Dicerna to Report Second Quarter 2020 Financial Results and Host Virtual Research and Development Event on Aug. 6, 2020

July 24, 2020

LEXINGTON, Mass.--(BUSINESS WIRE)--Jul. 24, 2020--

Dicerna Pharmaceuticals, Inc. (Nasdaq: DRNA) (the “Company” or “Dicerna”), a leading developer of investigational ribonucleic acid interference (RNAi) therapeutics, today announced that the Company will release its second quarter 2020 financial results before the market open on Thursday, Aug. 6, 2020. In lieu of a call to discuss its quarterly results, Dicerna will host a virtual research and development (R&D) event on Aug. 6 from 10:00 a.m. to noon ET.

Highlights of this year’s event will include:

- **Phase 1 proof-of-concept data for RG6346**, an investigational GalXC™ RNAi treatment candidate for chronic hepatitis B virus (HBV) infection in development with Roche
- **Multidose data from the PHYOX™3 trial of nedosiran**, an investigational GalXC RNAi treatment candidate for primary hyperoxaluria (PH)
- **First preclinical data** on Dicerna’s RNAi technology in extrahepatic tissues

In addition to Dicerna’s management, featured presenters will include:

- **Man-Fung Yuen, D.Sc., M.D., Ph.D.**, Chair Professor of Gastroenterology and Hepatology, Li Shu Fan Medical Foundation, Professor in Medicine, Chief of the Division of Gastroenterology and Hepatology, Deputy Head, Department of Medicine, the University of Hong Kong
- **Bernd Hoppe, M.D.**, Professor of Pediatrics, Head of the German Hyperoxaluria Center in Bonn, Germany, former Head of the Division of Pediatric Nephrology in the Department of Pediatrics at the University Hospital in Bonn, Germany and Vice President, Global Medical Affairs at Dicerna

Virtual Event Details

The virtual presentation will be webcast beginning at 10:00 a.m. ET on Thursday, Aug. 6, 2020, and may be accessed by visiting the “Investors & Media” section of the Dicerna website, www.dicerna.com. A conference line can be accessed by dialing (800) 708-4539 or +1 (847) 619-6396 and referencing conference ID 49860522. A replay of the webcast will be archived on Dicerna’s website following the conclusion of the live event.

About Dicerna Pharmaceuticals, Inc.

Dicerna Pharmaceuticals, Inc. (Nasdaq: DRNA) is a biopharmaceutical company focused on discovering, developing and commercializing medicines that are designed to leverage ribonucleic acid interference (RNAi) to selectively silence genes that cause or contribute to disease. Using our proprietary RNAi technology platform, GalXC™, Dicerna is committed to developing RNAi-based therapies with the potential to treat both rare and more prevalent diseases. By silencing disease-causing genes, Dicerna’s GalXC platform has the potential to address conditions that are difficult to treat with other modalities. Dicerna’s first gene targets are located within hepatocytes; however, continually innovating, Dicerna is also exploring new applications of RNAi technology beyond the liver, targeting additional tissues and enabling new therapeutic applications. In addition to our own pipeline of core discovery and clinical candidates, Dicerna has established collaborative relationships with some of the world’s leading pharmaceutical companies, including Novo Nordisk A/S, Roche, Eli Lilly and Company, Alexion Pharmaceuticals, Inc., Boehringer Ingelheim International GmbH and Alnylam Pharmaceuticals, Inc. Between Dicerna and our collaborative partners, we currently have more than 20 active discovery, preclinical or clinical programs focused on rare, cardiometabolic, viral, chronic liver and complement-mediated diseases, as well as neurodegeneration and pain. At Dicerna, our mission is to interfere – to silence genes, to fight disease, to restore heath. For more information, please visit www.dicerna.com.

Cautionary Note on Forward-Looking Statements

This press release includes forward-looking statements. Such forward-looking statements are subject to risks and uncertainties that could cause actual results to differ materially from those expressed or implied in such statements. Examples of forward-looking statements include, among others, statements we make regarding: (i) Phase 1 proof-of-concept data for RG6346, an investigational GalXC™ RNAi treatment candidate for chronic hepatitis B virus (HBV) infection in development with Roche; (ii) multidose data from the PHYOX™3 trial of nedosiran, an investigational GalXC RNAi treatment candidate for primary hyperoxaluria (PH), (iii) first preclinical data on Dicerna’s RNAi technology in extrahepatic tissues; (iv) the therapeutic and commercial potential of nedosiran and (v) clinical development timelines and review related to nedosiran and continued alignment on the regulatory pathway to approval. The process by which investigational therapies, such as nedosiran, could potentially lead to an approved product is long and subject to highly significant risks. Applicable risks and uncertainties include those relating to Dicerna’s clinical research and other risks identified under the heading "Risk Factors" included in the Company’s most recent filings on Forms 10-K and 10-Q and in other future filings with the Securities and Exchange Commission. These risks and uncertainties include, among others, the cost, timing and results of preclinical studies and clinical trials and other development activities by us and our collaborative partners; the likelihood of Dicerna’s clinical programs being executed on timelines provided and reliance on the Company’s contract research organizations and predictability of timely enrollment of subjects and patients to advance Dicerna’s clinical trials; the reliance of Dicerna on contract manufacturers to supply its products for research and development and the risk of

supply interruption from a contract manufacturer; the potential for future data to alter initial and preliminary results of early-stage clinical trials; the impact of the ongoing COVID-19 pandemic on our business operations, including the conduct of our research and development activities; the regulatory review and unpredictability of the duration and results of the regulatory review of Investigational New Drug applications (INDs) and Clinical Trial Applications (CTAs) that are necessary to continue to advance and progress the Company's clinical programs; the timing, plans and reviews by regulatory authorities of marketing applications such as New Drug Applications (NDAs) and comparable foreign applications for one or more of Dicerna's product candidates; the ability to secure, maintain and realize the intended benefits of collaborations with partners; market acceptance for approved products and innovative therapeutic treatments; competition; the possible impairment of, inability to obtain, and costs to obtain intellectual property rights; possible safety or efficacy concerns that could emerge as new data are generated in R&D; and general business, financial, and accounting risks and litigation. The forward-looking statements contained in this press release reflect Dicerna's current views with respect to future events, and Dicerna does not undertake and specifically disclaims any obligation to update any forward-looking statements.

View source version on businesswire.com: https://www.businesswire.com/news/home/20200724005179/en/

Media:
Amy Trevvett, Dicerna Pharmaceuticals, Inc.
+1 617-612-6253
atrevvett@dicerna.com

Investors:
Lauren Stival, Dicerna Pharmaceuticals, Inc.
+1 617-514-0461
lstival@dicerna.com

Source: Dicerna Pharmaceuticals, Inc.