MEI Pharma to Report 2023 Fiscal Year End Financial Results and Provide Corporate Overview on September 26, 2023

September 21, 2023

SAN DIEGO--(BUSINESS WIRE)--Sep. 21, 2023-- MEI Pharma, Inc. (NASDAQ: MEIP), a clinical-stage pharmaceutical company focused on advancing new therapies for cancer, today announced that the Company will release its 2023 fiscal year end financial results after the close of the U.S. financial markets on September 26, 2023. Company management will host a conference call and webcast that day at 5:00 p.m. ET to discuss the results and provide a corporate overview.

Conference Call & Webcast Information

- When: September 26, 2023, 5:00 p.m. ET
- Dial-in: 1-833-974-2378 (United States) or 1-412-317-5771 (International)
- Please ask to join into the MEI Pharma earnings call

Please join the conference call at least 10 minutes early to register. You can access the live webcast here or under the investor relations section of MEI's website at: www.meipharma.com. A replay of the conference call will be archived for at least 30 days after the call.

About MEI Pharma

MEI Pharma, Inc. (Nasdaq: MEIP) is a pharmaceutical company focused on developing potential new therapies for cancer. MEI Pharma's portfolio of drug candidates includes clinical stage candidates with differentiated mechanisms of action intended to address unmet medical needs and deliver improved benefit to patients in combination with other therapeutic options. For more information, please visit www.meipharma.com. Follow us on X (formerly Twitter) @MEI_Pharma and on LinkedIn.

Forward-Looking Statements

Certain information contained in this press release that are not historical in nature are "forward-looking statements" within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995 including, without limitation, statements regarding: the potential, safety, efficacy, and regulatory and clinical progress of our product candidates, including the anticipated timing for initiation of clinical trials and release of clinical trial data and our expectations surrounding potential regulatory submissions, approvals and timing thereof, our business strategy and plans; the sufficiency of our cash, cash equivalents and short-term investments to fund our operations. You should be aware that our actual results could differ materially from those contained in the forward-looking statements, which are based on management's current expectations and are subject to a number of risks and uncertainties, including, but not limited to our failure to successfully commercialize our product candidates; the availability or appropriateness of utilizing the FDA's accelerated approval pathway for our product candidates; final data from our pre-clinical studies and completed clinical trials may differ materially from reported interim data from ongoing studies and trials; costs and delays in the development and/or FDA approval, or the failure to obtain such approval, of our product candidates; uncertainties or differences in interpretation in clinical trial results; uncertainty regarding the impact of rising inflation and the increase in interest rates as a result; potential economic downturn; activist investors; our inability to maintain or enter into, and the risks resulting from, our dependence upon collaboration or contractual arrangements necessary for the development, manufacture, commercialization, marketing, sales and distribution of any products; competitive factors; our inability to protect our patents or proprietary rights and obtain necessary rights to third party patents and

intellectual property to operate our business; our inability to operate our business without infringing the patents and proprietary rights of others; general economic conditions; the failure of any products to gain market acceptance; our inability to obtain any additional required financing; technological changes; government regulation; changes in industry practice; and one-time events. We do not intend to update any of these factors or to publicly announce the results of any revisions to these forward-looking statements. Under U.S. law, a new drug cannot be marketed until it has been investigated in clinical studies and approved by the FDA as being safe and effective for the intended use.

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David A. Walsey
MEI Pharma
Tel: 858-369-7104
investor@meipharma.com

Joele Frank, Wilkinson Brimmer Katcher
Dan Katcher / Aaron Palash
212-355-4449
MEIP-jf@joelefrank.com

Source: MEI Pharma, Inc.