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): **June 17, 2013**

AMICUS THERAPEUTICS, INC.
(Exact name of registrant as specified in its charter)

Delaware	**001-33497**	**71-0869350**
(State or other Jurisdiction of Incorporation)	(Commission File Number)	(IRS Employer Identification No.)
1 Cedar Brook Drive, Cranbury, NJ		**08512**
(Address of Principal Executive Offices)		(Zip Code)

Registrant's telephone number, including area code: **(609) 662-2000**

(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:

o Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

o Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

o Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

o Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Item 7.01. Regulation FD Disclosure.

On June 17, 2013, Amicus Therapeutics, Inc. (the "Company") hosted a conference call and webcast to discuss regulatory strategy for migalastat HCl monotherapy for Fabry disease following a recent Type C meeting with the U.S. Food and Drug Administration (FDA). A copy of the conference call presentation materials is attached hereto as Exhibit 99.1.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits: The Exhibit Index annexed hereto is incorporated herein by reference.

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.

AMICUS THERAPEUTICS, INC.

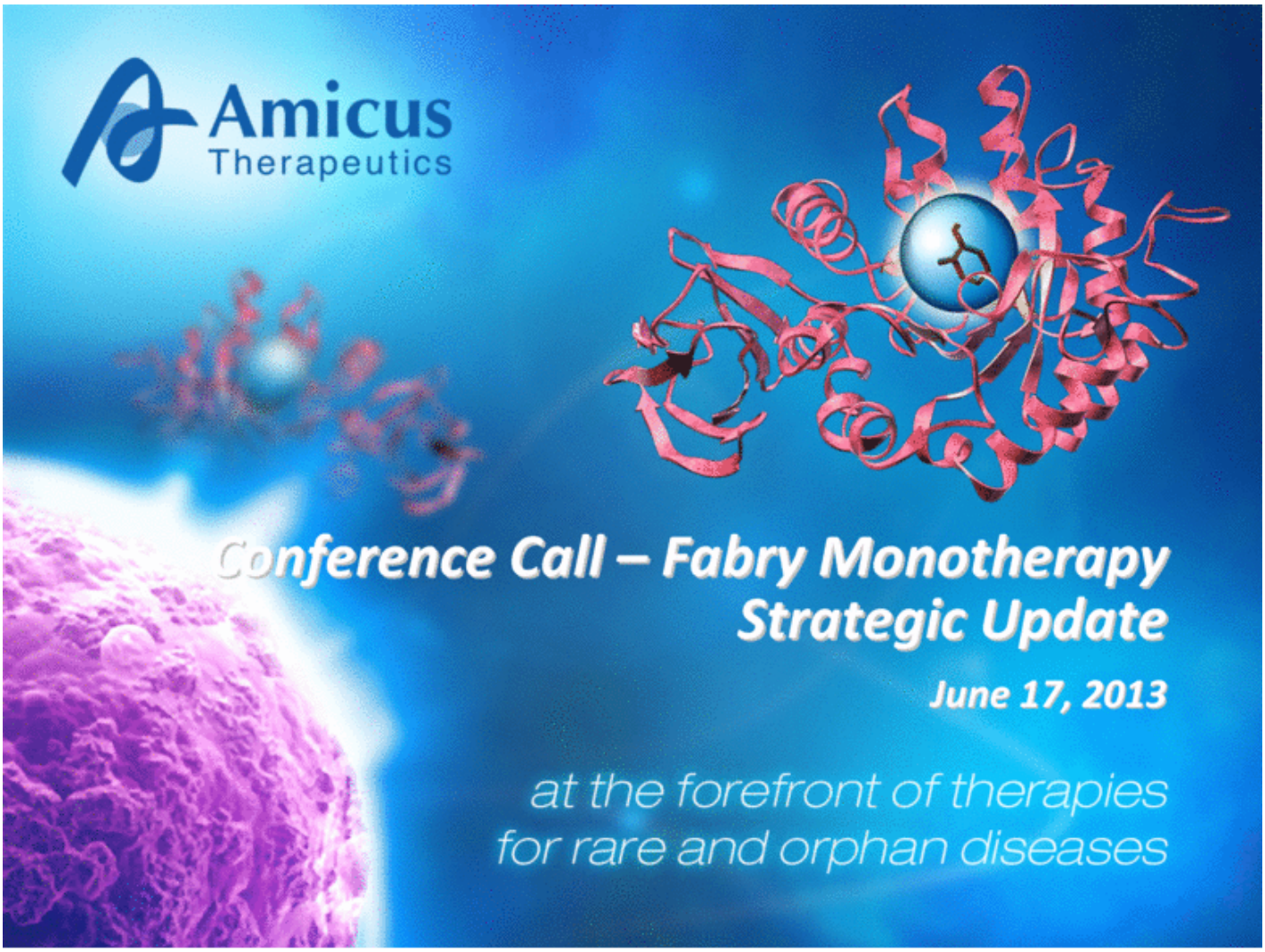
Date: June 17, 2013

By: /s/ PETER M. MACALUSO
Name: Peter M. Macaluso
Title: Secretary

EXHIBIT INDEX

Exhibit No.	Description
99.1	Presentation Materials

Exhibit 99.1


Amicus
Therapeutics
Conference Call – Fabry Monotherapy
Strategic Update
June 17, 2013
at the forefront of therapies
for rare and orphan diseases

Safe Harbor

This presentation contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995 relating to business, operations and financial conditions of Amicus including but not limited to preclinical and clinical development of Amicus' candidate drug products, the timing and reporting of results from clinical trials evaluating Amicus' candidate drug products. Words such as, but not limited to, "look forward to," "believe," "expect," "anticipate," "estimate," "intend," "plan," "would," "should" and "could," and similar expressions or words, identify forward-looking statements. Although Amicus believes the expectations reflected in such forward-looking statements are based upon reasonable assumptions, there can be no assurance that its expectations will be realized. Actual results could differ materially from those projected in Amicus' forward-looking statements due to numerous known and unknown risks and uncertainties, including the "Risk Factors" described in our Annual Report on Form 10-K for the year ended December 31, 2012. All forward-looking statements are qualified in their entirety by this cautionary statement, and Amicus undertakes no obligation to revise or update this presentation to reflect events or circumstances after the date hereof.

