

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): **March 7, 2016**

**ARCH THERAPEUTICS, INC.**

(Exact name of registrant as specified in its charter)

**Nevada**  
(State or other jurisdiction  
of incorporation)

**000-54986**  
(Commission  
File Number)

**46-0524102**  
(I.R.S. Employer  
Identification No.)

**235 Walnut Street, Suite 6**  
**Framingham, Massachusetts**  
(Address of principal executive offices)

**01702**  
(Zip Code)

Registrant's telephone number, including area code: **(617) 431-2313**

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 (see General Instruction A.2. below):

- 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))
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**Item 8.01 Other Events.**

On March 7, 2016, Arch Therapeutics, Inc. (the “**Company**”) issued a press release announcing that the Company has commenced enrollment and treatment in its first clinical trial in Western Europe. The text of the press release is attached hereto as Exhibit 99.1 and is incorporated by reference herein.

**Item 9.01 Financial Statements and Exhibit**

(d) Exhibits

<b>Exhibit</b>	<b>Description</b>
99.1	Press Release issued by Arch Therapeutics, Inc. on March 7, 2016

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**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.

**ARCH THERAPEUTICS, INC.**

Dated: March 7, 2016

By: /s/ Terrence W. Norchi, M.D.  
Name: Terrence W. Norchi, M.D.  
Title: President, Chief Executive Officer

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## Exhibit List

<b>Exhibit</b>	<b>Description</b>
99.1	Press Release issued by Arch Therapeutics, Inc. on March 7, 2016

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## Arch Therapeutics Clinical Trial Actively Enrolling Patients and Providing Investigational Treatment

### *Major Milestone Along Path Toward Commercialization of AC5*

**FRAMINGHAM, MA – March 7, 2016** — Arch Therapeutics, Inc. (OTCQB: ARTH) (“Arch” or the “Company”), developer of the AC5 Surgical Hemostatic Device™ (“AC5™”), has initiated patient enrollment and treatment in its first clinical trial in Western Europe.

Terrence W. Norchi, MD, President and CEO of Arch, said, “The start of this trial marks a major milestone for the Company. It is firmly underway with patients being actively screened, enrolled, and treated or scheduled for treatment. Our team has put in tremendous effort to get us to this point and we are pleased with this progress.”

The randomized controlled single-blind investigation is designed to assess safety and performance of AC5 during the course of a dermatological procedure performed in up to approximately 45 patients, of whom 10 will be taking an antiplatelet medication (commonly referred to as a “blood thinner”) during the study period. Patients participating in the study will have at least two dermatological lesions removed surgically. One of the wounds created as a result of the excision will receive AC5 while the other will receive a control treatment.

The endpoints include product related adverse effects and time to hemostasis. Follow-up assessments of patients are planned for seven and 30 days following the procedure. As previously stated, data are projected to be available within two quarters of the start of the trial.

Dr. Norchi added, “This clinical trial is a key component of our commercialization plan, representing the first time that AC5 has been used on humans in a surgical setting. Further, it highlights that Arch has completed a number of important achievements that are required before a patient could be treated with AC5. As we are getting closer to bringing AC5 to market, we continue to advance our pre-clinical work and clinical-regulatory planning for additional indications for AC5 and other products, in each case with the ultimate goal of making a difference in patients’ lives.”

#### **About Arch Therapeutics, Inc.**

Arch Therapeutics, Inc. is a biotechnology company developing a novel approach to stop bleeding (hemostasis) and control leaking (sealant) during surgery and trauma care. Arch is developing products based on an innovative self-assembling peptide technology platform to make surgery and interventional care faster and safer for patients. Arch's flagship development stage product candidate, known as the AC5 Surgical Hemostatic Device™, is being designed to achieve hemostasis in minimally invasive and open surgical procedures and is intended to be regulated as a medical device.

Find out more at [www.archtherapeutics.com](http://www.archtherapeutics.com).

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**Notice Regarding Forward-Looking Statements**

This news release contains "forward-looking statements" as that term is defined in Section 27(a) of the Securities Act of 1933, as amended, and Section 21(e) of the Securities Exchange Act of 1934, as amended. Statements in this press release that are not purely historical are forward-looking statements and include any statements regarding beliefs, plans, expectations or intentions regarding the future. Such forward-looking statements include, among other things, references to novel technologies and methods, our business and product development plans and projections, or market information. Actual results could differ from those projected in any forward-looking statements due to numerous factors. Such factors include, among others, the inherent uncertainties associated with developing new products or technologies and operating as a development stage company, our ability to retain important members of our management team and attract other qualified personnel, our ability to raise the additional funding we will need to continue to pursue our business and product development plans, our ability to develop and commercialize products based on our technology platform, and market conditions. These forward-looking statements are made as of the date of this news release, and we assume no obligation to update the forward-looking statements, or to update the reasons why actual results could differ from those projected in the forward-looking statements. Although we believe that any beliefs, plans, expectations and intentions contained in this press release are reasonable, there can be no assurance that any such beliefs, plans, expectations or intentions will prove to be accurate. Investors should consult all of the information set forth herein and should also refer to the risk factors disclosure outlined in the reports and other documents we file with the SEC, available at [www.sec.gov](http://www.sec.gov).

On Behalf of the Board,  
Terrence W. Norchi, MD  
Arch Therapeutics, Inc.

**Contact:**

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