomes in Ebola Zaire and Marburg Musoke virus infections and may prove applicable to other viral targets such as HCV or Dengue viruses. For more information, visit www.avibio.com.

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"Safe Harbor" Statement under the Private Securities Litigation Reform Act of 1995: The statements that are not historical facts contained in this release are forward-looking statements that involve risks and uncertainties, including, but not limited to, the results of research and development efforts, the results of preclinical and clinical testing, the effect of regulation by the FDA and other agencies, the impact of competitive products, product development, commercialization and technological difficulties, and other risks detailed in the company's Securities and Exchange Commission filings.