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

April 9, 2014
Date of Report (Date of earliest event reported)

ACURA PHARMACEUTICALS, INC.

(Exact Name of Registrant as Specified in Charter)

State of New York	**1-10113**	**11-0853640**
(State of Other Jurisdiction of Incorporation)	(Commission File Number)	(I.R.S. Employer Identification Number)

616 N. North Court, Suite 120
Palatine, Illinois 60067
(Address of principal executive offices) (Zip Code)

(847) 705-7709
(Registrant's telephone number, including area code)

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 (17CFR 240.14a-12)

☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17CFR240.14d- 2(b))

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

Item 1.01 Entry into a Material Definitive Agreement.

The contents of Item 1.02 are incorporated herein by reference.

Item 1.02 Termination of a Material Definitive Agreement.

On April 9, 2014 we entered into a letter agreement ("Letter Agreement") with King Pharmaceuticals Research & Development, Inc. ("King"), now a subsidiary of Pfizer Inc., (collectively "Pfizer") terminating the License, Development and Commercialization Agreement dated October 30, 2007 (the "License Agreement") between King and us. In September, 2012, we had entered into another letter agreement with Pfizer pursuant to which Pfizer returned to us all products licensed to it under the License Agreement except for OXECTA® (oxycodone hydrochloride) Tablets CII. Pfizer received U.S. Food and Drug Administration approval for its New Drug Application ("NDA") for OXECTA in June 2011. With this Letter Agreement we regain all rights to the products licensed to Pfizer under the License Agreement.

Pursuant to the terms of and the Letter Agreement:

- Our license grant to Pfizer for our AVERSION® Technology will terminate effective April 9, 2014 and Pfizer will no longer have a right to develop, market or sell any products using our technology;
- We will own all rights to OXECTA® (oxycodone hydrochloride) Tablets CII, except for the OXECTA® trademark, which will be retained by Pfizer;
- Pfizer will transfer to us all studies, data, regulatory filings (including the NDA) and other information relating to OXECTA® (oxycodone hydrochloride) Tablets CII pursuant to a transition process described in the Letter Agreement;
- We will pay Pfizer a one-time payment of $2,000,000 in consideration of the termination of the License Agreement;
- Neither party will have any royalty obligations to the other, except for Pfizer's royalty payment obligations to us relating to OXECTA® product sold before the effective date of the Letter Agreement; and
- Each of Pfizer and Acura waives any claims and releases the other party against any liability relating to the License Agreement, except for any claims and obligations under the Letter Agreement and certain sections of the License Agreement that survive termination in accordance with the terms of the Letter Agreement.

Item 2.03 Creation of a Direct Financial Obligation or an Obligation under an Off-Balance Sheet Arrangement of a Registrant.

Under the terms of the Letter Agreement we must pay Pfizer $2,000,000 in two installments: $1,000,000 by April 19, 2014 and $1,000,000 within five (5) days of Pfizer notifying the FDA of a transfer of the new drug application for OXECTA®. Item 1.02 of this Report is incorporated herein by reference.

Item 8.01 Other Events

On April 10, 2014 we issued a press release relating to the Letter Agreement, which press release is attached hereto as Exhibit 99.1.

Item 9.01 Financial Statements and Exhibits

Exhibit Number	**Description**
99.1	Press Release dated April 10, 2014 Announcing Termination of License Agreement between Pfizer and Acura.

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 thereunto duly authorized.

ACURA PHARMACEUTICALS, INC.

By: /s/ Peter A. Clemens
Peter A. Clemens
Senior Vice President & Chief Financial

Date: April 9, 2014

Exhibit Number	Description
99.1	Press Release dated April 10, 2014 Announcing Termination of License Agreement between Pfizer and Acura.

Acura Pharmaceuticals Announces Return of Product Rights

Palatine, IL - (April 10, 2014) - Acura Pharmaceuticals, Inc. (NASDAQ: ACUR), a specialty pharmaceutical company developing products intended to address medication abuse and misuse, announced today a letter agreement with Pfizer Inc. providing for the termination of Pfizer's license to Acura's AVERSION® Technology and the return to Acura of the FDA approved Oxecta® (oxycodone HCl) product. The letter agreement provides that Acura will make a one-time payment of $2.0 million to Pfizer. The license termination is effective April 9, 2014. The AVERSION® Technology utilizes a proprietary mixture of inactive ingredients to discourage tampering of a product for abusive purposes.

