UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, DC 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 17, 2020

MIMEDX GROUP, INC.
(Exact name of registrant as specified in charter)

Florida	**001-35887**	**26-2792552**
(State or other jurisdiction of incorporation)	**(Commission File Number)**	**(IRS Employer Identification No.)**

1775 West Oak Commons Ct., NE, Marietta GA 30062
(Address of principal executive offices) (Zip Code)

Registrant's telephone number, including area code: (770) 651-9100

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))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
None	**n/a**	**n/a**

Indicate by check mark whether the registrant is an emerging growth company 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 ☐

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 7.01 Regulation FD Disclosure

On September 17, 2020, MiMedx Group, Inc. (the "***Company***" or the "***Registrant***") issued a press release announcing the conclusion of enrollment for a Phase 3 study of plantar fasciitis. A copy of the press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

All information in the press release speaks as of the date thereof, and MiMedx does not assume any obligation to update such information in the future. In addition, MiMedx disclaims any inference regarding the materiality of such information which otherwise may arise as a result of its furnishing such information under Item 7.01 of this Current Report on Form 8-K.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No.	Description of Exhibit
99.1	Press release dated September 17, 2020.
104	The cover page from this Current Report on Form 8-K, formatted in Inline XBRL.

SIGNATURES

Pursuant to the requirements of the Exchange Act, the Registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

MIMEDX GROUP, INC.

Date: September 17, 2020

By: /s/ Peter M. Carlson
Peter M. Carlson
Chief Financial Officer

Exhibit 99.1

MiMedx Concludes Enrollment for Phase 3 Plantar Fasciitis Trial

Research in Plantar Fasciitis Advances Understanding of Amniotic Tissue as Potential Treatment for Chronic Musculoskeletal Pain and Function Disorders

MARIETTA, Ga., September 17, 2020 — MiMedx Group, Inc. (OTC PINK: MDXG) ("MiMedx" or "the Company"), an industry leader in advanced wound care and an emerging therapeutic biologics company, today announced the conclusion of enrollment for a Phase 3 study of plantar fasciitis, an inflammation of the fibrous tissue along the bottom of the foot that results in intense pain. This key clinical trial explores how placental science may address and treat musculoskeletal pain and function disorders. The study has met its timelines.

"Plantar fasciitis is an all-too-common, debilitating and painful foot condition that challenges both patients and clinicians. Several months to years of treatment may be required with conservative therapies before symptoms subside, and I believe plantar fasciitis represents a significant unmet patient need," said Stuart Miller, M.D., Principal Investigator, Department of Orthopaedic Surgery, MedStar Union Memorial Hospital, and Assistant Professor, Department of Orthopaedic Surgery, Johns Hopkins University School of Medicine, Baltimore, Maryland. "This is a landmark study that will help us advance the science and elevate the standard of care for millions of patients; I look forward to analyzing and publishing the data regarding treatment efficacy for this musculoskeletal condition. Our progress to date is all the more gratifying given the dampening effect of the pandemic on patient enrollment. This study is designed to provide statistically significant evidence of efficacy for this biologic treatment to reduce pain and improve function."

More than two million people are treated for plantar fasciitis inflammation in the United States annually. In 10% of patients treated with traditional measures, the condition progresses to chronic plantar fasciitis-related pain – recovery from which is lengthy and recurrence of which is very common, with an estimated $284 million annual national economic burden. The current treatment algorithm aims to maintain arch shape, modify foot loading and/or improve shock absorbency of the heel through night splints and orthotics. While they may assist in reducing pain associated with plantar fasciitis, these treatments do not address the root cause of the condition, which is thought to be both degenerative and inflammatory.

Timothy R. Wright, MiMedx Chief Executive Officer, commented, "Given the variability of efficacy, cost, and potential side effects of available plantar fasciitis treatment options, additional evidence-based alternatives are needed urgently. As a pioneer in the development of placental tissue technology, following through on clinical research is part of our mission to improve patient outcomes. Using our placental science platform to address the unmet need posed by plantar fasciitis is just one of the critical ways that we are exploring its application to improve people's lives. We look forward to sharing the results of this trial in 2021."

About the MiMedx Plantar Fasciitis Trial

This study is a Phase 3, prospective, double-blinded, randomized controlled trial of the micronized dehydrated Human Amnion Chorion Membrane (dHACM) injection as compared to saline placebo injection in the treatment of plantar fasciitis. The trial enrolled 276 patients between the ages of 21 and 79 years, with an investigator-confirmed diagnosis of plantar fasciitis for ³ 1 month (30 days) and £ 18 months. Patients were required to have a Visual Analog Scale (VAS) Pain scale of ³ 45 mm at randomization and be receiving conservative treatment for ³ 1 month (30 days), including any of the following modalities: Rest, Ice, Compression, Elevation (RICE); stretching exercises; NSAIDs or orthotics. The primary endpoints are change in VAS for Pain at 90 Days and incidence of related adverse events at 180 days, serious adverse events and unanticipated events during the first 12 months post-injection. Secondary endpoints include self-reported responses to the Foot Function Index – Revised (FFI-R) at 90 days.

Important Cautionary Statement

This press release includes forward-looking statements. Statements regarding expectations of study results, the timing of study results, and the expected timing of a Biologics License Application (BLA) filing are forward-looking. Additional forward-looking statements may be identified by words such as “believe,” “expect,” “may,” “plan,” “potential,” “will,” “preliminary,” and similar expressions, and are based on management’s current beliefs and expectations. Forward-looking statements are subject to risks and uncertainties, and the Company cautions investors against placing undue reliance on such statements. Actual results may differ materially from those set forth in the forward-looking statements. The Company describes additional risks and uncertainties in the Risk Factors section of its most recent annual report and quarterly reports filed with the Securities and Exchange Commission. Any forward-looking statements speak only as of the date of this press release and the Company assumes no obligation to update any forward-looking statement.

About MiMedx

MiMedx® is an industry leader in advanced wound care and an emerging therapeutic biologics company developing and distributing human placental tissue allografts with patent-protected processes for multiple sectors of healthcare. The Company processes the human placental tissue utilizing its proprietary PURION® process methodology, among other processes, to produce allografts by employing aseptic processing techniques in addition to terminal sterilization. MiMedx has supplied more than 1.9 million allografts to date. For additional information, please visit www.mimedx.com.

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