Vanda Receives Negative Opinion for Marketing Authorization from the European Medicines Agency on Fanaptum™ for the Treatment of Schizophrenia

November 10, 2017

WASHINGTON, Nov. 10, 2017 /PRNewswire/ -- Vanda Pharmaceuticals Inc. (Vanda) (NASDAQ: VNDA) announced today the refusal of the marketing authorization application (MAA) of Fanaptum™ (oral iloperidone tablets) for the treatment of schizophrenia in adult patients in the European Union. In July 2017, the Committee for Medicinal Products for Human Use (CHMP) adopted a negative opinion, recommending the refusal of the Fanaptum™ MAA. Vanda requested a re-examination of the initial opinion. After considering the grounds for this request, the CHMP re-examined the opinion and confirmed the refusal of the Fanaptum™ MAA on November 9, 2017.

About Vanda Pharmaceuticals Inc.
Vanda is a global biopharmaceutical company focused on the development and commercialization of innovative therapies to address high unmet medical needs and improve the lives of patients. For more on Vanda Pharmaceuticals Inc., please visit www.vandapharma.com.

About Fanapt®

For full U.S. Prescribing Information for Fanapt®, including indication, Boxed Warnings and Important Safety Information, visit our Web site at www.fanapt.com.

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