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): January 9, 2017

HALOZYME THERAPEUTICS, INC.

(Exact name of registrant as specified in its charter)

Delaware	**001-32335**	**88-0488686**
(State or other jurisdiction of incorporation)	**(Commission File Number)**	**(IRS Employer Identification No.)**

11388 Sorrento Valley Road, San Diego, California	**92121**
(Address of principal executive offices)	**(Zip Code)**

Registrant's telephone number, including area code: (858) 794-8889

Not Applicable
(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))

Item 2.02. Results of Operations and Financial Condition.

On January 9, 2017, Halozyme Therapeutics, Inc., a Delaware corporation ("Halozyme") issued a press release (the "Press Release") which contained information related to Halozyme's expected 2016 revenue for partner reimbursed R&D expense. A copy of the press release is attached hereto as Exhibit 99.1.

On January 9, 2017, Halozyme presented at the annual JP Morgan Healthcare Conference to provide a corporate update on certain strategic programs and to provide financial guidance for 2017. The presentation contained information related to Halozyme's expected 2016 royalty revenue and 2016 revenue for partner reimbursed R&D expense. A copy of the slides from the presentation containing this information is attached hereto as Exhibit 99.2.

Exhibits 99.1 and 99.2 are furnished under Item 2.02 of this report and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, (the "Exchange Act") or otherwise subject to the liabilities of that section, nor shall they be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, regardless of any general incorporation language in such filing.

Item 7.01 Regulation FD Disclosure.

The Press Release and certain slides from the JP Morgan Healthcare Conference presentation attached hereto as Exhibits 99.1 and 99.2, respectively, also provided a corporate update on certain strategic programs and provided financial guidance for 2017.

This information is being furnished pursuant to Item 7.01 of this Report and shall not be deemed to be "filed" for the purposes of Section 18 of the Securities and Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that section and will not be incorporated by reference into any registration statement filed by Halozyme, under the Securities Act of 1933, as amended, unless specifically identified as being incorporated therein by reference. This Report will not be deemed an admission as to the materiality of any information in this Report that is being disclosed pursuant to Regulation FD.

Please refer to the press release attached hereto as Exhibit 99.1 for a discussion of certain forward-looking statements included therein and the risks and uncertainties related thereto.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
99.1	Press release dated January 9, 2017
99.2	Certain slides from Halozyme corporate update presentation dated January 9, 2017

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.

Halozyme Therapeutics, Inc.

January 9, 2017

By: */s/ Harry J. Leonhardt Esq.*

Harry J. Leonhardt, Esq.

Senior Vice President, General Counsel,
Chief Compliance Officer and Corporate Secretary

Exhibit Index

Exhibit No.	Description
99.1	Press release dated January 9, 2017
99.2	Certain slides from Halozyme corporate update presentation dated January 9, 2017

EXHIBIT 99.1

Contacts:
Jim Mazzola
858-704-8122
ir@halozyme.com

Chris Burton
858-704-8352
ir@halozyme.com

HALOZYME PROVIDES PROGRAM UPDATES, 2017 FINANCIAL GUIDANCE AT 35 TH ANNUAL JP MORGAN HEALTHCARE CONFERENCE

SAN DIEGO, Jan. 9, 2017 — Halozyme Therapeutics, Inc. (NASDAQ: HALO), a biotechnology company developing novel oncology and drug-delivery therapies, today provided program updates and its annual financial guidance at the 35 th annual JP Morgan Healthcare Conference.

“We enter the year with strongly supportive data from our HALO 202 study in metastatic pancreatic cancer patients and momentum in our HALO 301 global registration trial for our investigational new drug PEGPH20 in a similar, targeted HA-High patient population,” said Dr. Helen Torley, president and chief executive officer. “In 2017 we expect to make continued progress demonstrating the pan-tumor potential of PEGPH20, as we also focus on near-term catalysts in our revenue-generating ENHANZE™ platform, including the potential approval of rituximab SC in the U.S. and progressing other partnered programs in the clinic.”

In addition to HALO 301, the company has ongoing studies of PEGPH20 in combination with Merck’s KEYTRUDA ® (pembrolizumab) in gastric and non-small-cell lung cancer patients and in collaboration with Eisai in combination with HALAVEN ® (eribulin) in breast cancer patients. Two new studies exploring four tumor types are planned to start in 2017 as part of a recently announced clinical collaboration with Genentech. Under the collaboration, Genentech will evaluate PEGPH20 in combination with its anti-PDL1 TECENTRIQ ® (atezolizumab) in pancreas and gastric cancer and Halozyme will evaluate the same combination in gallbladder cancer and cholangiocarcinoma. Halozyme has also been included in an innovative, patient-centered clinical trial planned for initiation in 2017 called Precision Promise, led by the Pancreatic Cancer Action Network.

In the Halozyme ENHANZE™ platform business, the company announced in November 2016 that the U.S. Food and Drug Administration (FDA) has accepted Genentech’s Biologics License Application for a subcutaneous formulation of rituximab in multiple blood cancer indications with an action date in June. This is a co-formulation with Halozyme's proprietary recombinant human

hyaluronidase enzyme (ENHANZE platform), approved and marketed under the MabThera ® SC brand in countries outside the U.S.

