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): September 27, 2018

ADMA BIOLOGICS, INC.

(Exact name of registrant as specified in its charter)

Delaware	001-36728	56-2590442
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)
465 State Route 17, Ramsey, New Jersey		07446
(Address of principal executive offices)		(Zip Code)

Registrant's telephone number, including area code: (201) 478-5552

(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 (*see* General Instruction A.2. below):

o 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 as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ý

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 5.02 Departure of Directors or Certain Officers; Election of Directors; Appointment of Certain Officers; Compensatory Arrangements of Certain Officers.

On September 27, 2018, the Board of Directors (the "Board") of ADMA Biologics, Inc. (the "Company") approved payments in 2019 of the following temporary living expenses of certain of the Company's named executive officers: (i) a continuation payment of $5,000 per month in temporary housing expenses, plus appropriate tax gross-up, through December 2019, for Brian Lenz, the Company's Executive Vice President and Chief Financial Officer; and (ii) a continuation payment of $8,000 per month in temporary living expenses, plus appropriate tax gross-up, through December 2019, for Dr. James Mond, the Company's Executive Vice President, Chief Scientific Officer and Chief Medical Officer. As previously disclosed by the Company in its Current Report on Form 8-K filed with the Securities and Exchange Commission on February 15, 2018, the Board previously approved the temporary housing expense payments of $5,000 and $8,000 to Mr. Lenz and Dr. Mond, respectively, plus appropriate tax gross-ups, through December 2018.

Item 8.01 Other Events.

On October 3, 2018, the Company issued a press release announcing that its wholly-owned subsidiary, ADMA Bio Centers Georgia Inc., has received U.S. Food and Drug Administration ("FDA") approval for its third plasma collection center, located at 166 Ernest W Barrett Parkway, NW, Marietta, Georgia. The facility commenced operations and initiated source plasma collection in December 2017, and is now FDA licensed to collect and enter into interstate commerce to sell and use the human source plasma for further manufacturing in the United States.

The full text of the press release is filed as Exhibit 99.1 to this Current Report on Form 8-K.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
99.1	ADMA Biologics, Inc. Press Release, dated October 3, 2018.

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.

October 3, 2018

ADMA Biologics, Inc.

By: /s/ Brian Lenz

Name: Brian Lenz

Title: Executive Vice President and Chief Financial Officer

ADMA BioCenters Receives FDA Approval for Third Plasma Collection Center

RAMSEY, NJ & MARIETTA, GA – October 3, 2018 – ADMA Biologics, Inc. (NASDAQ: ADMA) ("ADMA" or the "Company"), a vertically integrated commercial biopharmaceutical company that develops, manufactures and markets specialty plasma-based biologics for the treatment of Primary Immune Deficiency Disease ("PIDD") and the prevention and treatment of certain infectious diseases, announces that the Company's wholly-owned subsidiary, ADMA BioCenters, has received U.S. Food and Drug Administration ("FDA") approval for its third plasma collection center, located at 166 Ernest W Barrett Parkway, NW, Marietta, Georgia. The facility commenced operations and initiated source plasma collection in December 2017, and is now FDA licensed to collect and enter into interstate commerce to sell and use the human source plasma for further manufacturing in the U.S.

Additionally, ADMA BioCenters has received FDA approval for a license supplement to collect whole blood from donors with a rare blood type to produce source plasma that contains high levels of Anti-D antibodies which is the basis for producing Rho_o (D) Immune Globulin (Anti-D Ig). Other plasma products manufacturers use this Anti-D high-titer plasma to make a hyperimmune globulin which is used to prevent and treat conditions associated with Rh negative blood type. Market prices for Anti-D plasma can be as much as two times that of normal source plasma.

"FDA approval of our third plasma collection facility is an important milestone that enables us to continue to internally supply a portion of our raw material, as well as collect other hyperimmune source plasma for the manufacturing of ADMA's products in our Boca Raton, FL facility or for sale to other customers," stated Adam Grossman, President and CEO of ADMA Biologics. "As a vertically-integrated plasma products manufacturer, operating raw-material source plasma collection facilities allows us to continue to leverage our know-how and potentially expand our ADMA BioCenters business in the future to meet our growing demands. I am pleased that we obtained approval of this facility and our license supplement well ahead of our expected timeline and we are grateful to our dedicated and knowledgeable staff for meeting this important milestone."

