



March 1, 2018

The Honorable Scott Y. Nishimoto  
House Committee on Judiciary  
Hawaii State Capitol, Conference Room 325  
Honolulu, HI 96813

Dear Chair Nishimoto and Members of the House Committee on Judiciary:

On behalf of the Epilepsy Foundation and our local affiliate, Epilepsy Foundation of Hawaii, we urge you to support House Bill 1893, as amended by HD1, which would allow therapies derived from cannabidiol (CBD) and approved by the Food and Drug Administration (FDA) to become available to patients. Access to new therapies is particularly important for the one third of people living with epilepsy who experience intractable or uncontrolled seizures and are living with rare epilepsies, and the many more who experience significant adverse effects from their current medication.

The FDA is currently reviewing at least one CBD derived therapy that shows promise for the treatment of Dravet and Lennox-Gastaut syndromes (LGS), tuberous sclerosis complex (LSC) and potentially other rare epilepsies. This potential treatment option has both Orphan Drug Designation and Fast Track Designation from the FDA and could be approved as soon as summer 2018. After FDA approval, the Drug Enforcement Administration (DEA) would schedule the therapy through administrative action and the medication would become available for patients. However, since CBD is a Schedule I substance under the state drug schedule, state action is needed to ensure proper rescheduling of FDA-approved therapies derived from CBD. Unless Hawaii acts, patients will not have access to these new therapies. This is an issue of creating access to FDA-approved prescription drugs and we strongly urge your support for House Bill 1893, as amended by HD 1.

The Epilepsy Foundation is the leading national voluntary health organization that speaks on behalf of the at least 3.4 million Americans, including nearly 13,000 Hawaii residents, with epilepsy and seizures. We foster the wellbeing of children and adults affected by seizures through research programs, educational activities, advocacy, and direct services. Epilepsy is a medical condition that produces seizures affecting a variety of mental and physical functions. Approximately 1 in 26 Americans will develop epilepsy at some point in their lifetime. There is no "one size fits all" treatment for epilepsy, and about a third of people living with epilepsy suffer from uncontrolled or intractable seizures, with many more living with significant side-effects, despite available treatments. Uncontrolled seizures can lead to disability, injury, and even death.

The Epilepsy Foundation and the Epilepsy Foundation of Hawaii are committed to supporting physician-directed care, and to exploring and advocating for all potential treatment options for epilepsy. Bureaucratic processes should not stand in the way of patients gaining access to proven and potentially lifesaving treatment once they have been reviewed and approved by FDA. We urge your support of House Bill 1893, as amended by HD 1. Please do not hesitate to contact Angela Ostrom, Chief Legal Officer & Vice President Public Policy, at 301-918-3766 or [aostrom@efa.org](mailto:aostrom@efa.org) with any questions or concerns.

Sincerely,

A handwritten signature in blue ink that reads "Naomi Manuel".

Naomi Manuel  
Executive Director  
Epilepsy Foundation of Hawaii

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A handwritten signature in blue ink that reads "Philip M. Gattone".

Philip M. Gattone, M.Ed.  
President & CEO  
Epilepsy Foundation

The Epilepsy Foundation is your unwavering ally on your journey with epilepsy and seizures. The Foundation is a community-based, family-led organization dedicated to improving the lives of all people impacted by seizures.