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 5, 2017

ANTRIABIO, INC.
(Name of registrant in its charter)

Delaware	**000-54495**	**27-3440894**
(State or jurisdiction of incorporation or organization)	(Commission File Number)	(IRS Employer Identification No.)

1450 Infinite Drive
Louisville, CO 80027
(Address of principal executive offices)

(303) 222-2128
(Registrant's telephone number)

(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 7.01. Regulation FD Disclosure.

On January 5, 2017, we issued a shareholder letter attached hereto as Exhibit 99.1. In accordance with General Instruction B.2 of Form 8-K, the information set forth herein is deemed to be "furnished" and shall not be deemed to be "filed" for purposes of the Securities Exchange Act of 1934, as amended. The information set forth in Item 7.01 of this Current Report on Form 8-K shall not be deemed an admission as to the materiality of any information in this Current Report on Form 8-K that is required to be disclosed solely to satisfy the requirements of Regulation FD.

Item 9.01 Financial Statements and Exhibits

EXHIBIT	**DESCRIPTION**
99.1	Shareholder Letter of AntriaBio, Inc. dated January 5, 2017 **

** The following exhibit relating to Item 7.01 is intended to be furnished to, not filed with, the SEC pursuant to Regulation FD.

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.

ANTRIABIO, INC.

DATE: January 5, 2017

By: /s/ Morgan Fields
Morgan Fields
Chief Accounting Officer

EXHIBIT INDEX

EXHIBIT	DESCRIPTION
99.1	Shareholder Letter of AntriaBio, Inc. dated January 5, 2017 **

** The following exhibit relating to Item 7.01 is intended to be furnished to, not filed with, the SEC pursuant to Regulation FD.

EXHIBIT 99.1

January 5, 2017

Dear Shareholders:

We are writing to highlight a few recent developments in the competitive landscape of the once-weekly basal insulin market. AntriaBio closely tracks the progress of companies with pipeline products that we view as potential competition to AB101 market share. The three programs management believes are the most relevant are the weekly insulin programs from Hanmi Pharmaceutical Co., Ltd. (^{LAPS}Insulin 115), PhaseBio Pharmaceuticals (PE0139) and Novo Nordisk (LAI287). We believe AB101 is in a position to be a differentiated product as it is the only product that uses native, unmodified insulin in its drug substance. This may confer a safety and regulatory advantage over all three of our competitors whose weekly basal insulin candidates are insulin analogs or modified forms of insulin.

Late last week, Hanmi announced it was amending the licensing agreement it signed with Sanofi in November 2015. Under the original agreement, Sanofi licensed the development and commercialization rights for three of Hanmi's long-acting diabetes products: efpeglenatide (weekly/monthly GLP-1 agonist), ^{LAPS}Insulin 115 (weekly basal insulin) and ^{LAPS}Insulin Combo (GLP-1 agonist + basal insulin combination). The agreement stipulated that Hanmi would receive an upfront payment of 400 million euros and up to 3.5 billion euros in development, registration and sales milestones, in addition to double-digit royalties on net sales.

Under the terms of the revised licensing agreement, Sanofi returned development rights for ^{LAPS}Insulin 115 to Hanmi. As a result, Hanmi will return 196 million euros to Sanofi by December 2018 and Sanofi's commercial milestone payments to Hanmi have been reduced by 798 million euros. Hanmi will also now be responsible for part of the development expenses for efpeglenatide and receive reduced milestone payments for the drug.

In 2014, Hanmi terminated their lead clinical stage weekly insulin compound, HM12460A, for failing to meet the target product profile of a once-weekly subcutaneous injection. Though Hanmi initiated Phase 1 clinical studies of their follow-on compound, ^{LAPS}Insulin 115 in February 2015, neither Hanmi nor Sanofi has announced or presented data from these trials.

Based on the preclinical PK/PD data Hanmi presented at the American Diabetes Association 74 th Scientific Sessions® in San Francisco in 2014, we believe ^{LAPS}Insulin 115 fails to meet the target product profile of a once-weekly subcutaneous injection. We believe the data showed an acute release of insulin with significant glucose lowering within the first 24 hours. Peak insulin levels and glucose lowering occurred at Day 2-3 and resolved by Day 5-6, which leads us to believe that the ^{LAPS}Insulin 115 formulation may fall short of once-weekly dosing. We believe any issues related to the efficacy of ^{LAPS}Insulin 115, if they exist, are unique to ^{LAPS}Insulin 115 and are not applicable to all once-weekly insulin therapies.

In contrast, single dose and repeat weekly dose preclinical studies of AB101 in multiple animal species demonstrated no acute insulin release or glucose reduction and also demonstrated slow onset, sustained insulin increases and corresponding glucose reductions over the entire course of the weekly dosing interval.

AntriaBio continues to believe in the promise of the once-weekly insulin class to deliver a basal insulin with fewer side effects and greater accessibility to patients needing insulin, given the right formulation and product. Due to its potential safety advantages and consistent weekly profile in animals, we believe AB101 is uniquely positioned to meet this promise. Recent events strengthen our competitive advantage. In addition to our view of AB101 as a best in class therapy, we also believe AB101 remains in a position to be first in class to take share of the current ~$11 billion basal insulin market. We look forward to commencing our first clinical study this year and we are optimistic we will demonstrate human proof of concept for AB101 as a once-weekly insulin therapy.

Best Regards,

Noopur Liffick
Vice President, Corporate Development

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Forward-Looking Statements

This shareholder letter, like many written and oral communications presented by AntriaBio, Inc., and our authorized officers, may contain certain forward-looking statements regarding our prospective performance and strategies within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended. We intend such forward-looking statements to be covered by the safe harbor provisions for forward-looking

statements contained in the Private Securities Litigation Reform Act of 1995, and are including this statement for purposes of said safe harbor provisions. Forward-looking statements, which are based on certain assumptions and describe future plans, strategies, and expectations of the Company, are generally identified by use of words "anticipate," "believe," "estimate," "expect," "intend," "plan," "project," "seek," "strive," "try," or future or conditional verbs such as "could," "may," "should," "will," "would," or similar expressions. Our ability to predict results or the actual effects of our plans or strategies is inherently uncertain. Accordingly, actual results may differ materially from anticipated results. Readers are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date of this release. Except as required by applicable law or regulation, AntriaBio undertakes no obligation to update these forward-looking statements to reflect events or circumstances that occur after the date on which such statements were made.