Momenta Pharmaceuticals to Host R&D Day on October 11, 2018

October 4, 2018

CAMBRIDGE, Mass., Oct. 04, 2018 (GLOBE NEWSWIRE) -- Momenta Pharmaceuticals, Inc. (Nasdaq: MNTA) today announced that it will host a live webcast of its R&D Day on Thursday, October 11, 2018 beginning at 8:30 am ET. The event is being held in New York, NY.

Momenta's leadership team plans to highlight the clinical development strategy for its pipeline of novel drug candidates for immune-mediated disorders. Presentations will include:

- Data from the Phase 1 study in healthy volunteers for M281, a potentially best-in-class anti-FcRn antibody engineered to reduce circulating pathogenic IgG antibodies by blocking endogenous IgG recycling via FcRn, and the indications for the planned Phase 2 proof of concept studies. Momenta will be joined by a key opinion leader to discuss one of the indications;
- Overview of the Phase 1/2 proof of concept study in idiopathic thrombocytopenic purpura (ITP) for M254, a hyper-sialylated immunoglobulin (hsIgG) designed as a high potency alternative to intravenous immunoglobulin (IVIg) to remediate limitations of that therapeutic approach; and
- Overview of Momenta's research platform and additional pipeline expansion opportunities.

The live webcast of this event will be accessible through the "Investors" section of the company's website located at www.momentapharma.com. A webcast replay of the presentation will be posted on the Momenta website approximately two hours after the event.

About Momenta Pharmaceuticals
Momenta is a biotechnology company with a validated innovative scientific platform focused on discovering and developing novel therapeutics to treat rare, immune-mediated diseases. Momenta's lead product candidate, M281, is a potentially best-in-class anti-FcRn antibody; M254, is a hyper-sialylated human immunoglobulin (hsIgG) designed as a high potency alternative to intravenous immunoglobulin (IVIg); and M230 (CSL730), is a potential first-in-class novel recombinant Fc multimer being developed in collaboration with CSL. Momenta also has a focused pipeline of two biosimilar candidates: M710, a proposed biosimilar to EYLEA® being developed in collaboration with Mylan, and M923, Momenta's wholly-owned proposed biosimilar to HUMIRA®. Momenta's two FDA-approved complex generic products, enoxaparin sodium injection and Glatopa® (glatiramer acetate injection), are marketed by its collaboration partner, Sandoz.

To learn more about Momenta, please visit www.momentapharma.com, which does not form a part of this press release.

Forward-Looking Statements

Statements in this press release regarding management's future expectations, beliefs, intentions, goals, strategies, plans or prospects, are forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, including but not limited to statements about the timing and content of our proposed presentations regarding Phase 1 data and indications for Phase 2 studies for M28, plans regarding timing of studies for M254, and potential research and pipeline expansions. Forward-looking statements may be identified by words such as "believe," "continue," "plan to", "potential," "will," and other similar words or expressions, or the negative of these words or similar words or expressions. Such forward-looking statements involve known and unknown risks, uncertainties and other important factors, including the risk that we are unable to make such presentations or that the content of such presentations will change; and those referred to under the section "Risk Factors" in the Company's Quarterly Report on Form 10-Q for the quarter ended June 30, 2018 filed with the Securities and Exchange Commission, as well as other documents that may be filed by the Company from time to time with the Securities and Exchange Commission. As a result of such risks, uncertainties and factors, the Company's actual results may differ materially from any future results, performance or achievements discussed in or implied by the forward-looking statements contained herein. The Company is providing the information in this press release as of this date and assumes no obligations to update the information included in this press release or revise any forward-looking statements, whether as a result of new information, future events or otherwise.

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