Clearside Biomedical to Report Second Quarter 2018 Financial Results on August 8, 2018 – Conference Call to Follow

August 1, 2018

ALPHARETTA, Ga., Aug. 01, 2018 (GLOBE NEWSWIRE) -- Clearside Biomedical, Inc. (NASDAQ:CLSD), a biopharmaceutical company dedicated to developing treatments that restore and preserve vision for people with serious eye diseases, announced today that its second quarter 2018 financial results will be released at approximately 7:00 a.m. ET on Wednesday, August 8, 2018. Following the release, Clearside will host a live conference call and webcast at 8:30 a.m. ET to discuss its financial results and provide a general business and strategic review.

The live webcast and a replay may be accessed by visiting the "Investor Relations" section at www.clearsidebio.com. Alternately, please call (844) 263-8310 (U.S.) or (213) 358-0959 (international) to participate in the live conference call. The conference ID number for the live call is 6390517. Please dial in approximately 10 minutes prior to the call. An archive of the webcast will be available until September 8, 2018.

About Clearside

Clearside Biomedical, Inc. is a biopharmaceutical company dedicated to developing treatments that restore and preserve vision for people with serious eye diseases. Clearside's proprietary suprachoroidal treatment approach offers unprecedented access to the back of the eye where sight-threatening disease often occurs. The company's unique platform for eye disease treatments is inherently flexible and intended to work with established medicines, new formulations of medicines, as well as future innovations. Clearside's pipeline includes advanced and pre-clinical product candidates in diseases where macular edema is a common complication, including uveitis, retinal vein occlusion ("RVO") and diabetic macular edema ("DME"). Clearside's most advanced program is in non-infectious uveitis and it expects to submit a New Drug Application to the U.S. Food and Drug Administration for use of suprachoroidal CLS-TA for the treatment of macular edema associated with non-infectious uveitis by the end of 2018. The company is also conducting two ongoing Phase 3 trials of suprachoroidal CLS-TA with an intravitreal anti-VEGF agent in patients with RVO. In addition, Clearside recently announced positive topline results from a Phase 2 clinical trial of suprachoroidal CLS-TA used with EYLEA® (aflibercept) in patients with DME, and is continuing to analyze additional data from the trial as it becomes available. Clearside is headquartered in Alpharetta, GA. For more information, please visit http://www.clearsidebio.com. Follow @clearsidebio on Twitter and Linkedin.

Contacts:

Stephen Kilmer
Investor Relations
(678) 430-8206
stephen.kilmer@clearsidebio.com

Charles Deignan
Chief Financial Officer
(678) 270-4005
charlie.deignan@clearsidebio.com

Primary Logo

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