THE GENERAL ASSEMBLY OF PENNSYLVANIA

HOUSE RESOLUTION

No. 1099 Session of 2015

INTRODUCED BY BAKER, DEAN, REGAN, BIZZARRO, TOOHIL, VEREB, DIGIROLAMO, KNOWLES, COOK-ARTIS, CAUSER, SCHLOSSBERG, HENNESSEY, O'BRIEN, KIRKLAND, DRISCOLL, SAYLOR, MILLARD, PICKETT, DUSH, SONNEY, WATSON, FREEMAN, KINSEY, LONGIETTI, COHEN, YOUNGBLOOD, D. COSTA, ROSS, McCARTER, WARD, MAHONEY, DELOZIER, READSHAW, MARSICO, GILLEN, MILNE AND PAYNE, OCTOBER 24, 2016

INTRODUCED AS NONCONTROVERSIAL RESOLUTION UNDER RULE 35, OCTOBER 24, 2016

A RESOLUTION

- 1 Recognizing November 12, 2016, as "International CJD Awareness 2 Day" in Pennsylvania.
- 3 WHEREAS, Creutzfeldt-Jakob disease (CJD) is a rare
- 4 neurodegenerative disease categorized as a prion disease; and
- 5 WHEREAS, Prions represent an unusual class of infectious
- 6 agents composed entirely of misfolded prion proteins; and
- 7 WHEREAS, Abnormal prion proteins in the brain and spinal
- 8 fluid trigger significant mental and physical disabilities; and
- 9 WHEREAS, In its early stages, CJD causes failing memory,
- 10 behavioral changes, lack of coordination and visual
- 11 disturbances; and
- 12 WHEREAS, Mental deterioration becomes pronounced as the
- 13 illness progresses; and
- 14 WHEREAS, Involuntary movements, blindness, weakness of
- 15 extremities and coma may occur; and

- 1 WHEREAS, CJD symptoms typically appear in later life; and
- 2 WHEREAS, Individuals with CJD become increasingly dependent
- 3 on caregivers; and
- 4 WHEREAS, CJD is usually fatal within months of onset; and
- 5 WHEREAS, No specific treatment exists to eradicate the
- 6 infectious agent or slow the progression from disability to
- 7 death; and
- 8 WHEREAS, CJD is reported worldwide, distributed without
- 9 geographic clustering at a rate of one person per million per
- 10 year; and
- 11 WHEREAS, Approximately 300 cases per year are diagnosed in
- 12 the United States; and
- WHEREAS, Sporadic CJD, representing approximately 85% of
- 14 cases, occurs with no apparent predisposing factors; and
- 15 WHEREAS, Inherited CJD, representing approximately 10% of
- 16 cases, features a predisposition to mutation of the prion
- 17 protein coding gene; and
- 18 WHEREAS, Acquired CJD is transmitted through the use of
- 19 hormones or tissues obtained from an infected cadaver donor; and
- 20 WHEREAS, Diagnosis of CJD involves clinical and
- 21 neuropathological examination to rule out other diseases; and
- 22 WHEREAS, Conclusive diagnosis follows brain biopsy or autopsy
- 23 only; and
- 24 WHEREAS, Research focuses on treatment options, infection
- 25 control, palliative care and presymptomatic testing; and
- 26 WHEREAS, The Creutzfeldt-Jakob Disease Foundation, Inc., a
- 27 nonprofit volunteer organization, offers a toll-free HelpLine,
- 28 facilitates referrals and support groups and hosts an annual
- 29 family conference for the benefit of affected persons and
- 30 medical professionals; and

- 1 WHEREAS, The House of Representatives joins in urging greater
- 2 awareness, collaboration and support relating to this invariably
- 3 fatal prion disease; therefore be it
- 4 RESOLVED, That the House of Representatives recognize
- 5 November 12, 2016, as "International CJD Awareness Day" in
- 6 Pennsylvania.