Congress of the United States Washington, DC 20515

August 2, 2024

The Honorable Joseph R. Biden, Jr. President of the United States The White House 1600 Pennsylvania Avenue NW Washington, D.C. 20500

Dear President Biden:

We write today to express our sense of urgency to address the post-traumatic stress disorder (PTSD) and suicide epidemic among Veterans and servicemembers, and the grave need for new and better treatment options like MDMA-assisted therapy.

As you know, between 17 and 44 Veterans take their lives each and every day, and nearly 150,000 have done so since 9/11 – more than 21 times as many American lives lost in warzones over that same time. PTSD and suicide have been exponentially more lethal for those who served in uniform than combat.

Despite billions of federal dollars spent on addressing PTSD and mental health, the number of Veteran lives lost to suicide has not decreased. This is due in part to the fact that the FDA has not approved any new treatments for PTSD in nearly 25 years. This astounding lack of innovation has undoubtedly contributed to our rising Veteran suicide rates over the past two decades.

Fortunately, hope is on the horizon. Promising new treatments like MDMA-assisted therapy (MDMA-AT) have the clinically proven potential to bring Veterans who have long suffered with PTSD a treatment that works. MDMA-AT has shown enormous promise in privately funded research for decades and was officially designated a "Breakthrough Therapy" by the FDA in 2017.

Under this designation, the FDA has worked closely with the drug's sponsor to design clinical trials that included strategies to ensure the study's validity and safety, and which ultimately proved MDMA-AT's remarkable efficacy.

In Phase three confirmatory trials, 71.2% of participants no longer qualified for a PTSD diagnosis after three eight-hour sessions of MDMA-AT. Moreover, 86.5% of participants experienced "clinically significant" improvements in their PTSD symptoms. These results are nearly double those of existing PTSD treatments, rendering MDMA-AT the most effective PTSD treatment ever developed.

We understand that the FDA has a statutory obligation to carefully weigh both the risks and benefits of any new drug application. This duty is undoubtedly more complicated when dealing with a novel treatment like MDMA-assisted therapy, which calls on the FDA to consider not only the risks and benefits of a novel drug, but also the unique combination of a pharmaceutical when combined with psychotherapy interventions.

It is not our intent to urge the FDA in any direction related to its decision, which must be based solely on the data and evidence. However, we are aware that as this application has made its way through the regulatory review process, certain groups and individuals have voiced criticism of the application. It is our understanding that while these critics may be well-intentioned, their criticism is not necessarily reflective of the science, but rather their personal ideological beliefs and biases related to the medicalization of substances like MDMA.

While all Americans are free to voice their views on this and other topics of public interest, scientific evidence should not be ignored in favor of those who have been on a mission to discredit this promising treatment at all costs – especially when that would come at the cost of our servicemembers and Veterans.

Thousands of Veterans suffering from PTSD continue to take their lives each year. Current treatments clearly are not working well enough, and our Veterans can no longer wait.

If the data and evidence show that MDMA-AT is safe and effective at relieving the suffering that many Veterans face, then we owe it to them and the millions of others who are living with PTSD to make this treatment option available to them.

We appreciate your consideration of our views on this matter and look forward to the FDA's decision on this application.

Sincerely,

Jack Bergman Member of Congress

Morgan Juttre Member of Congress

J. Luis Correa Member of Congress

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David G. Valadao Member of Congress

Dan Crenshaw Member of Congress

Mariannette & Thicker Macker

Mariannette Miller-Meeks, M.D. Member of Congress

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Magazine

Member of Congress

David J. Trone Member of Congress

long Lambon

Doug Lamborn Member of Congress

Susan Wild Member of Congress

Jim Costa Member of Congress

Patrick Ryan Member of Congress

Mark Pocan

Member of Congress

Jernah Vm Orden

Derrick Van Orden Member of Congress

Bill Huizenga Member of Congress

Jared Mosk witz_

Member of Congress

1. Cho

Tom McClintock Member of Congress

Darren Soto Member of Congress

Michael C. Burgess, M.D. Member of Congress

Michelle Steel Member of Congress

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Don Bacon Member of Congress

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Brittany Pettersen Member of Congress

Brian Fitzpatrick Member of Congress

Brian J. Mast Member of Congress

5W

Pete Sessions Member of Congress

Ruben Gallego Member of Congress

Kevin Kiley Member of Congress

Tony Cardenes

Tony Cárdenas Member of Congress

in Swelwell

Eric Swalwell Member of Congress

Sanford D. Bishop, Jr.

Member of Congress

Som

Julia Brownley Member of Congress

and Mace

Nancy Mace Member of Congress

Jan Schakowsky Member of Congress

O.C.L

Salud Carbajal Member of Congress

Jasmine Member of Congress

Mikie Sherrill Member of Congress

Unette N. Clarke

Wrette D. Clarke Member of Congress

Robert Garcia Member of Congress

landia Tenney

Claudia Tenney Member of Congress

Jame hber o Congress

Lois Frankel Member of Congress

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Neal P. Dunn, M.D. Member of Congress

Member of Congress

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David Rouzer Member of Congress

Alexandria Ocasio-Cortez Member of Congress

Young Kim Member of Congress

abert Menenley

Robert J. Menendez Member of Congress

High & Menherge

Abigail Davis Spanberger Member of Congress

MU

Rich McCormick, MD, MBA Member of Congress

Austin Scott Member of Congress

Jason Crow Member of Congress

Adriano Espaillat Member of Congress

Marilyn Strickland Member of Congress

Cabe Amo

Member of Congress

Bolad & W. ulmai

Robert J. Wittman Member of Congress

LM

Doug LaMalfa Member of Congress

Elissa Slotkin Member of Congress

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Jonathan L. Jackson Member of Congress

CC:

The Honorable Denis McDonough Secretary U.S. Department of Veterans Affairs 810 Vermont Avenue, N.W. Washington, D.C. 20420

The Honorable Xavier Becerra Secretary U.S. Department of Health and Human Services 200 Independence Ave SW Washington DC 20201

The Honorable Robert M. Califf, M.D. Commissioner U.S. Food and Drug Administration 10903 New Hampshire Avenue Silver Spring, MD 20993