Plasma can be used for the manufacture of a variety of live-saving and life-sustaining therapies. Plasma centers provide local economies with financial resources through employment opportunities and compensation to local residents for their generous plasma donations. At over 12,000 square feet, the new ADMA BioCenters facility is anticipated to support approximately 50 donor beds at peak capacity and is expected to employ up to 50 staff members. Plasma donors have an opportunity to potentially earn up to $400 per month or more by donating at an ADMA Bio Center. For information about plasma donation or to become a donor with ADMA BioCenters, please visit www.admabiocenters.com.

About ADMA BioCenters
ADMA BioCenters is a wholly-owned subsidiary of ADMA Biologics, which operates as a source plasma collection business. ADMA BioCenters currently holds FDA, German Health Authorities (GHA), and Korean Ministry of Food and Drug Safety (MFDS) licenses to operate as a source plasma collection organization for both U.S. based and foreign fractionators' therapeutic plasma products manufacturing. A typical plasma collection center can collect between 30,000 to 50,000 liters of source plasma annually. Plasma collected from ADMA BioCenters' FDA approved facilities that is not used to manufacture ADMA's products or development-stage candidates is sold to customers under an existing supply agreement or in the open "spot" market generating revenues for the Company.

About ADMA Biologics, Inc. (ADMA)
ADMA Biologics is a vertically integrated commercial biopharmaceutical company that manufactures, markets and develops specialty plasma-based biologics for the treatment of Primary Immune Deficiency Disease (“PIDD”) and the prevention and treatment of certain infectious diseases. ADMA's mission is to develop and commercialize plasma-derived, human immune globulins targeted to niche patient populations for the treatment and prevention of certain infectious diseases. The target patient populations include immune-compromised individuals who suffer from an underlying immune deficiency disease, or who may be immune-compromised for other medical reasons. ADMA has received U.S. Patents 9,107,906, 9,714,283, 9,815,886 and 9,969,793 related to certain aspects of its lead product candidate, RI-002. For more information, please visit www.admabiologics.com.

Cautionary Note Regarding Forward-Looking Statements
This press release contains "forward-looking statements" pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995, about ADMA Biologics, Inc. ("we“, “our” or the “Company”). Forward-looking statements include, without limitation, any statement that may predict, forecast, indicate, or imply future results, performance or achievements, and may contain the words "estimate," “project,” "intend," “forecast,” "target," ”anticipate,” “plan,” “planning,” “expect,” “believe,” “will," “is likely,” “will likely,” “should,” “could,” "would," "may," or, in each case, their negative, or words or expressions of similar meaning. These forward-looking statements also include, but are not limited to, statements concerning our plans to develop, manufacture, market, launch and expand our own commercial infrastructure and commercialize our current products and future products, the safety, efficacy and expected timing of, and our ability to, obtain and maintain regulatory approvals of our current products and product candidates, and the labeling or nature of any such approvals, the success of our work with our third party vendors and the U.S. Food and Drug Administration (the “FDA”) in furtherance of and progress towards an approval of our Biologics License Application for specialty plasma-based biologics and the ability of such third parties to respond adequately or in a timely manner to the issues raised by the FDA, our ability to successfully pursue commercialization and prelaunch activities, the timeframe within which we may receive approval from the FDA for specialty plasma-based biologics, if at all, the potential of our specialty plasma-based biologics to provide meaningful clinical improvement for patients living with Primary Immune Deficiency Disease or other indications and our ability to realize increased prices for plasma growth in the plasma collection industry. Actual events or results may differ materially from those described in this document due to a number of important factors. Current and prospective security holders are cautioned that there also can be no assurance that the forward-looking statements included in this press release will prove to be accurate. In light of the significant uncertainties inherent in the forward-looking statements included herein, the inclusion of such information should not be regarded as a representation or warranty by ADMA or any other person that the objectives and plans of ADMA will be achieved in any specified time frame, if at all. Except to the extent required by applicable laws or rules, ADMA does not undertake any obligation to update any forward-looking statements or to announce revisions to any of the forward-looking statements. Forward-looking statements are subject to many risks, uncertainties and other factors that could cause our actual results, and the timing of certain events, to differ materially from any future results expressed or implied by the forward-looking statements, including, but not limited to, the risks and uncertainties described in our filings with the U.S. Securities and Exchange Commission, including our most recent reports on Form 10-K, 10-Q and 8-K, and any amendments thereto.

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