



FINANCIAL TEAR SHEET

CORPORATE PROFILE

CRISPR Therapeutics is a leading gene-editing company focused on the development of transformative gene-based medicines for serious diseases using its proprietary CRISPR/Cas9 gene-editing platform. CRISPR/Cas9 is a revolutionary technology that allows for precise, directed changes to genomic DNA. Our multi-disciplinary team of world-class researchers and drug developers is working to translate this technology into breakthrough human therapeutics in a number of serious diseases. Our lead program, CTX001, aims to treat Sickle cell disease and β -thalassemia using an ex vivo approach. We are also pursuing oncology indications with our allogeneic CAR-T platform and liver, lung, and neuromuscular indications using in vivo approaches. In addition, we have established strategic collaborations with Bayer AG and Vertex Pharmaceuticals to develop CRISPR-based therapeutics in other diseases with high unmet need. We have licensed the foundational CRISPR/Cas9 patent estate for human therapeutic use from our scientific founder, Dr. Emmanuelle Charpentier, who co-invented the application of CRISPR/Cas9 for gene editing. We are headquartered in Zug, Switzerland with R&D operations in Cambridge, Massachusetts, USA and select business operations in London, United Kingdom.

PRIMARY IR CONTACT

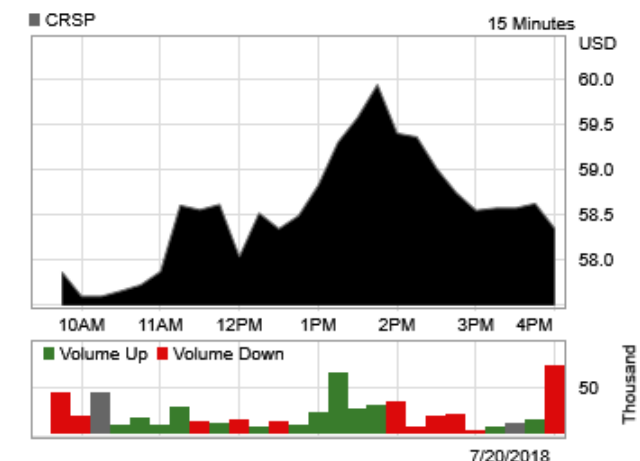
Chris Brinzey
Westwicke Partners
Phone: 339-970-2843
E-mail:
chris.brinzey@westwicke.com

STOCK PERFORMANCE

CRSP (Common Stock)

Exchange	NASDAQ (US Dollar)
Price	\$58.35
Change (%)	▼ 0.04 (0.07%)
Volume	687,323
52 Week Low	\$16.15
Market Cap	\$2,723,825,555
Rolling EPS	-1.63
PE Ratio	0
Shares Outstanding	46,680,815

Data as of 07/20/18 4:00 p.m. ET



RECENT HEADLINES & EVENTS

Jun-19-2018 - 9:05 a.m.
CRISPR Therapeutics, Intellia Therapeutics and
Caribou Biosciences Announce Grant of U.S.
Patent for CRISPR/Cas9 Genome Editing

There are currently no events scheduled.

Jun-05-2018 - 8:00 a.m.
CRISPR Therapeutics to Participate in Three June
Conferences

May-30-2018 - 4:30 p.m.
CRISPR Therapeutics and Vertex Provide Update
on FDA Review of Investigational New Drug
Application for CTX001 for the Treatment of Sickle
Cell Disease

SEC FILINGS

Filing Date	Form
07/05/18	4
06/19/18	4
06/18/18	4
06/15/18	4

Data provided by Nasdaq. Minimum 15 minutes delayed. View Attributions and Sources