## MAINE STATE LEGISLATURE

The following document is provided by the LAW AND LEGISLATIVE DIGITAL LIBRARY at the Maine State Law and Legislative Reference Library http://legislature.maine.gov/lawlib



Reproduced from electronic originals (may include minor formatting differences from printed original)

From: Snyder, Sharon

To: Legislature: All; housesupplements@lists.legislature.maine.gov

**Subject:** House Supplement No. 3

**Date:** Thursday, May 04, 2017 9:21:46 AM

State of Maine
House of Representatives
128th Legislature
First Regular Session
HOUSE ADVANCE JOURNAL AND CALENDAR
http://legislature.maine.gov/legis/house/hcalfr.htm
Thursday, May 4, 2017

## **SUPPLEMENT NO. 3**

## **ORDERS**

(4-1) On motion of Representative KINNEY of Knox, the following Joint Resolution: (H.P. 1089) (Cosponsored by President THIBODEAU of Waldo and Representatives: AUSTIN of Gray, BLACK of Wilton, CRAIG of Brewer, GILLWAY of Searsport, HIGGINS of Dover-Foxcroft, TALBOT ROSS of Portland, TUELL of East Machias, Senator: DAVIS of Piscataquis)

## JOINT RESOLUTION RECOGNIZING MAY 2017 AS WILLIAMS SYNDROME AWARENESS MONTH

WHEREAS, Williams syndrome is a genetic disorder that remains virtually unknown to the general public, educators and many doctors; and

WHEREAS, Williams syndrome affects one in 10,000 people worldwide, including an estimated 20,000 to 30,000 people in the United States; and

WHEREAS, Williams syndrome is characterized by medical and developmental delays, cardiovascular disease and learning disabilities; and

WHEREAS, despite these problems, people with Williams syndrome often have unique abilities, including advanced verbal skills, highly sociable personalities and an affinity for music; and

WHEREAS, children with Williams syndrome need costly ongoing medical care and early interventions, such as speech or occupational therapy, that may not be covered by insurance or state funding; and

WHEREAS, as adults, most people with Williams syndrome need supportive housing to live to their fullest potential; and

WHEREAS, the Williams Syndrome Association sponsors activities that work to increase community awareness and provide financial support for critical research and programs, leading to further insights about the disorder and new treatments for the future; now, therefore, be it

RESOLVED: That We, the Members of the One Hundred and Twenty-eighth Legislature now assembled in the First Regular Session, on behalf of the people we represent, take this opportunity to recognize May 2017 as Williams Syndrome Awareness Month and encourage citizen participation in volunteer activities to increase local community awareness of Williams syndrome and its effects.

\_\_\_\_\_