



## **BIO-PATH HOLDINGS REPORTS FIRST QUARTER 2019 FINANCIAL RESULTS**

*Conference Call to be Held Today at 8:30 A.M. ET*

**HOUSTON—May 16, 2019** – Bio-Path Holdings, Inc., (NASDAQ:BPTH), a biotechnology company leveraging its proprietary DNAbilize® antisense RNAi nanoparticle technology to develop a portfolio of targeted nucleic acid cancer drugs, today announced its financial results for the first quarter ended March 31, 2019 and provided an update on recent corporate developments.

“During the first quarter, we continued to execute on our strategic initiatives to build our DNAbilize® technology and advance our clinical development programs. We were delighted to present data from preclinical studies supporting the potential of BP1003, a novel liposome-incorporated STAT3 oligodeoxynucleotide inhibitor for the treatment of pancreatic cancer, non-small cell lung cancer (NSCLC) and acute myelogenous leukemia (AML) at the American Association for Cancer Research (AACR) Annual Meeting before an audience of world-leading oncologists,” said Peter Nielsen, President and Chief Executive Officer of Bio-Path Holdings. “We look forward to completing Investigational New Drug (IND) enabling studies in 2019 through early 2020, and to file an IND application for a Phase 1 study of BP1003 for the treatment of pancreatic cancer in 2020.”

### **Recent Corporate Highlights**

- **Presented Preclinical Data at American Association for Cancer Research Annual Meeting 2019.** In April 2019, Bio-Path presented data from preclinical studies supporting the potential of BP1003, a novel liposome-incorporated STAT3 oligodeoxynucleotide inhibitor, for the treatment of pancreatic cancer, non-small cell lung cancer (NSCLC) and acute myelogenous leukemia (AML). These data were presented in a poster at the American Association for Cancer Research (AACR) Annual Meeting 2019 in Atlanta, GA.
- **Raised \$21.3 Million During the First Quarter.** In March 2019, Bio-Path issued 712,910 shares of its common stock at a price of \$25.95 per share, for gross proceeds of approximately \$18.5 million in one transaction. In addition, Bio-Path completed two smaller offerings during the first quarter.

### **Financial Results for the First Quarter Ended March 31, 2019**

- The Company reported a net loss of \$1.5 million, or \$0.89 per share, for the three months ended March 31, 2019, compared to a net loss of \$1.9 million, or \$3.38 per share, for the three months ended March 31, 2018.

- Research and development expenses for the three months ended March 31, 2019 decreased to \$0.4 million, compared to \$0.9 million for the three months ended March 31, 2018 primarily due to decreased clinical trial expenses as we modified operations between Stage 1 and Stage 2 of our Phase 2 clinical trial in AML to include venetoclax combination treatment with prexigebersen.
- General and administrative expenses for the three months ended March 31, 2019 increased to \$1.1 million, compared to \$1.0 million for the three months ended March 31, 2018 primarily due to increased legal fees and insurance costs.
- As of March 31, 2019, the Company had cash of \$19.3 million, compared to \$1.0 million at December 31, 2018. Net cash used in operating activities for the three months ended March 31, 2019 was \$2.0 million compared to \$1.7 million for the comparable period in 2018. Net cash provided by financing activities for the three months ended March 31, 2019 was \$20.3 million.

### **Conference Call and Webcast Information**

Bio-Path Holdings will host a conference call and webcast today at 8:30 a.m. ET to review these first quarter 2019 financial results and to provide a general update on the Company. To access the conference call please dial (844) 815-4963 (domestic) or (210) 229-8838 (international) and refer to the conference ID 4956638. A live audio webcast of the call and the archived webcast will be available in the Media section of the Company's website at [www.biopathholdings.com](http://www.biopathholdings.com).

### **About Bio-Path Holdings, Inc.**

Bio-Path is a biotechnology company developing DNabilize<sup>®</sup>, a novel technology that has yielded a pipeline of RNAi nanoparticle drugs that can be administered with a simple intravenous transfusion. Bio-Path's lead product candidate, prexigebersen (BP1001, targeting the Grb2 protein), is in a Phase 2 study for the treatment of blood cancers and is in the process of filing an IND for a Phase 1 clinical trial for solid tumors. The Company is also developing BP1002, which targets the Bcl-2 protein and is expected to be evaluated for the treatment of lymphoma and solid tumors. In addition, BP1003, a novel liposome-incorporated STAT3 antisense oligodeoxynucleotide developed by Bio-Path as a specific inhibitor of STAT3, is expected to enter Phase 1 studies in 2020.

For more information, please visit the Company's website at <http://www.biopathholdings.com>.

### **Forward-Looking Statements**

This press release contains forward-looking statements that are made pursuant to the safe harbor provisions of the federal securities laws. These statements are based on management's current expectations and accordingly are subject to uncertainty and changes

in circumstances. Any express or implied statements contained in this press release that are not statements of historical fact may be deemed to be forward-looking statements. Any statements that are not historical facts contained in this release are forward-looking statements that involve risks and uncertainties, including Bio-Path's ability to raise needed additional capital on a timely basis in order for it to continue its operations, Bio-Path's ability to have success in the clinical development of its technologies, the timing of enrollment and release of data in such clinical studies and the accuracy of such data, limited patient populations of early stage clinical studies and the possibility that results from later stage clinical trials with much larger patient populations may not be consistent with earlier stage clinical trials, the maintenance of intellectual property rights, risks relating to maintaining Bio-Path's listing on the Nasdaq Capital Market and such other risks which are identified in Bio-Path's most recent Annual Report on Form 10-K, in any subsequent quarterly reports on Form 10-Q and in other reports that Bio-Path files with the Securities and Exchange Commission from time to time. These documents are available on request from Bio-Path Holdings or at [www.sec.gov](http://www.sec.gov). Bio-Path disclaims any intention or obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise.

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