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

 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): November 13, 2017

Arbutus Biopharma Corporation

(Exact Name of Registrant as Specified in Charter)

BRITISH COLUMBIA, CANADA (State or Other Jurisdiction of Incorporation)

[]

[]

001-34949 (Commission File Number)

980597776

(I.R.S. Employer Identification Number)

100-8900 Glenlyon Parkway, Burnaby, British Columbia, Canada V5J 5J8

(Address of Principal Executive Offices) (Zip Code)

(604) 419-3200

(Registrant's telephone number, including area code)

(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))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (17 CFR §230.405) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR §240.12b-2). Emerging growth company []
If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 8.01. Other Events.

On November 13, 2017, the Registrant issued a press release, a copy of which is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits.

Exhibit 99.1. Press release dated November 13, 2017

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.

Arbutus Biopharma Corporation

Date: November 13, 2017 By: <u>/s/ Bruce G. Cousins</u>

Bruce G. Cousins

Executive Vice President and Chief Financial Officer

Arbutus' LNP Licensee Alnylam Receives Accelerated Assessment of Patisiran from European Medicines Agency (EMA)

VANCOUVER, British Columbia and WARMINSTER, Pa., Nov. 13, 2017 (GLOBE NEWSWIRE) -- Arbutus Biopharma Corporation (Nasdaq:ABUS), an industry-leading Hepatitis B Virus (HBV) therapeutic solutions company, announced today that the Company's lipid nanoparticle (LNP) licensee Alnylam Pharmaceuticals, Inc. (Nasdaq:ALNY), announced the grant of an accelerated assessment from the European Medicines Agency (EMA) for patisiran, an investigational RNAi therapeutic being developed for patients with hereditary ATTR (hATTR) amyloidosis. Accelerated assessment may provide a reduced review timeline from 210 to 150 days once the marketing authorization application (MAA) is filed and validated, which Alnylam intends to file by year-end 2017.

Dr. Mark J Murray, Arbutus' President and CEO said, "We are very pleased to see our licensing partner, Alnylam, advance it's LNP-enabled RNAi asset, patisiran, one step closer to final regulatory approval. Arbutus could see its first royalty payment from patisiran as early as next year, pending final regulatory approvals."

About Arbutus

Arbutus Biopharma Corporation is a biopharmaceutical company dedicated to discovering, developing, and commercializing a cure for patients suffering from chronic HBV infection. Arbutus is headquartered in Vancouver, BC, and has facilities in Warminster, PA. For more information, visit www.arbutusbio.com.

Forward-Looking Statements and Information

This press release contains forward-looking statements within the meaning of the Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934, and forward-looking information within the meaning of Canadian securities laws (collectively, "forward-looking statements"). Forward-looking statements in this press release include statements about the filing of an MAA by Alnylam; a reduced timeline for the assessment of patisiran; Arbutus receiving its first royalty payment from patisiran as early as next year; and discovering, developing and commercializing a cure for chronic hepatitis B virus (HBV) infection.

With respect to the forward-looking statements contained in this press release, Arbutus has made numerous assumptions regarding, among other things: the speed of regulatory approvals; continued and timely positive preclinical and clinical efficacy data; the continued demand for Arbutus' assets, including its LNP technology; and the stability of economic and market conditions. While Arbutus considers these assumptions to be reasonable, these assumptions are inherently subject to significant business, economic, competitive, market and social uncertainties and contingencies.

Additionally, there are known and unknown risk factors which could cause Arbutus' actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements contained herein. Known risk factors include, among others: patisiran may not receive regulatory approval on a timely basis, or at all; Arbutus may not receive timely royalty payments, or at all; and market shifts may require a change in strategic focus.

A more complete discussion of the risks and uncertainties facing Arbutus appears in Arbutus' Annual Report on Form 10-K and Arbutus' continuous disclosure filings, which are available at www.sedar.com and at www.sec.gov. All forward-looking statements herein are qualified in their entirety by this cautionary statement, and Arbutus disclaims any obligation to revise or update any such forward-looking statements or to publicly announce the result of any revisions to any of the forward-looking statements contained herein to reflect future results, events or developments, except as required by law.

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