



## News Release

### **Puma Biotechnology to Present at B. Riley Securities’ 3<sup>rd</sup> Annual Oncology Conference**

**LOS ANGELES, Calif., Jan. 12, 2023** –Puma Biotechnology, Inc. (NASDAQ: PBYI), a biopharmaceutical company, announced that Alan H. Auerbach, Chairman, Chief Executive Officer, President and Founder of Puma, will provide an overview of the Company at 11:00 a.m. PST/2:00 p.m. EST on January 19 at the virtual B. Riley Securities’ 3<sup>rd</sup> Annual Oncology Conference. A live webcast of Mr. Auerbach’s presentation will be accessible via registration on the event website at <https://brileyoncology22.sequireevents.com/>. A replay of the presentation will be available at the same address.

#### **About Puma Biotechnology**

Puma Biotechnology, Inc. is a biopharmaceutical company with a focus on the development and commercialization of innovative products to enhance cancer care. Puma in-licensed the global development and commercialization rights to PB272 (neratinib, oral), PB272 (neratinib, intravenous) and PB357. Neratinib, oral was approved by the U.S. Food and Drug Administration in 2017 for the extended adjuvant treatment of adult patients with early stage HER2-overexpressed/amplified breast cancer, following adjuvant trastuzumab-based therapy, and is marketed in the United States as NERLYNX® (neratinib) tablets. In February 2020, NERLYNX was also approved by the FDA in combination with capecitabine for the treatment of adult patients with advanced or metastatic HER2-positive breast cancer who have received two or more prior anti-HER2-based regimens in the metastatic setting. NERLYNX was granted marketing authorization by the European Commission in 2018 for the extended adjuvant treatment of adult patients with early stage hormone receptor-positive HER2-overexpressed/amplified breast cancer and who are less than one year from completion of prior adjuvant trastuzumab-based therapy. NERLYNX is a registered trademark of Puma Biotechnology, Inc.

In September 2022, Puma entered into an exclusive license agreement for the development and commercialization of the anti-cancer drug alisertib, a selective, small molecule, orally administered inhibitor of aurora kinase A. Initially, Puma intends to focus the development of alisertib on the treatment of small cell lung cancer and breast cancer.

Further information about Puma Biotechnology may be found at <https://www.pumabiotechnology.com>.

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