

**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): August 10, 2020**

**MIMEDX GROUP, INC.**

(Exact name of registrant as specified in charter)

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

**001-35887**  
(Commission  
File Number)

**26-2792552**  
(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
<b>None</b>	<b>n/a</b>	<b>n/a</b>

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.

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**Item 7.01 Regulation FD Disclosure**

On August 10, 2020, MiMedx Group, Inc. (the “**Company**” or the “**Registrant**”) issued a press release announcing the hiring of Robert Stein as Executive Vice President – Research and Development. A copy of the press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits.

<b>Exhibit No.</b>	<b>Description of Exhibit</b>
99.1	<a href="#">Press release dated August 10, 2020.</a>
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: August 10, 2020

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

**MiMedx Welcomes Robert B. Stein, M.D., Ph.D., to Head Revitalized R&D Efforts***Pharmaceutical and Biotech Executive to Focus on Product and Pipeline Development Priorities*

MARIETTA, Ga., August 10, 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 addition of Robert B. Stein, M.D., Ph.D., as Executive Vice President, Research and Development, effective August 10, 2020. Noted for significant contributions and accomplishments in medical research and development, Dr. Stein brings to MiMedx more than 40 years of drug discovery and development experience across multiple pharmaceutical and biotechnology companies. He has served as a lead contributor in the discovery and registration of eight marketed drugs, and is experienced in product development, including in the areas of molecular and cellular biology, biochemistry, animal pharmacology, drug metabolism, and safety assessment, among others.

Timothy R. Wright, MiMedx Chief Executive Officer, commented, “The science of regenerative tissue technology holds tremendous promise for the millions of people who suffer with chronic, difficult-to-heal wounds – and our commitment is to advance category science and demonstrate value so that patients can access the care they need. Dr. Stein’s arrival at MiMedx validates our commitment to be a category leader. A highly regarded and respected industry scientist, he brings the extensive development, regulatory and registration expertise we need to advance the rigor of placenta-based science, and accelerate the transition of our category in line with recent FDA guidance. MiMedx is investing in Research and Development as core to our future portfolio and pipeline, and Dr. Stein is known for creating and advancing best-in-class treatment options, with a patient-centric focus on achieving the best outcomes.”

Dr. Stein added, “Throughout my career I’ve been drawn to research efforts and companies that are taking on meaningful and challenging health problems. MiMedx is taking the next steps to further the science and better understand the clinical potential of placental tissue to bring improved healing to more people in need. My role is to continually enhance the rigor of our strong scientific foundation that will more clearly inform our pipeline and expand the Company’s potential to address significant areas of unmet clinical need.”

**About Robert B. Stein, M.D., Ph.D.**

After completing his Bachelors of Science with honors, earning a double major in both biology and chemistry at Indiana University, Dr. Stein received his M.D. and Ph.D. in Physiology and Pharmacology and completed his internship, residency and Board Certification in Anatomic and Clinical Pathology, all at Duke University. Following residency, Dr. Stein worked at Merck, with contributions to Cozaar, Sustiva, and Gardasil. He was then recruited as the first head of Research and Development for Ligand Pharmaceuticals, with responsibility for building the Research and Development organization and programs targeting various nuclear hormone receptors. This work led to eight pharmaceutical partnerships and six marketed medicines, including two SERMs (Fablyn and Viviant), three novel retinoids (Panretin, Targetin gel and capsules), and Promacta, the small molecule Thrombopoietin mimetic.

After six years at Ligand, he became Executive Vice President, Research and Pre-clinical Development for DuPont-Merck and DuPont pharmaceuticals, leading to the registration of Sustiva and Innohep and the discovery and advancement of blockbuster Eliquis, subsequently registered by Bristol Myers Squibb Company. Following the acquisition of DuPont by Bristol Myers Squibb, Dr. Stein joined Incyte Pharmaceuticals as President, R&D and Chief Scientific Officer, spearheading the transition from genomics to drug discovery and development. Following Incyte, Dr. Stein became President of Roche Palo Alto LLC, where he built Roche's Translational Medicine capabilities, served on the Global Early Development Committee, Global Biologics Steering Committee, and Joint Roche-Genentech Steering Committee. Dr. Stein then joined Kinemed, a translational medicine company as Chief Executive Officer.

Following Kinemed, Dr. Stein served as President, R&D, for Agenus, an immuno-oncology company, with responsibility for the development of their R&D organization, and the advancement of four checkpoint modulatory antibodies and a personalized neo-epitope-directed cancer vaccine to Phase 1 trials. He also led the generation of a best-in-class TCR discovery and optimization effort. He served as a full time Senior Advisor, R&D, to Agenus and AgenTus from March 2017 to October 2019. During this time, Agenus advanced 13 monoclonal antibodies into the clinic and formed significant partnerships with Incyte, Merck, UCB, and Gilead. The two most advanced Agenus products, AGEN2034 (PD-1 antagonist monoclonal antibody) and AGEN1884 (CTLA-4 antagonist monoclonal antibody) are on track for potential BLA submission in 2020.

Dr. Stein is deeply experienced in the lab, including in the areas of molecular and cellular biology, biochemistry, enzymology, animal pharmacology, virology, drug metabolism, and safety assessment. He also has extensive experience in Translational Medicine and Early Clinical Development. He has led groups of over 1,000 scientists and physicians for over 15 years, supervising work in all the major therapeutic areas. In addition, Dr. Stein serves on the board of directors for a number of private and public company boards, and acts as an advisor to a multiple clients and academic institutions.

### **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](http://www.mimedx.com).

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