# Revance to Release First Quarter 2020 Financial Results on Thursday, May 7, 2020

April 30, 2020

*Conference Call Scheduled for Thursday, May 7, 2020 at 4:30 p.m. ET*

NEWARK, Calif.--(BUSINESS WIRE)--Apr. 30, 2020-- Revance Therapeutics, Inc. (Nasdaq: RVNC), a biotechnology company focused on innovative aesthetic and therapeutic offerings, including its next-generation neuromodulator product, DaxibotulinumtoxinA for Injection, today announced that the company will release first quarter 2020 financial results on Thursday, May 7, 2020 after the close of market. Revance will host a corresponding conference call and a live webcast at 1:30 p.m. PT/4:30 p.m. ET on the same day to discuss the results and provide a business and pipeline update.

Individuals interested in listening to the conference call may do so by dialing (855) 453-3827 for domestic callers, or (484) 756-4301 for international callers and reference conference ID: 4799936; or from the webcast link in the investor relations section of the company's website at: www.revance.com.

A replay of the call will be available beginning May 7, 2020 at 4:30 p.m. PT/7:30 p.m. ET to May 8, 2020 at 4:30 p.m. PT/7:30 p.m. ET. To access the replay, dial (855) 859-2056 or (404) 537-3406 and reference conference ID: 4799936. The live webcast can be access here and will be available in the investor relations section on the company's website for 30 days following the completion of the call.

In light of reduced call center resources during this time of required social-distancing, Revance requests listeners not planning on participating in Q&A to listen to the live webcast rather than dialing in by phone.

**About Revance Therapeutics, Inc.**

Revance Therapeutics, Inc. is a biotechnology company focused on innovative aesthetic and therapeutic offerings, including its next-generation neuromodulator product, DaxibotulinumtoxinA for Injection. DaxibotulinumtoxinA for Injection combines a proprietary stabilizing peptide excipient with a highly purified botulinum toxin that does not contain human or animal-based components. Revance has successfully completed a Phase 3 program for DaxibotulinumtoxinA for Injection in glabellar (frown) lines and is pursuing U.S. regulatory approval in 2020. Revance is also evaluating DaxibotulinumtoxinA for Injection in the full upper face, including glabellar lines, forehead lines and crow's feet, as well as in three therapeutic indications - cervical dystonia, adult upper limb spasticity and plantar fasciitis. Beyond DaxibotulinumtoxinA for Injection, Revance gained exclusive rights to commercialize TEOXANE SA's Resilient Hyaluronic Acid® (RHA®) line of fillers in the U.S., the first and only range of FDA-approved dermal fillers for correction of dynamic facial wrinkles and folds. Revance has also begun development of a biosimilar to BOTOX®, which would compete in the existing short-acting neuromodulator marketplace. Revance is dedicated to making a difference by transforming patient experiences. For more information or to join our team visit us at www.revance.com.



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