Tricida to Report First Quarter 2019 Financial Results and Host Conference Call and Webcast on Wednesday, May 8, 2019

April 30, 2019

SOUTH SAN FRANCISCO, Calif.--(BUSINESS WIRE)--Apr. 30, 2019-- Tricida, Inc. (Nasdaq: TCDA) today announced that it will report its first quarter 2019 financial results after the close of market on Wednesday, May 8, 2019. Tricida will host a conference call and webcast at 4:30 pm Eastern Time to discuss its financial results and business progress. The call or webcast may be accessed as follows:

Tricida First Quarter 2019 Conference Call

Wednesday, May 8, 2019
4:30 pm Eastern Time

Webcast:	IR.Tricida.com
Dial-in:	**(877) 377-5478**
International:	(629) 228-0740
Conference ID:	**5661605**

A replay of the webcast will be available on Tricida's website approximately two hours following the completion of the call and will be available for up to 90 days following the presentation.

About Tricida

Tricida, Inc. is a pharmaceutical company focused on the development and commercialization of its drug candidate, TRC101, a non-absorbed, orally-administered polymer designed to treat metabolic acidosis in patients with 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 has successfully completed all of the clinical trials that it planned to complete prior to submission of an NDA to the U.S. Food and Drug Administration (FDA). Tricida plans to submit an NDA, in the second half of 2019, seeking approval of TRC101 through the FDA's Accelerated Approval Program.

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 our ability to submit an NDA for TRC101 under the FDA's 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, the timing of Tricida's NDA submission; 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 will find that our clinical trials have provided evidence of clinical benefit; 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 TRC101. 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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Source: Tricida, Inc.

Jackie Cossmon, IRC
Tricida, Inc.
Vice President of Investor Relations and Communications
IR@Tricida.com