

Mustang Bio Reports Third Quarter 2019 Financial Results and Recent Corporate Highlights

NEW YORK, Nov. 12, 2019 (GLOBE NEWSWIRE) -- Mustang Bio, Inc. ("Mustang") (NASDAQ: MBIO), a clinical-stage biopharmaceutical company focused on translating today's medical breakthroughs in cell and gene therapies into potential cures for hematologic cancers, solid tumors and rare genetic diseases, today announced financial results and recent corporate highlights for the third quarter ended September 30, 2019.

Manuel Litchman, M.D., President and Chief Executive Officer of Mustang, said, "We continue to be pleased with the progress of our gene therapy and CAR T programs. In the third quarter, MB-107 (lentiviral gene therapy) was granted the Regenerative Medicine Advanced Therapy (RMAT) designation by the U.S. Food and Drug Administration (FDA), which we hope will translate into expedited development and approval of this very important and potentially curative lentiviral gene therapy for the treatment of X-linked severe combined immunodeficiency (XSCID), also known as bubble boy disease. We look forward to a strong presence at the upcoming 61st American Society of Hematology (ASH) Annual Meeting, where additional data pertaining to MB-107 (lentiviral gene therapy) and information pertaining to MB-106 (CD20-targeted CAR T cell therapy) will be presented. With a strong balance sheet and the continued advancement of our gene therapy and CAR T programs, we are well positioned to execute on our goals for the rest of 2019 and into 2020."

Financial Results:

- As of September 30, 2019, Mustang's cash, cash equivalents, short-term investments (certificates of deposit) and restricted cash totaled \$73.3 million, compared to \$83.1 million as of June 30, 2019 and \$34.6 million as of December 31, 2018, a decrease of \$9.8 million for the guarter and an increase of \$38.7 million year-to-date.
- Research and development expenses were \$7.3 million for the third quarter of 2019, compared to \$5.3 million for the third quarter of 2018. Non-cash, stock-based compensation expenses included in research and development were \$0.7 million for third quarter of 2019, compared to \$0.7 million for the third quarter of 2018.
- Research and development expenses from license acquisitions totaled \$0.7 million for the third guarter of 2019, compared to \$1.0 million for the third guarter of 2018.
- General and administrative expenses were \$2.0 million for the third quarter of 2019, compared to \$1.3 million for the third quarter of 2018. Non-cash, stock-based compensation expenses included in general and administrative expenses were \$0.4

- million for the third quarter of 2019, compared to \$0.2 million for the third quarter of 2018.
- Net loss attributable to common stockholders was \$10.2 million, or \$0.25 per share, for the third quarter of 2019, compared to \$7.5 million, or \$0.28 per share, for the third quarter of 2018.

Recent Corporate Highlights:

- In July 2019, the FDA granted Orphan Drug Designation to MB-102 (CD123-targeted CAR T cell therapy) for the treatment of acute myeloid leukemia (AML).
- In August 2019, Mustang announced that the FDA approved its Investigational New Drug (IND) application to initiate a multi-center Phase 1/2 clinical trial of MB-102 in AML, blastic plasmacytoid dendritic cell neoplasm and high-risk myelodysplastic syndrome.
- In August 2019, MB-107, a lentiviral gene therapy for the treatment of X-linked severe combined immunodeficiency (XSCID), also known as bubble boy disease, was granted the RMAT designation by the FDA.
- Also in August 2019, Mustang entered into a license agreement with CSL Behring for the Cytegrity[™] stable producer cell line, which will be used to produce the viral vector for the MB-107 lentiviral gene therapy program for the treatment of XSCID.
- Additionally in August 2019, the California Institute for Regenerative Medicine (CIRM) granted City of Hope \$9.28 million to fund an ongoing Phase 1 clinical trial of MB-103 (HER2-targeted CAR T cell therapy) for the treatment of HER2-positive breast cancer with brain metastases.
- In September 2019, Mustang announced that City of Hope opened and has initiated patient treatments in a Phase 1 clinical trial of MB-105 (PSCA-targeted CAR T cell therapy) for the treatment of prostate cancer.
- In October 2019, Mustang announced that City of Hope received \$4.1 million in grant awards for a clinical trial of MB-101 (IL13Rα2-targeted CAR T cell therapy) in combination with nivolumab (commercial name: Opdivo®) and ipilimumab (commercial name: Yervoy®) in patients with recurrent malignant glioma. The trial, which is now enrolling patients, is the first human study to combine IL13Rα2-targeted CAR T cell therapy with checkpoint inhibitors, as well as the first to locally deliver CAR T cells with combination treatment with systemic nivolumab treatment.
- Also in October 2019, Mustang announced that the first participant was dosed in a Phase 1 clinical trial to determine the safety and efficacy of MB-108 (oncolytic virus C134), an attenuated herpes simplex virus type 1, in recurrent glioblastoma multiforme.
- In November 2019, Mustang announced that MB-107 lentiviral gene therapy for XSCID clinical data were accepted for <u>oral</u> and <u>poster</u> presentations at the 61st ASH Annual Meeting in December 2019. In addition, Mustang's collaborator Fred Hutchinson Cancer Research Center will present a <u>poster</u> about the design of the ongoing Phase 1/2 clinical trial investigating the safety and efficacy of MB-106 CD20-targeted CAR T cell therapy for high-risk B-cell non-Hodgkin lymphomas.

