Tricida to Present at the 38th Annual J. P. Morgan Healthcare Conference

January 7, 2020

SOUTH SAN FRANCISCO, Calif.--(BUSINESS WIRE)--Jan. 7, 2020-- Tricida, Inc. (Nasdaq: TCDA) today announced that it will present at the 38th Annual J. P. Morgan Healthcare Conference on Tuesday, January 14, 2020 at 9:30 am PT / 12:30 pm ET. Gerrit Klaerner, Ph.D., Tricida's Founder, Chief executive officer and President, will provide a company overview, business update and progress on the company's key initiatives.

A live webcast of the presentation will be accessible on the Tricida website at IR.Tricida.com. An archive of the webcast will be available for 90 days following the presentation.

About Tricida

Tricida, Inc. is a pharmaceutical company focused on the development and commercialization of its drug candidate, veverimer (TRC101), a non-absorbed, orally-administered polymer designed to treat metabolic acidosis in patients with chronic kidney disease (CKD). Metabolic acidosis is a condition commonly caused by CKD that is believed to accelerate the progression of kidney deterioration. It is estimated to pose a health risk to approximately three million patients with CKD in the United States. Tricida is currently conducting its confirmatory postmarketing trial, VALOR-CKD, of veverimer. The Tricida New Drug Application (NDA) for veverimer has been accepted for review by the U.S. Food and Drug Administration (FDA) under the Accelerated Approval Program. The FDA has assigned a Prescription Drug User Fee Act (PDUFA) goal date of August 22, 2020.

For more information about Tricida, please visit www.Tricida.com.

Cautionary Note on Forward-Looking Statements

This press release includes forward-looking statements, including for example, statements about the assigned PDUFA goal date of August 22, 2020, and the potential availability of the Accelerated Approval Program. Forward-looking statements involve known and unknown risks, uncertainties, assumptions and other factors that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. These risks and uncertainties include, among others, that we may not be able to achieve upcoming milestones; the cost, timing and results of clinical trials and other studies; that many drug candidates that have completed Phase 3 trials do not become approved drugs on a timely or cost effective basis or at all; there can be no assurance that the FDA would approve an NDA under the Accelerated Approval Program, or at all, and even if approval for a drug is obtained, there can be no assurance that it will be adopted in the market or accepted as a benefit to patients and healthcare providers; possible safety and efficacy concerns; and that we completely rely on third-party suppliers to manufacture our clinical drug supply. These and other factors that may affect our future results of operations are identified and described in more detail in our filings with the Securities and Exchange Commission (SEC), including our Quarterly Reports on Form 10-Q and our Annual Report on Form 10-K. The forward-looking statements contained in this press release reflect Tricida's current views with respect to future events, and Tricida does not undertake and specifically disclaims any obligation to update any forward-looking statements.

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