



I-MAB
BIOPHARMA

I-Mab to Hold Investor Call to Present In-Depth Phase 1 Clinical Data on Highly Differentiated CD73 Antibody Uliledlimab

June 2, 2021

SHANGHAI and GAITHERSBURG, Md., June 2, 2021 /PRNewswire/ -- I-Mab (the "Company") (Nasdaq: IMAB), a clinical stage biopharmaceutical company committed to the discovery, development and commercialization of novel biologics, today announced that it will hold a call with investors on Monday, June 7 at 8:00 a.m. ET to provide an in-depth clinical data analysis of its U.S. phase 1 study of uliledlimab in combination with atezolizumab (TECENTRIQ®) in patients with advanced cancers.

I-Mab Conference Call Information

Investors and analysts are invited to join the conference call on June 7 at 8:00 a.m. ET via Zoom:

Meeting URL: <https://zoom.us/j/99293081819?pwd=aHJjQ0o4Uk9kUEtzeHl1dngzUGREUT09>
Meeting ID: 992 9308 1819
Password: 942873

About Uliledlimab (TJD5)

Uliledlimab (TJD5) is a differentiated, humanized antibody against CD73, an ecto-enzyme expressed on stromal cells and tumors that converts extracellular adenosine monophosphate (AMP) to adenosine. Adenosine in turn binds to adenosine receptors on relevant immune cells and inhibits anti-tumor immune responses in tumor microenvironment. Uliledlimab is expected to offer clinical benefit by suppressing tumor growth in concert with checkpoint therapies such as PD-(L)1 antibodies. Uliledlimab is effective in anti-tumor activities through a unique intra-dimer binding, leading to differentiated and favorable functional properties as evident in preclinical studies.

About I-Mab

I-Mab (Nasdaq: IMAB) is an innovation-driven global biotech company focusing on discovery, development and soon commercialization of novel and highly differentiated biologics in immuno-oncology therapeutic area. The Company's mission is to bring transformational medicines to patients around the world through drug innovation. I-Mab's globally competitive pipeline of more than 15 clinical and pre-clinical stage drug candidates is driven by its internal R&D capability and global licensing partnerships, based on the Company's unique Fast-to-Proof-of-Concept and Fast-to-Market pipeline development strategies. The Company is now rapidly progressing from a clinical stage biotech company to a fully integrated global biopharmaceutical company with cutting-edge global R&D capabilities, a world-class GMP manufacturing facility and commercialization capability. I-Mab has established its global footprint in Shanghai (headquarters), Beijing, Hangzhou and Hong Kong in China, and Maryland and San Diego in the United States. For more information, please visit <http://ir-i-mabbiopharma.com> and follow I-Mab on [LinkedIn](#), [Twitter](#) and [WeChat](#).

For more information, please contact:

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