"We are pleased that we have been able to reach agreement on acceptable terms for the license termination," said Bob Jones, President and Chief Executive Officer of Acura Pharmaceuticals. Mr. Jones further added, "We are currently evaluating our strategic options for the returned product and our other AVERSION® Technology products in development, which may include a re-launch under a new brand name in partnership with another pharmaceutical company."

The Company will host a conference call to discuss our strategies for the product going forward on Friday, April 11, 2014 at 8:30 am ET. To participate in the live conference call, please dial 1-888-556-4997 (U.S. and Canada) five to ten minutes prior to the start of the call. The participant passcode is 9666652

About Acura Pharmaceuticals
Acura Pharmaceuticals is a specialty pharmaceutical company engaged in the research, development and commercialization of product candidates intended to address medication abuse and misuse, utilizing its proprietary AVERSION® and IMPEDE® Technologies. AVERSION® contains polymers that cause the drug to gel when dissolved; it also contains compounds that irritate the nasal passages. IMPEDE® is designed to disrupt the processing of pseudoephedrine from tablets into methamphetamine.

In June 2011, the U.S. Food and Drug Administration approved OXECTA® (oxycodone HC1 tablets) which incorporates the AVERSION® technology. The Company has a development pipeline of additional AVERSION® technology products containing other opioids.

Forward-Looking Statements

Certain statements in this press release constitute "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995. Such forward-looking statements involve known and unknown risks, uncertainties and other factors which may cause our actual results, performance or achievements to be materially different from any future results, performance, or achievements expressed or implied by such forward-looking statements. Forward-looking statements may include, but are not limited to, our ability to enter into a license agreement for our Aversion oxycodone product, our and our licensee's ability to successfully launch and commercialize our products and technologies including our Aversion oxycodone and NEXAFED® Tablets, the price discounting that may be offered by our licensee for Aversion oxycodone, our or our licensee's ability to obtain commercial supplies of Aversion oxycodone from a third party contract manufacturer, our and our licensee's ability to obtain necessary regulatory approvals and commercialize products utilizing our technologies and the market acceptance of and competitive environment for any of our products, the willingness of wholesalers and pharmacies to stock NEXAFED® Tablets, expectations regarding the terms and payments under any license agreement for our products, expectations regarding potential market share for our products and the timing of first sales, our ability to enter into additional license agreements for our AVERSION® Technology product candidates, our exposure to product liability and other lawsuits in connection with the commercialization of our products, the increasing cost of insurance and the availability of product liability insurance coverage, the ability to avoid infringement of patents, trademarks and other proprietary rights of third parties, and the ability of our patents to protect our products from generic competition, our ability to protect and enforce our patent rights in any paragraph IV patent infringement litigation, and the ability to fulfill the FDA requirements for approving our product candidates for commercial manufacturing and distribution in the United States, including, without limitation, the adequacy of the results of the laboratory and clinical studies completed to date, the results of laboratory and clinical studies we may complete in the future to support FDA approval of our product candidates and the sufficiency of our development to meet OTC Monograph standards as applicable, the adequacy of the development program for our product candidates, including whether additional clinical studies will be required to support FDA approval of our product candidates, changes in regulatory requirements, adverse safety findings relating to our product candidates, whether the FDA will agree with our analysis of our clinical and laboratory studies and how it may evaluate the results of these studies or whether further studies of our product candidates will be required to support FDA approval, whether or when we are able to obtain FDA approval of labeling for our product candidates for the proposed indications and will be able to promote the features of our abuse discouraging technologies, whether our product candidates will ultimately deter abuse in commercial settings and whether our Impede technology will disrupt the processing of pseudoephedrine into methamphetamine. In some cases, you can identify forward-looking statements by terms such as "may," "will," "should," "could," "would," "expects," "plans," "anticipates," "believes," "indicates," "estimates," "projects," "predicts," "potential" and similar expressions intended to identify forward-looking statements. These statements reflect our current views with respect to future events and are based on assumptions and subject to risks and uncertainties. Given these uncertainties, you should not place undue reliance on these forward-looking statements. We discuss many of these risks in greater detail in our filings with the Securities and Exchange Commission.

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