

Sanofi
Form 6-K
March 19, 2019

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER
PURSUANT TO RULE 13a-16 OR 15d-16
UNDER THE SECURITIES EXCHANGE ACT OF 1934

For the month of March 2019

Commission File Number: 001-31368

SANOFI

(Translation of registrant's name into English)

54, rue La Boétie, 75008 Paris, FRANCE

(Address of principal executive offices)

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Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F.

Form 20-F

Form 40-F

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):

Indicate by check mark whether the registrant by furnishing the information contained in this Form is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes

No

If Yes marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b):

82-_____

In March 2019, Sanofi issued the press releases attached hereto as Exhibit 99.1, 99.2, 99.3 and 99.4 which are incorporated herein by reference.

Exhibit List

| Exhibit | |
|--------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| No. | Description |
| Exhibit 99.1 | Press release dated March 15, 2019: Praluent® (alirocumab) now approved in European Union to reduce the risk of cardiovascular events in patients with established cardiovascular disease |
| Exhibit 99.2 | Press release dated March 13, 2019: Sanofi successfully prices EUR 2 billion of bond issues |
| Exhibit 99.3 | Press release dated March 8, 2019: Filing of the 2018 U.S. Form 20-F and French « Document de Référence » containing the Annual Financial Report |
| Exhibit 99.4 | Press release dated March 8, 2019: FDA to undertake priority review of Dupixent® (dupilumab) for adults with inadequately controlled severe chronic rhinosinusitis with nasal polyps |

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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, thereunto duly authorized.

Dated: March 19, 2019

SANOFI

By /s/ Alexandra Roger
Name: Alexandra Roger
Title: Head of Securities Law and Capital Markets