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Sanofi Form 6-K December 27, 2016

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 December 2016

Commission File Number: 001-31368

SANOFI

(Translation of registrant s name into English)

54, rue La Boétie, 75008 Paris, FRANCE

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(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or I	Form 40	n 4
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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 November and December 2016, Sanofi issued the press releases attached hereto as Exhibit 99.1 and 99.2 which is incorporated herein by reference.

Exhibit List

No.	Description	
Exhibit 99.1	Press release dated November 21, 2016: Sanofi Receives FDA Approval of Soliqua Treatment of Adults with Type 2 Diabetes	100/33, for the
Exhibit 99.2	Press release dated December 8, 2016: Sanofi and Regeneron Announce Marketing Application for Dupixent® (dupilumab) Accepted for Review by the EMA	Authorization

Exhibit

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: December 27, 2016 SANOFI

By /s/ Alexandra Roger Name: Alexandra Roger

Title: Head of Securities Law and Capital Markets

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Exhibit Index

Exhibit

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Exhibit 99.1 Press release dated November 21, 2016: Sanofi Receives FDA Approval of Soliqua 100/33, for the Treatment of Adults with Type 2 Diabetes

Exhibit 99.2 Press release dated December 8, 2016: Sanofi and Regeneron Announce Marketing Authorization Application for Dupixent® (dupilumab) Accepted for Review by the EMA