- 1 HJR20
- 2 171787-1
- 3 By Representative Williams (JD)
- 4 RFD: Rules
- 5 First Read: 10-SEP-15

1	171787-1:n:09/09/2015:JEC/jec LRS2015-2889
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8	URGING AWARENESS OF SPINAL MUSCULAR ATROPHY AND
9	RECOGNIZING AUGUST AS SPINAL MUSCULAR ATROPHY AWARENESS MONT
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11	WHEREAS, Spinal Muscular Atrophy is a debilitating,
12	horrific disease that renders infected individuals physically
13	weak and unable to walk, eat, breathe, or swallow; and
14	WHEREAS, Spinal Muscular Atrophy, which attacks the
15	motor nerve cells in the spinal cord, affects one in 10,000
16	babies; and
17	WHEREAS, about one in every 50 Americans is a
18	genetic carrier, and Spinal Muscular Atrophy is blind to race
19	and gender; and
20	WHEREAS, currently, no approved treatment for this
21	nightmare of a disease exists; and
22	WHEREAS, however, researchers understand the causes
23	of Spinal Muscular Atrophy and what route to take to develop
24	effective therapies; and
25	WHEREAS, scientists are on the verge of major
26	breakthroughs that will strengthen the bodies of infected

children, extend life, and eventually produce a cure; now therefore,

BE IT RESOLVED BY THE LEGISLATURE OF ALABAMA, BOTH

HOUSES THEREOF CONCURRING, That we urge Alabamians to unite in the cause of raising awareness for Spinal Muscular Atrophy and bolstering support for all entities searching for effective treatment and a cure for this disease and recognize August 2015 as Spinal Muscular Atrophy Awareness Month.