Amicus Therapeutics

Migalastat HCl Monotherapy for Fabry Disease

"We remain very committed to advancing migalastat HCl monotherapy, which we are developing with GlaxoSmithKline, toward a potential U.S. approval for patients with Fabry disease who have amenable mutations."

-John F. Crowley, Chairman and CEO

Amicus Therapeutics

FDA Feedback: Type C Meeting*

- Study 011 missed 6-month primary endpoint therefore Stage 1 (6-month) data not sufficient for conditional approval
- Acknowledged trends towards improvement in kidney interstitial capillary GL-3 in Stage 1, although not statistically significant
- Recognized merit of proposed additions to Study 011 Stage 2 (12-month) statistical analysis plan, however, will not consider them pre-specified
- Agreed to follow-up meeting to review full data from Phase 2 and Phase 3 studies

Migalastat HCl Monotherapy: Development Strategy

Assembling Robust Dataset to Maximize Chances for Potential U.S. Approval of Migalastat HCl Monotherapy for Fabry Patients with Amenable Mutations

- We remain fully committed to Fabry collaboration
- Study 011 Stage 2 (12-month) top-line data expected 4Q13
- Study 012 top-line data anticipated 2H14
- FDA meeting anticipated 2H14 to discuss U.S. approval pathway

Migalastat HCl Monotherapy: Proposed NDA Submission

Data from >100 Patients to Support Potential U.S. Approval

- Study 011 (primary endpoint – kidney interstitial capillary GL-3)
  - Stage 1 and Stage 2 kidney biopsies and clinical assessments (6- and 12-month data)
  - Clinical assessments from 12-month treatment extension (no biopsies)
- Study 012 (primary endpoint – measured GFR, no biopsies)
  - Analysis from 18-month primary treatment period
  - Clinical assessments from 12-month treatment extension
- Ongoing extension studies
  - Long-term data for patients rolling over from Phase 2 and Phase 3 treatment extensions

Amicus Therapeutics

Anticipated Milestones

Building Shareholder Value

2H13	Top-Line 12-Month Data from Phase 3 Fabry Monotherapy Study 011
	Initiation of Phase 2 Repeat-Dose Pompe Co-Administration Study
	Disclosure of CHART™ Program for Additional LSD target
1H14	Initiation of Phase 1/2 Fabry Co-Formulation Study
	Initial Data from Phase 2 Repeat-Dose Pompe Co-Administration Study
2H14	Top-Line Data from Phase 3 Monotherapy Study 012
	FDA Meeting to Discuss Fabry Monotherapy Approval Pathway
	Phase 1 Data from Phase 1/2 Fabry Co-Formulation Study

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