



Alnylam to Webcast Conference Call Discussing First Quarter 2019 Financial Results

April 23, 2019

CAMBRIDGE, Mass.--(BUSINESS WIRE)--Apr. 23, 2019-- [Alnylam Pharmaceuticals, Inc.](http://www.alnylam.com) (Nasdaq: ALNY), the leading RNAi therapeutics company, announced today that it will report financial results for the first quarter ending March 31, 2019 on Wednesday, May 1, 2019, before the U.S. financial markets open.

Management will provide an update on the Company and discuss first quarter 2019 results as well as expectations for the future via conference call on Wednesday, May 1, 2019 at 8:30 am ET. To access the call, please dial 800-289-0438 (domestic) or 323-794-2423 (international) five minutes prior to the start time and refer to conference ID 4935120. A replay of the call will be available beginning at 11:30 am ET on the day of the call. To access the replay, please dial 888-203-1112 (domestic) or 719-457-0820 (international) and refer to conference ID 4935120.

A live audio webcast of the call will be available on the Investors section of the Company's website, www.alnylam.com. An archived webcast will be available on the Alnylam website approximately two hours after the event.

About Alnylam Pharmaceuticals

Alnylam (Nasdaq: ALNY) is leading the translation of RNA interference (RNAi) into a whole new class of innovative medicines with the potential to transform the lives of people afflicted with rare genetic, cardio-metabolic, hepatic infectious, and central nervous system (CNS)/ocular diseases. Based on Nobel Prize-winning science, RNAi therapeutics represent a powerful, clinically validated approach for the treatment of a wide range of severe and debilitating diseases. Founded in 2002, Alnylam is delivering on a bold vision to turn scientific possibility into reality, with a robust discovery platform. Alnylam's first U.S. FDA-approved RNAi therapeutic is ONPATTRO® (patisiran) lipid complex injection available in the U.S. for the treatment of the polyneuropathy of hereditary transthyretin-mediated (hATTR) amyloidosis in adults. In the EU, ONPATTRO is approved for the treatment of hATTR amyloidosis in adults with stage 1 or stage 2 polyneuropathy. Alnylam has a deep pipeline of investigational medicines, including five product candidates that are in late-stage development. Looking forward, Alnylam will continue to execute on its "*Alnylam 2020*" strategy of building a multi-product, commercial-stage biopharmaceutical company with a sustainable pipeline of RNAi-based medicines to address the needs of patients who have limited or inadequate treatment options. Alnylam employs over 1,000 people worldwide and is headquartered in Cambridge, MA. For more information about our people, science and pipeline, please visit www.alnylam.com and engage with us on Twitter at [@Alnylam](https://twitter.com/Alnylam) or on [LinkedIn](https://www.linkedin.com/company/alnylam).

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