In addition, Halozyme plans in 2017 to support ongoing development of subcutaneous formulations for Roche's PERJETA ® and Janssen's DARZALEX ® , work with existing ENHANZE platform partners to advance development of additional licensed targets, and seek to sign new global licensing and collaboration agreements.

The company also provided financial guidance for 2017 of:

- **Revenue of $115 million to $130 million** , excluding revenue from any new ENHANZE global collaboration and licensing agreements that may be signed during the year. The company expects to report $20 million in 2016 revenue for reimbursed partner R&D expenses that will not recur in 2017;

- **Operating Expenses of $240 million to $250 million** , supporting the ongoing Phase 3 study in metastatic pancreatic cancer patients and the continued execution of clinical programs to study the pan-tumor potential of PEGPH20;

- **Year-end cash balance of $100 million to $110 million** .

Dr. Torley will present at 3 p.m. PST, Jan. 9 at the conference. Her presentation will be webcast through the "Investors" section of www.halozyme.com, and a recording will be made available for 90 days following the event. To access the live webcast, please log on approximately fifteen minutes prior to the presentation to register and download any necessary audio software.

About Halozyme
Halozyme Therapeutics is a biotechnology company focused on developing and commercializing novel oncology therapies that target the tumor microenvironment. Halozyme's lead proprietary program, investigational drug PEGPH20, applies a unique approach to targeting solid tumors, allowing increased access of co-administered cancer drug therapies to the tumor in animal models. PEGPH20 is currently in development for metastatic pancreatic cancer, non-small cell lung cancer, gastric cancer, metastatic breast cancer and has potential across additional cancers in combination with different types of cancer therapies. In addition to its proprietary product portfolio, Halozyme has established value-driving partnerships with leading pharmaceutical companies including Roche, Baxalta, Pfizer, Janssen, AbbVie and Lilly for its ENHANZE™ drug delivery platform. Halozyme is headquartered in San Diego. For more information visit www.halozyme.com.

###

Safe Harbor Statement
In addition to historical information, the statements set forth above include forward-looking statements concerning the Company's future expectations and plans for 2017, timing and results of clinical trials, the development and commercialization of product candidates and the potential

benefits and attributes of such product candidates (including, without limitation, statements concerning the possible activity, benefits and attributes of PEGPH20, the possible method of action of PEGPH20, its potential application to improve cancer therapies and statements concerning future actions relating to the development of PEGPH20). These statements also include forward-looking statements concerning the possible activity, benefits and attributes of ENHANZE™, the possible method of action of ENHANZE, its potential application to aid in the dispersion and absorption of other injected therapeutic drugs, the number of collaborative targets actually chosen, the product development efforts of our ENHANZE partners, whether such products are ultimately developed or commercialized, whether milestones triggering milestone payments will be achieved, and statements concerning facilitating more rapid delivery of injectable medications through subcutaneous delivery. The forward looking statements also include the Company's expected financial outlook for 2017 and statements concerning expectations for 2016 revenue for reimbursed partner R&D expenses. These forward looking statements involve risk and uncertainties that could cause actual results to differ materially from those in the forward-looking statements. The forward-looking statements are typically, but not always, identified through use of the words "believe," "enable," "may," "will," "could," "intends," "estimate," "anticipate," "plan," "predict," "probable," "potential," "possible," "should," "continue," and other words of similar meaning. Actual results could differ materially from the expectations contained in forward-looking statements as a result of several factors, including unexpected expenditures and costs, unexpected fluctuations or changes in revenues from collaborators or product sales, audited 2016 financial results differing from the financial results stated above, unexpected clinical trial delays or results, including enrollment delays, unexpected results or delays in development and regulatory review, regulatory approval requirements, unexpected adverse events and competitive conditions. These and other factors that may result in differences are discussed in greater detail in the Company's most recent Annual and Quarterly Reports filed with the Securities and Exchange Commission.

EXHIBIT 99.2

Significant Milestones Achieved in 2016

ENHANZE Platform

- ✓ Royalties grew 65% versus 2015
- ✓ Rituximab SC BLA filed in the U.S.
- ✓ Darzalex SC progressing toward Phase 3 study
- ✓ Roche initiates study with Perjeta

Oncology Pipeline

- ✓ Dosed first patient, initiated ~85% of sites in HALO-301 Study
- ✓ Initiated dose expansion in Keytruda trial
- ✓ Initiated HALO-Eisai Study in Breast Cancer Patients
- ✓ Signed Roche I-O/PEGPH20 clinical collaboration
- ✓ Expanded pipeline with novel preclinical assets

Halozyme

2017 Financial Guidance

	2016	2017	Notes
Net Revenue	$145M to $150M	$115M to $130M	• 2017 guidance excludes new ENHANZE partnerships • Robust royalty growth projected to continue in 2017 • 2016 included ~$20M of one-time reimbursed R&D expenses
Operating Expenses	$240M to $245M	$240M to $250M	• Continued investment in Phase 3 pancreas and pan-tumor studies
Year-end Cash	$180M to $190M	$100M to $110M	• Royalty-backed loan repayment starts 2017