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) November 15, 2010

A.P. Pharma, Inc.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation) 001-33221 (Commission File Number) 94-2875566 (I.R.S. Employer Identification No.)

123 Saginaw Drive Redwood City CA (Address of principal executive offices)

94063 (Zip Code)

Registrant's telephone number, including area code (650) 366-2626

N/A (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))

INFORMATION TO BE INCLUDED IN THE REPORT

Item 2.02 Results of Operations and Financial Condition Item 9.01 Financial Statements and Exhibits Signature Exhibit Index EX-99.1

ITEM 2.02 Results of Operations and Financial Condition

On November 15, 2010, the Company issued a press release announcing its financial results for the quarter ended September 30, 2010. A copy of this press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K (the "Report"). The press release should be read in conjunction with the note regarding forward-looking statements, which is included in the text of the press release.

The information in this Item 2.02 and attached as Exhibit 99.1 to this Report will not be treated as "filed" for the purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section. This information will not be incorporated by reference into any filing under the Securities Act of 1933, as amended, or into another filing under the Exchange Act, unless that filing expressly incorporates this information by reference.

ITEM 9.01 Financial Statements and Exhibits

(d) Exhibits

99.1 Press Release of A. P. Pharma, Inc., dated November 15, 2010

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 hereunto duly authorized.

Date: November 15, 2010

A.P. Pharma, Inc.

/s/ John B. Whelan

John B. Whelan Acting Chief Executive Officer and Chief Financial Officer

Exhibit	Description
99.1	Press Release of A. P. Pharma, Inc., dated November 15, 2010



A.P. Pharma Announces Third Quarter 2010 Financial Results

REDWOOD CITY, Calif. – November 15, 2010 – A.P. Pharma, Inc. (NASDAQ: APPA), a specialty pharmaceutical company, today reported financial results for its third quarter ended September 30, 2010.

Operational Highlights

"A. P. Pharma has been working diligently to address the issues raised in the U.S. Food and Drug Administration's Complete Response Letter of March 2010," said John Whelan, A.P. Pharma's acting chief executive officer. "Our goal is to schedule what we hope will be a productive End of Review meeting with the agency as soon as the necessary information is available. The primary focus of a meeting with the agency would be to discuss the next steps for the APF530 New Drug Application."

In November, A.P. Pharma received notice that it was awarded a non-taxable grant from the United States government under the Qualifying Therapeutics Discovery Project (QTDP) program in the amount of \$244,479. Grants were awarded to projects that show reasonable potential to produce new therapies, address unmet medical needs, and reduce the long-term growth of health care costs in the U.S. The QTDP program is a part of the Patient Protection and Affordable Health Care Act of 2010.

Results of Operations

A.P. Pharma's net loss for the third quarter of 2010 was \$1.7 million, or \$0.04 per share, compared with a net loss of \$1.2 million, or \$0.04 per share, for the third quarter of 2009. Net loss was higher in the current fiscal quarter primarily due to \$1.0 million of income recognized in the prior year quarter in connection with the termination of a license agreement for APF530 with RHEI Pharmaceuticals, N.V. Revenue for the third quarter of 2010 was \$0.4 million compared with revenue of \$1.1 million for the third quarter of 2009. Revenue in 2010 was primarily related to research and development work performed under an agreement with Merial Limited entered into in September 2009 for a long-acting pain management product for companion animals. The \$0.7 million decrease in revenue in the current fiscal quarter was partially offset by \$0.3 million of lower spending resulting from continuing cost containment actions undertaken by the Company.

Cash and cash equivalents as of September 30, 2010 were \$3.0 million, compared with \$7.6 million at December 31, 2009.

In March 2010, A.P. Pharma received a Complete Response Letter from the U.S. Food and Drug Administration (FDA) on the APF530 New Drug Application (NDA). The full extent of activities, costs and time required to address the FDA's questions is not currently known; however, A.P. Pharma expects to clarify the actions required for resubmission and approval of its NDA at an End of Review meeting. Based on a current analysis of anticipated expenses to prepare for an End of Review meeting, the Company believes it has sufficient cash resources to fund operations into the first quarter of 2011. A.P. Pharma is currently seeking debt or equity financing to fund its operations. Multiple factors, including market conditions, may prevent the Company from obtaining adequate financing to support its operations, or obtaining financing that will be on terms favorable to A.P. Pharma or its stockholders.

