HOUSE RESOLUTION

URGING COMMUNITY AWARENESS OF AMYOTROPHIC LATERAL SCLEROSIS, ALSO KNOWN AS ALS OR LOU GEHRIG'S DISEASE, AND COMMUNITY SUPPORT FOR FINDING A CURE FOR ALS AND THE PROVISION OF ASSISTANCE TO ALS PATIENTS.

WHEREAS, amyotrophic lateral sclerosis (ALS), also known as Lou Gehrig's disease, is a progressive, fatal neurodegenerative disease that affects nerve cells in the brain and the spinal cord; and

WHEREAS, early symptoms of ALS often include increasing muscle weakness, especially involving the arms and legs, speech, swallowing, or breathing; and

WHEREAS, the progressive degeneration of the motor neurons in an ALS patient eventually leads to death; and

WHEREAS, ALS does not affect a patient's mental capacity, and thus, the patient remains alert and aware of the loss of motor functions and the inevitable outcome of deterioration and death; and

WHEREAS, on average, ALS patients survive only two to five years after the ALS diagnosis is made; and

WHEREAS, ALS has no known cause, means of prevention, or cure; and

WHEREAS, research indicates that military veterans are about twice as likely to develop ALS as compared to those who have not served in the military; and

WHEREAS, the United States Department of Veterans Affairs implemented regulations to establish a presumption of service connection for ALS, thereby presuming that the development of ALS was incurred or aggravated by a veteran's service in the military; and

 WHEREAS, the National ALS Patient Registry, which is administered by the Centers for Disease Control and Prevention, is a congressionally-mandated registry for persons in the United States with ALS, and may be the single largest ALS research project ever created; and

WHEREAS, due to his affliction with ALS, Lou Gehrig, the pride of the Yankees, bid farewell to baseball in a famous speech given at Yankee Stadium on July 4, 1939; and

WHEREAS, 2014 is the 75th Anniversary of that speech, and this anniversary will be observed during the National ALS Association's advocacy gathering in May 2014, in Washington, D.C., with generous support from Major and Minor League Baseball across the country; and

WHEREAS, Hawaii has joined with other groups nationally to increase public awareness of:

(1) The circumstances and experiences which ALS patients face in combatting this fatal disease;

(2) The terrible impact ALS has not only on ALS patients but on their families; and

(3) The research being done to treat and eradicate this rapidly progressive and fatal disease; now, therefore,

 BE IT RESOLVED by the House of Representatives of the Twenty-seventh Legislature of the State of Hawaii, Regular Session of 2014, that this body hereby urges community support for, and awareness in, finding a cure for ALS and the provision of assistance to ALS patients; and

BE IT FURTHER RESOLVED that the community is further urged to support the "Walk to Defeat ALS," scheduled for Saturday, September 6, 2014, at 11:00 a.m., at Kapiolani Regional Park in Honolulu, Hawaii, as a joint community awareness event; and

BE IT FURTHER RESOLVED that the ALS support group and the ALS Association Golden West Chapter in Agoura Hills, California, be recognized for their joint efforts to achieve the goal of a Hawaii ALS Registry; and

- 1 BE IT FURTHER RESOLVED that certified copies of this
- 2 Resolution be transmitted to the ALS Association Golden West
- 3 Chapter and the ALS Association.