NEWS RELEASE

Alpine Immune Sciences to Present Updated Clinical Data for Povetacicept in IgA Nephropathy at the World Congress of Nephrology 2024

4/1/2024

-- The Company will present updated data in a late-breaking poster session for povetacicept in IgA nephropathy patients --

-- The Company will host an investor event the same day concurrently with the poster presentation on April 15th --

SEATTLE--(BUSINESS WIRE)-- **Alpine Immune Sciences, Inc.** (NASDAQ: ALPN), a leading clinical-stage immunotherapy company focused on developing innovative treatments for autoimmune and inflammatory diseases, has announced that the Company will present updated clinical data for povetacicept from RUBY-3, a phase 1b/2a study of povetacicept in autoimmune glomerulonephritis, including IgA nephropathy, in a late breaking poster presentation at the World Congress of Nephrology April 13-16, 2024 in Buenos Aires, Argentina. Concurrently with the poster presentation, the Company will host an investor event on Monday, April 15th.

World Congress of Nephrology Late-Breaking Poster

Date/Time: Monday, April 15, 2024, 5:45 pm local time (GMT-3)/4:45 pm ET/1:45 pm PT

Poster Title: Updated Results from the RUBY-3 study of Povetacicept, an Enhanced Dual BAFF/APRIL Antagonist in IgA Nephropathy

Poster Number: MON-304

Session Name: Late-Breaking Abstract

Presenter: James Tumlin, M.D., Professor of Medicine at Emory University School of Medicine, Founder and CEO of NephroNet Clinical Trials Consortium

Location: Exhibition Hall and Main Foyer, Buenos Aires Convention Center, Buenos Aires, Argentina

Investor Event

Date/Time: Monday, April 15th at 5:45 pm local time (GMT-3)/4:45 pm ET/1:45 pm PT

Link to Webcast: **https://events.q4inc.com/attendee/228395132**

About **Povetacicept (ALPN-303)**

Povetacicept (ALPN-303) is a dual antagonist of the BAFF (B cell activating factor) and APRIL (a proliferation inducing ligand) cytokines, which play key roles in pathogenesis of multiple autoimmune diseases via their roles in the activation, differentiation and/or survival of B cells, particularly antibody-secreting cells, as well as T cells and innate immune cells. Based upon an engineered TACI (transmembrane activator and CAML interactor) domain, povetacicept has exhibited greater potency in preclinical studies versus other inhibitors of BAFF and/or APRIL alone and B cell depletion and has demonstrated initial activity in patients with IgA nephropathy. Povetacicept is in development for multiple autoimmune diseases, including IgA nephropathy and other autoimmune kidney diseases, systemic lupus erythematosus, and autoimmune cytopenias.

About RUBY-3

RUBY-3 (**NCT05732402**) is a multiple ascending dose, multi-cohort, open label, phase 1b/2a study of povetacicept in autoimmune glomerulonephritis, including IgA nephropathy, primary membranous nephropathy, lupus nephritis, and renal ANCA-associated vasculitis, where povetacicept is being administered subcutaneously for up to 104 weeks. Key endpoints include proteinuria, eGFR, renal response, and disease-related autoantibodies.

About Alpine Immune Sciences

Alpine Immune Sciences is committed to leading a new wave of immune therapeutics. With world-class research and development capabilities, a highly productive scientific platform, and a proven management team, Alpine is seeking to create first- or best-in-class multifunctional immunotherapies via unique protein engineering technologies to improve patients' lives. Alpine has entered into strategic collaborations with leading global

biopharmaceutical companies and has a diverse pipeline of clinical and preclinical candidates in development. For more information, visit **www.alpineimmunesciences.com**. Follow @AlpineImmuneSci on **X** and **LinkedIn**.

Forward-Looking Statements

This release contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, Section 21E of the Securities Exchange Act of 1934 and the Private Securities Litigation Reform Act of 1995. These forward-looking statements are not based on historical fact and include statements regarding our platform technology and potential therapies; the timing of and results from clinical trials and preclinical development activities; clinical and regulatory objectives and the timing thereof; expectations regarding the sufficiency of cash, including cash equivalents and restricted cash, and investments to fund our planned operations; our ability to achieve additional milestones in our collaborations and proprietary programs; the progress and potential of our development programs; future development plans and clinical and regulatory milestones and objectives, including the timing and achievement thereof; the efficacy of our clinical trial designs; anticipated enrollment in our clinical trials and the timing thereof; expectations regarding the anticipated reporting of data from our ongoing and planned clinical trials and potential publication of future clinical data; our ability to potentially advance povetacicept directly into a pivotal trial as well as a phase 2 study in systemic lupus erythematosus, and the potential efficacy, safety profile, addressable market, regulatory success and commercial or therapeutic potential of our product candidates. Forward-looking statements generally include statements that are predictive in nature and depend upon or refer to future events or conditions and include words such as “may,” “will,” “should,” “would,” “expect,” “plan,” “intend,” and other similar expressions, among others. These forward-looking statements are based on current assumptions that involve risks, uncertainties, and other factors that may cause actual results, events, or developments to be materially different from those expressed or implied by such forward-looking statements. These risks and uncertainties, many of which are beyond our control, include, but are not limited to: clinical trials may not demonstrate safety and efficacy of any of our product candidates; our ongoing discovery and preclinical efforts may not yield additional product candidates; our discovery stage and preclinical programs may not advance into the clinic or result in approved products; any of our product candidates may fail in development, may not receive required regulatory approvals, or may be delayed to a point where they are not commercially viable; we may not achieve additional milestones in our proprietary or partnered programs; the impact of competition; adverse conditions in the general domestic and global economic markets; we may be unable to advance povetacicept directly into a pivotal trial or a phase 2 study in systemic lupus erythematosus; the impact of pandemics, or other related health crises on our business, research and clinical development plans and timelines and results of operations, including the impact on our clinical trial sites, collaborators, and contractors who act for or on our behalf; as well as the other risks identified in our filings with the Securities and Exchange Commission. These forward-looking statements speak only as of the date hereof and we undertake no obligation to update forward-looking statements, and readers are cautioned not to place undue reliance on such forward-looking

statements.

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