About APF530

A.P. Pharma's lead product candidate, APF530, is being developed for the prevention of both acute and delayed onset chemotherapy-induced nausea and vomiting (CINV). APF530 contains the 5-HT3 antagonist, granisetron, formulated in the Company's proprietary Biochronomer[™] drug delivery system, which allows therapeutic drug levels to be maintained for five days with a single subcutaneous injection. Intravenous and oral formulations containing granisetron are approved for the prevention of acute-onset CINV, but not for delayed-onset CINV. Granisetron was selected because it is widely prescribed by physicians based on a well-established record of safety and efficacy.

About A.P. Pharma

A.P. Pharma is a specialty pharmaceutical company developing products using its proprietary Biochronomer[™] polymer-based drug delivery technology. The Company's primary focus is on its lead product, APF530, for the prevention of CINV. A.P. Pharma received a Complete Response Letter on the APF530 NDA in March 2010 and is in the process of preparing a resubmission responsive to the deficiencies listed in the Complete Response Letter. The Company has additional clinical and preclinical stage programs in the area of pain management, all of which utilize its bioerodible injectable and implantable delivery systems. Further work on these programs has been deferred while the Company focuses on the approval of APF530. For further information, visit the Company's web site at www.appharma.com.

(financial tables follow)

A.P. Pharma, Inc. Condensed Statements of Operations (in thousands, except per share amounts) (Unaudited)									
		Three Months Ended September 30,		Nine Months Ended September 30,					
		2010	2009	2010	2009				
Contract revenue	<u>\$</u>	351	\$ 1,117	\$ 1,122	\$ 1,139				
Operating expenses:									
Research and development		1,541	1,418	5,762	6,376				
General and administrative		445	912	3,561	2,905				
Total operating expenses		1,986	2,330	9,323	9,281				
Operating loss		(1,635)	(1,213)	(8,201)	(8,142)				
Gain on sale of royalty interest		-	-	2,500	-				
Interest income (expense), net		(1)	(1)		27				
Loss from continuing operations		(1,636)	(1,214)		(8,115)				
Loss from discontinued operations		(36)	-	(47)	-				
Net loss	\$	(1,672)	\$ (1,214)	\$ (5,750)	\$ (8,115)				
Basic and diluted net loss per share:									
Loss from continuing operations	\$	(0.04)	\$ (0.04)	\$ (0.14)	\$ (0.26)				
Net loss	\$	(0.04)	\$ (0.04)	\$ (0.15)	\$ (0.26)				
Shares used to compute basic and diluted net									
loss per share		39,507	31,234	39,481	31,041				

A.P. Pharma	Inc.								
Condensed Balance Sheets									
(in thousan	ds)								
		P	1 04						
	September 30, 2010	De	December 31, 2009						
	2010		2009						
Assets									
Current assets:									
Cash and cash equivalents	\$ 3,0	15 \$	7,593						
Accounts receivable		28	171						
Prepaid expenses and other current assets	2	86	549						
Total current assets	3,6	29	8,313						
Property and equipment, net	4	07	510						
Other long-term assets		53	128						
Total assets	\$ 4,0	89 \$	8,951						
Liabilities and Stockholders' Equity									
Current liabilities:									
Accounts payable		57 \$	162						
Accrued expenses		17	1,080						
Deferred revenue		95	92						
Accrued disposition costs		00	553						
Total current liabilities	1,3		1,887						
Deferred revenue		89	268						
Total liabilities	1,5	58	2,155						
Stockholders' equity:									
Common stock		01	394						
Additional paid-in capital	148,9		147,481						
Accumulated deficit	(146,8		(141,079)						
Total stockholders' equity	2,5		6,796						
Total liabilities and stockholders' equity	\$ 4,0	<u>89</u> <u></u>	8,951						

Forward-looking Statements

This news release contains "forward-looking statements" as defined by the Private Securities Litigation Reform Act of 1995. These forward-looking statements involve risks and uncertainties, including uncertainties associated with capital resources and liquidity, timely development and regulatory approval of product candidates, satisfactory completion of clinical studies, progress in research and development programs, launch and acceptance of new products and other risks and uncertainties identified in the Company's filings with the Securities and Exchange Commission. We caution investors that forward-looking statements reflect our analysis only on their stated date. We do not intend to update them except as required by law.

Contacts

Corporate Contact: A.P. Pharma, Inc. John B. Whelan, Acting Chief Executive Officer and Chief Financial Officer 650-366-2626

and

Investor and Media Relations:

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