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**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION**

Washington, D.C. 20549

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**FORM 8-K**

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**CURRENT REPORT  
Pursuant to Section 13 or 15(d)  
of the Securities Exchange Act of 1934**

**Date of Report (Date of earliest event reported): April 30, 2015**

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**FibroGen, Inc.**

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

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

**001-36740**  
(Commission  
File Number)

**77-0357827**  
(IRS Employer  
Identification No.)

**FibroGen, Inc.**  
**409 Illinois Street**  
**San Francisco, CA 94158**  
(Address of principal executive offices, including zip code)

**(415) 978-1200**  
(Registrant's telephone number, including area code)

**Not Applicable**  
(Former name or former address, if changed since last report.)

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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))
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**Item 7.01 Regulation FD Disclosure**

On April 30, 2015, FibroGen, Inc. (the Company) conducted meetings with prospective investors in which its corporate slide presentation was presented. The presentation included a discussion of whether, if approved, the Company's Phase 3 clinical trial product candidate, roxadustat, will be required to be included in the bundled payment system for dialysis center drugs and services by the Centers for Medicare and Medicaid Services (CMS).

Under the Medicare Improvements for Patients and Providers Act (MIPPA) a basic case-mix adjusted composite, or bundled, payment system commenced in January 2011 and transitioned fully by January 2014 to a single reimbursement rate for drugs and all services furnished by renal dialysis centers for Medicare beneficiaries with end-stage renal disease. Specifically, under MIPPA the bundle now covers drugs, services, lab tests and supplies under a single treatment base rate for reimbursement by the CMS based on the average cost per treatment, including the cost of ESAs and IV iron doses, typically without adjustment for usage. It is unknown whether roxadustat, if approved, will be included in the payment bundle. Under MIPPA, agents that have no IV equivalent in the bundle are currently expected to be excluded from the bundle until 2024. Based on roxadustat's differentiated mechanism of action and therapeutic effects, and discussions with its collaboration partner, the Company currently believes that roxadustat would not be included in the bundle, and would instead be reimbursed in this setting under Medicare Part D.

The information in this Item 7.01 is being furnished, not filed, pursuant to Regulation FD. Accordingly, the information in Item 7.01 of this report will not be incorporated by reference into any registration statement filed by the Company under the Securities Act of 1933, as amended, unless specifically identified therein as being incorporated therein by reference. The furnishing of the information in this report is not intended to, and does not, constitute a determination or admission by the Company that the information in this report is material or complete, or that investors should consider this information before making an investment decision with respect to any security of the Company.

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**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 hereunto duly authorized.

**FIBROGEN, INC.**

Dated: April 30, 2015

By: /s/ Michael Lowenstein

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