

ARENA PHARMACEUTICALS INC  
Form 8-K  
August 25, 2016

**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): August 24, 2016**

**Arena Pharmaceuticals, Inc.**

**(Exact name of registrant as specified in its charter)**

**Delaware**  
**(State or other jurisdiction**

**of incorporation)**

**000-31161**  
**(Commission**

**File Number)**

**6154 Nancy Ridge Drive, San Diego, California 92121**

**23-2908305**  
**(I.R.S. Employer**

**Identification No.)**

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**(Address of principal executive offices) (Zip Code)**

**858.453.7200**

**(Registrant's telephone number, including area code)**

**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:

- .. 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))

In this report, Arena Pharmaceuticals, Arena, Company, we, us and our refer to Arena Pharmaceuticals, Inc., one or more of our wholly owned subsidiaries, unless the context otherwise provides. Arena Pharmaceuticals® and Arena® are registered service marks of Arena Pharmaceuticals, Inc.

### **Item 8.01 Other Events.**

We and Eisai Inc. have received a Paragraph IV certification notification with respect to patents for BELVIQ® (lorcaserin hydrochloride tablets, 10 mg) listed in the FDA's Approved Drug Products with Therapeutic Equivalence Evaluations, or the Orange Book. The certification was from an Abbreviated New Drug Application (ANDA) for a proposed generic version of BELVIQ®.

The notification is currently under investigation. In accordance with the Hatch-Waxman Act, we and Eisai have 45 days from the receipt of the notification to file a patent infringement suit against the ANDA filer which would result in the stay of FDA approval of the ANDA under statutory guidelines.

We and Eisai intend to vigorously defend our intellectual property rights. We cannot predict the outcome of any litigation matter.

### **Forward-Looking Statements**

Certain statements in this Form 8-K are forward-looking statements that involve a number of risks and uncertainties. Such forward-looking statements include our and Eisai's plans to defend our intellectual property rights and any lawsuit. For such statements, we claim the protection of the Private Securities Litigation Reform Act of 1995. Actual events or results may differ materially from our expectations. Factors that could cause actual results to differ materially from the forward-looking statements include, but are not limited to, risks related to litigation, intellectual property rights and collaborations. Additional factors that could cause actual results to differ materially from those stated or implied by Arena's forward-looking statements are disclosed in our filings with the Securities and Exchange Commission. These forward-looking statements represent our judgment as of the time of this Form 8-K. We disclaim any intent or obligation to update these forward-looking statements, other than as may be required under applicable law.

**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: August 24, 2016

Arena Pharmaceuticals, Inc.

By: /s/ Amit Munshi  
Amit Munshi  
President and Chief Executive Officer