



Caribou Biosciences to Participate in Upcoming Investor Conferences

August 31, 2022

BERKELEY, Calif., Aug. 31, 2022 (GLOBE NEWSWIRE) -- [Caribou Biosciences, Inc.](#) (Nasdaq: CRBU), a leading clinical-stage CRISPR genome-editing biopharmaceutical company, today announced the company's participation in the following investor conferences:

- **Citi 17th Annual BioPharma Conference**, Boston
Panel participation on September 8, 2022, 11:20 am ET
[Webcast](#)
- **H.C. Wainwright 24th Annual Global Investment Conference**, New York City
Corporate presentation on September 13, 2022, 10:00 am ET
[Webcast](#)
- **Morgan Stanley 20th Annual Global Healthcare Conference**, New York City
Fireside chat on September 13, 2022, 4:40 pm ET
[Webcast](#)
- **Jefferies Cell and Genetic Medicines Summit**, New York City
September 29-30, 2022; see [Events](#) page for details

For more information, visit the [Events](#) page on Caribou's website. Webcasts will be available on the Caribou website for 30 days after the event.

About Caribou's Novel Next-Generation CRISPR Platform

CRISPR genome editing uses easily designed, modular biological tools to make DNA changes in living cells. There are two basic components of Class 2 CRISPR systems: the nuclease protein that cuts DNA and the RNA molecule(s) that guide the nuclease to generate a site-specific, double-stranded break, leading to an edit at the targeted genomic site. CRISPR systems are capable of editing unintended genomic sites, known as off-target editing, which may lead to harmful effects on cellular function and phenotype. In response to this challenge, Caribou has developed CRISPR hybrid RNA-DNA guides (chRDNA; pronounced "chardonnays") that direct substantially more precise genome editing compared to all-RNA guides. Caribou is deploying the power of its Cas12a chRDNA technology to carry out high efficiency multiple edits, including multiplex gene insertions, to develop CRISPR-edited therapies.

About Caribou Biosciences, Inc.

Caribou Biosciences is a clinical-stage CRISPR genome-editing biopharmaceutical company dedicated to developing transformative therapies for patients with devastating diseases. The company's genome-editing platform, including its proprietary Cas12a chRDNA technology, enables superior precision to develop cell therapies that are specifically engineered for enhanced persistence. Caribou is advancing a pipeline of off-the-shelf CAR-T and CAR-NK cell therapies for the treatment of patients with hematologic malignancies and solid tumors.

For more information about Caribou, visit www.cariboubio.com and follow the company @CaribouBio.

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