

October 18, 2017



Cerecor, Inc. Regains Compliance with NASDAQ Minimum Bid Price Requirement

BALTIMORE, MD -- (Marketwired) -- 10/18/17 -- Cerecor, Inc. (NASDAQ: CERC), a biopharmaceutical company developing innovative drug candidates for patients with neurologic and neuropsychiatric disorders, today announced that it received a letter from the Nasdaq Listing Qualifications Staff (the "**Staff**") confirming that it has regained compliance with the minimum bid price requirement under Nasdaq Listing Rule 5550(a)(2) (the "**Rule**") for continued listing on The Nasdaq Capital Market. In February 2017, the Staff notified the Company that its common stock failed to maintain a minimum bid price of \$1.00 over the previous 30 consecutive business days as required by the Rule. Since then, the Staff has determined that from October 3 to October 16, 2017, the closing bid price of the Company's common stock has been at \$1.00 per share or greater. In a letter dated October 17, 2017, the Staff notified Cerecor that it has regained compliance with the Rule and the matter is now closed.

About Cerecor

Cerecor is a biopharmaceutical company that is developing innovative drug candidates to make a difference in the lives of patients with neurologic and psychiatric disorders. Cerecor's lead drug candidate is CERC-301, which Cerecor currently intends to explore as a novel treatment for orphan neurological indications. Cerecor is also developing two pre-clinical stage compounds, CERC-611 and CERC-406. Cerecor's portfolio of product candidates is summarized below:

CERC-301 belongs to a class of compounds known as antagonists of the N-methyl-D-aspartate ("NMDA") receptor, a receptor subtype of the glutamate neurotransmitter system that is responsible for controlling neurological adaptation. Given its selective mechanism of action and tolerability profile, Cerecor believes CERC-301 may be well suited to address unmet medical needs in other neurological indications. Cerecor is now embarking on a pre-clinical and clinical program to explore the use of CERC-301 in orphan neurological conditions.

CERC-611 is a potent and selective transmembrane AMPA receptor regulatory proteins γ 8-dependent α -amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid ("AMPA") receptor antagonist, which Cerecor plans to develop as an adjunctive therapy for the treatment of partial-onset seizures with or without secondarily generalized seizures in patients with epilepsy.

CERC-406 is a brain penetrant catechol-O-methyltransferase inhibitor with potential pro-cognitive activity. Cerecor believes CERC-406 may have the potential to be developed for the treatment of residual cognitive impairment symptoms.

The Company plans both to evaluate its current portfolio for potential new indications, focusing on orphan neurologic diseases, and to identify potential new product candidates that could be in-licensed.

For more information about the Company and its products, please visit www.cerecor.com or contact

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Source: Cerecor, Inc.