About Mustang Bio

Mustang Bio, Inc. ("Mustang") is a clinical-stage biopharmaceutical company focused on translating today's medical breakthroughs in cell and gene therapies into potential cures for hematologic cancers, solid tumors and rare genetic diseases. Mustang aims to acquire rights to these technologies by licensing or otherwise acquiring an ownership interest, to fund

research and development, and to outlicense or bring the technologies to market. Mustang has partnered with top medical institutions to advance the development of CAR T and CRISPR/Cas9-enhanced CAR T therapies across multiple cancers, as well as a lentiviral gene therapy for XSCID. Mustang is registered under the Securities Exchange Act of 1934, as amended, and files periodic reports with the U.S. Securities and Exchange Commission. Mustang was founded by Fortress Biotech, Inc. (NASDAQ: FBIO). For more information, visit www.mustangbio.com.

Forward-Looking Statements

This press release may contain "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934, each as amended. Such statements include, but are not limited to, any statements relating to our growth strategy and product development programs and any other statements that are not historical facts. Forward-looking statements are based on management's current expectations and are subject to risks and uncertainties that could negatively affect our business, operating results, financial condition and stock value. Factors that could cause actual results to differ materially from those currently anticipated include: risks relating to our growth strategy; our ability to obtain, perform under and maintain financing and strategic agreements and relationships; risks relating to the results of research and development activities; risks relating to the timing of starting and completing clinical trials; uncertainties relating to preclinical and clinical testing; our dependence on third-party suppliers; our ability to attract, integrate and retain key personnel; the early stage of products under development; our need for substantial additional funds; government regulation; patent and intellectual property matters; competition; as well as other risks described in our SEC filings. We expressly disclaim any obligation or undertaking to release publicly any updates or revisions to any forward-looking statements contained herein to reflect any change in our expectations or any changes in events, conditions or circumstances on which any such statement is based, except as required by law.

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MUSTANG BIO, INC.

Condensed Balance Sheets (\$ in thousands, except for share and per share amounts)

	Se	eptember 30,	December 31,		
	2019 (Unaudited)			2018	
ASSETS					
Current Assets:					
Cash and cash equivalents	\$	72,794	\$	16,469	
Short-term investments (certificates of deposit)		-		17,604	
Other receivables - related party		55		-	
Prepaid expenses and other current assets		1,513		1,052	
Total current assets		74,362		35,125	
Property, plant and equipment, net		6,486		6,465	
Fixed assets - construction in process		1,011		393	
Restricted cash		500		500	
Other assets		250		271	
Operating lease right-of-use asset, net		1,226		-	
Total Assets	\$	83,835	\$	42,754	
LIABILITIES AND STOCKHOLDERS' EQUITY Current Liabilities:					
Accounts payable and accrued expenses	\$	5,770	\$	5,381	
Payables and accrued expenses - related party		284		236	
Operating lease liabilities - short-term		239		-	
Total current liabilities		6,293		5,617	
Deferred rent payable		-		741	
Notes payable		13,185		-	
Operating lease liabilities - long-term		1,909		-	
Total Liabilities		21,387		6,358	
Commitments and Contingencies					
Stockholders' Equity					
Preferred stock (\$0.0001 par value), 2,000,000 shares authorized, 250,000 shares of Class A preferred stock issued and outstanding as of September 30, 2019 and December 31, 2018 Common Stock (\$0.0001 par value), 85,000,000 shares authorized Class A common shares, 862,392 and 1,000,000 shares issued and outstanding as of September 30, 2019 and December 31,		-		-	
2018, respectively					

Total Liabilities and Stockholders' Equity	\$ 83,835	\$ 42,754	
Total Stockholders' Equity	 62,448	 36,396	
Accumulated deficit	 (109,250)	(79,070)	
Additional paid-in capital	171,694	113,378	
Common stock issuable, 0 and 709,314 shares as of September 30, 2019 and December 31, 2018, respectively	-	2,085	
Common shares, 39,578,371 and 26,610,183 shares issued an outstanding as of September 30, 2019 and December 31, 2018 respectively	4	3	

MUSTANG BIO, INC. Condensed Statements of Operations (\$ in thousands, except for share and per share amounts) (Unaudited)

	For the three months ended September 30,				For the nine months ended September 30,				
-		2019		2018		2019	2018		
Revenue - related party	\$	-	\$	-	\$	-	\$	50	
Operating expenses:									
Research and development		7,309		5,316		21,092		13,165	
Research and development – licenses acquired		700		1,000		1,350		1,075	
General and administrative		1,987		1,340		7,520		5,133	
Total operating expenses		9,996		7,656		29,962		19,373	
Loss from operations		(9,996)		(7,656)	-	(29,962)		(19,323)	
Other income (expense)									
Interest income		406		138		945		431	
Interest expense		(578)				(1,163)		-	
Total other income (expense)		(172)		138		(218)		431	
Net Loss	\$	(10,168)	\$	(7,518)	\$	(30,180)	\$	(18,892)	
Net loss per common share outstanding, basic and diluted	\$	(0.25)	\$	(0.28)	\$	(0.87)	\$	(0.70)	

Weighted average number of common shares outstanding, basic and diluted

39,875,209

27,146,721

34,752,938

26,871,505



Source: Mustang Bio, Inc.