

ASX/Media Announcement

## Immutep Announces the Start of an Investigator-Initiated Phase II Study in COVID-19 Patients

- Investigator-initiated, placebo controlled and randomised Phase II study evaluating efitlagimod alpha (“efti”) in up to 110 COVID-19 patients
- Immutep supplies efti to University Hospital Pilsen, Czech Republic
- Study tests whether the early administration of efti prevents COVID-19 disease progression in adult patients
- Initial interim results are expected to be reported from early 2021

**SYDNEY, AUSTRALIA – 23 October 2020** – [Immutep Limited](#) (ASX: IMM; NASDAQ: IMMP) (“Immutep” or “the Company”), is pleased to announce it has signed a Material Transfer Agreement (“Agreement”) with the University Hospital Pilsen, Czech Republic to enable an investigator-initiated randomised Phase II clinical trial evaluating its lead product candidate efitlagimod alpha (“efti” or “IMP321”) in hospitalised patients with COVID-19. The necessary approvals from the Czech Republic’s State Institute for Drug Control (SUKL-competent authority) and ethics committee have now been obtained, enabling the recruitment of patients to commence immediately. Initial interim results are expected to be reported from early 2021.

The study, called “Efitlagimod Alpha Treatment by immune modulation in **COVID-19** disease” or EAT COVID (EudraCT n° 2020-002009-25), aims to boost a patient’s immune response to prevent the patient from developing severe COVID-19 symptoms that require intensive care and can lead to respiratory failure and death. As an antigen presenting cell (APC) activator, efti could help to control the viral load in hospitalised patients by boosting CD8 effector T cells.

Under the Agreement, Immutep will provide efti at no cost to the University Hospital Pilsen which will fund the study. The trial will be led by Principal Investigator, Professor Martin Matejovic who is the Head of Medical Department at University Hospital Pilsen, Professor of Medicine at University Hospital Pilsen and Charles University Medical School. The trial will also be conducted in collaboration with Dr. Dalibor Sedlacek, Associate Professor of Medicine and Head of the Department of Infectious Diseases, along with Dr. Marek Nalos, Associate Professor of Medicine and Head Medical ICU at Department of Intensive Care Medicine of the Nepean Hospital, Sydney.

The study will be a placebo controlled, 1:1 randomised, double blinded Phase II clinical trial involving up to 110 adult patients hospitalised with COVID-19 at University Hospital Pilsen. Patients will receive subcutaneous injections of efti (10 mg) on days 1, 3 and 7, in addition to standard care. The primary endpoint is the patient’s clinical status at day 15 as per the WHO recommended evaluation scale.

**Immutep CEO, Marc Voigt, said:** “We were delighted to be approached for this study by Dr Marek Nalos and Professor Martin Matejovic. It is highly encouraging that ehti, with its broad mechanism of action, attracts keen interest from clinicians who are willing and able to fund the research and commit their time to further exploring ehti’s potential to fight diseases. While Immutep’s focus remains on cancer and autoimmune disease, this new trial gives us the opportunity to contribute to the global efforts to beat COVID-19. We believe ehti is currently the only APC activator of its kind being evaluated against COVID-19 in a randomised Phase II trial and that the study will contribute to our further understanding of its potential in infectious diseases in general.”

**Immutep CSO and CMO, Dr Frederic Triebel said:** “The most effective anti-viral therapy for COVID-19 is likely an effective CD8 T cell that has been empowered by activated dendritic cells and given a *licence to kill* by these antigen presenting cells (APCs). This effector CD8 killer T cell is then able to detect and eliminate epithelial cells which serve as reservoirs where the virus can replicate. Indeed, the vast majority of patients infected with COVID-19 have strong innate immunity/adaptive immunity crosstalk and are able to fight the infection without the help of modern medicine. In older patients due to immuno-senescence or in patients with compromised innate immunity, restoring APC function with ehti may help drive an improved anti-COVID immune response before it becomes too late and the patient requires intensive care.”

**University Hospital Pilsen, Professor of Medicine and EAT COVID Principal Investigator, Professor Martin Matejovic said:** “Some hospital patients are developing severe COVID-19 disease after 7-10 days which requires intensive care and can be fatal. Ehti’s safety profile to date and its proven efficacy as an APC activator<sup>1</sup> gives us the confidence to initiate a Phase II study for our patients as soon as they are hospitalised and before their condition deteriorates.”

Recruitment will start with an open label safety run-in of 6 patients and then a first cohort of 26 randomized patients, with successive cohorts being recruited following a positive recommendation from the Data Monitoring Committee (DMC), which will evaluate safety and efficacy of the treatment.

Immutep has two existing patent families with claims drawn to the use of ehti in the treatment of infectious disease. For example, in September 2017 Immutep announced the grant of a Japanese patent and, in November 2018, announced the grant of a European patent from these families.

### **Ehti’s Mechanism of Action**

Ehti is a first-in-class antigen presenting cell (APC) activator currently being developed by Immutep for the treatment of cancer. Ehti binds to antigen presenting cells such as dendritic cells, monocytes and macrophages via MHC II molecules. This activates the APCs causing them to become professional antigen presenting cells, thereby presenting antigen to the adaptive immune system. This leads to activation and proliferation of CD4+ (helper) and CD8+ (cytotoxic) T cells. Thus, the aim of ehti is to “push the gas” on the body’s innate and

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<sup>1</sup> [https://en.wikipedia.org/wiki/APC\\_Activator](https://en.wikipedia.org/wiki/APC_Activator)

adaptive immune systems making it a suitable candidate to evaluate not only for the treatment of cancer, but also for the treatment of infectious diseases such as COVID-19.

### **About University Hospital Pilsen**

The University Hospital Pilsen provides patients with basic, specialized and so-called highly-specialized medical care in all areas, with the exception of burn medicine, bone marrow transplantation in children, the most complex organ transplants and heart surgery in children. In some fields, University Hospital Pilsen provides health care for residents of adjacent regions, especially southern and northern Bohemia, Karlovy Vary and parts of central Bohemia. But, for example, in the field of artificial insemination, bone marrow transplantation, liver surgery, prostate surgery or kidney tumors, the importance of the University Hospital Pilsen is nationwide. University hospital ranks among the top hospitals in the Czech republic and represents key opinion leader in many fields of clinical medicine. The hospital also has close links with the Faculty of Medicine, Charles University in Pilsen.

### **About Immutep**

Immutep is a globally active biotechnology company that is a leader in the development of LAG-3 related immunotherapeutic products for the treatment of cancer and autoimmune disease. Immutep is dedicated to leveraging its technology and expertise to bring innovative treatment options to market for patients and to maximize value to shareholders. Immutep is listed on the Australian Securities Exchange (IMM), and on the NASDAQ (IMMP) in the United States.

Immutep's current lead product candidate is efitlagimod alpha ("efti" or "IMP321"), a soluble LAG-3 protein (LAG-3lg) based on the LAG-3 immune control mechanism. This mechanism plays a vital role in the regulation of the T cell immune response. Efti is currently in a Phase IIb clinical trial as a chemoimmunotherapy for metastatic breast cancer termed AIPAC (clinicaltrials.gov identifier NCT02614833); a Phase II clinical trial being conducted in collaboration with Merck & Co., Inc., Kenilworth, NJ, USA (known as "MSD" outside the United States and Canada) referred to as TACTI-002 to evaluate a combination of efti with KEYTRUDA® (pembrolizumab) in several different solid tumours (clinicaltrials.gov identifier NCT03625323); a Phase I clinical trial being conducted in collaboration with Merck KGaA, Darmstadt, Germany and Pfizer Inc. referred to as INSIGHT-004 to evaluate a combination of efti with avelumab (clinicaltrials.gov identifier NCT03252938); and a Phase I combination therapy trial in metastatic melanoma termed TACTI-mel (clinicaltrials.gov identifier NCT02676869).

Additional LAG-3 products, including antibodies, for immune response modulation in autoimmunity and cancer are being developed by Immutep's large pharmaceutical partners. Immutep is also developing an agonist of LAG-3 (IMP761) for autoimmune disease.

Further information can be found on the Company's website [www.immutep.com](http://www.immutep.com) or by contacting:

### **Australian Investors/Media:**

Catherine Strong, Citadel-MAGNUS  
+61 (0)406 759 268; [cstrong@citadelmagnus.com](mailto:cstrong@citadelmagnus.com)

**U.S. Media:**

Tim McCarthy, LifeSci Advisors

+1 (212) 915.2564; [tim@lifesciadvisors.com](mailto:tim@lifesciadvisors.com)

This announcement was authorised for release by the board of Immutep Limited.