THE GENERAL ASSEMBLY OF PENNSYLVANIA

HOUSE RESOLUTION

No. 710

Session of 2010

INTRODUCED BY HICKERNELL, REED, ADOLPH, BAKER, BEAR, BELFANTI, BEYER, BOYD, BRADFORD, CALTAGIRONE, CAUSER, CIVERA, COHEN, CONKLIN, D. COSTA, CREIGHTON, DeWEESE, DIGIROLAMO, DONATUCCI, EVERETT, FAIRCHILD, GABLER, GEORGE, GINGRICH, GODSHALL, GOODMAN, GRUCELA, HELM, HENNESSEY, HESS, HUTCHINSON, KILLION, KORTZ, KOTIK, KULA, MAJOR, MARSICO, MELIO, MILLARD, MURT, MUSTIO, OBERLANDER, O'NEILL, PAYNE, PHILLIPS, QUIGLEY, READSHAW, REICHLEY, ROSS, SAINATO, SANTONI, SAYLOR, SCAVELLO, SIPTROTH, STERN, STEVENSON, SWANGER, TRUE, VULAKOVICH AND YOUNGBLOOD, MARCH 24, 2010

INTRODUCED AS NONCONTROVERSIAL RESOLUTION UNDER RULE 35, MARCH 24, 2010

A RESOLUTION

- 1 Designating the month of July 2010 as "MECP2 Duplication
- 2 Syndrome Awareness Month" in Pennsylvania.
- 3 WHEREAS, MECP2 duplication syndrome is a neurological
- 4 disorder that occurs almost exclusively in boys; and
- 5 WHEREAS, MECP2 duplication syndrome is usually caused by
- 6 duplication of DNA on the Xq28 region of the chromosome; and
- 7 WHEREAS, MECP2 duplication syndrome was not formally
- 8 recognized until 2005; and
- 9 WHEREAS, MECP2 duplication syndrome is most commonly
- 10 inherited in an X-linked manner, meaning that affected males
- 11 have, most commonly, inherited the MECP2 duplication from the
- 12 carrier mother, however, spontaneous duplications have been
- 13 reported, but, to date, no cases of a father transmitting the

- 1 duplication have been reported; and
- 2 WHEREAS, MECP2 is the gene known to cause the neurological
- 3 problems in MECP2 duplication syndrome; and
- 4 WHEREAS, While duplication of MECP2 causes many problems,
- 5 some boys have large duplications that include many other genes,
- 6 and the full extent of phenotypes due to duplication of other
- 7 genes is not completely understood at this time; and
- 8 WHEREAS, MECP2 duplication syndrome is characterized by
- 9 hypotonia, developmental delay, mental retardation, limited or
- 10 absent speech, constipation, reflux, ataxia, progressive
- 11 spasticity, stereotyped movements of hands, teeth grinding,
- 12 recurrent respiratory infections in 80%, epilepsy in 50% and
- 13 developmental regression occurring in some boys; and
- 14 WHEREAS, Recent studies have shown a link between MECP2
- 15 duplication syndrome and autism, and most boys with this
- 16 syndrome do have distinct features of autism; and
- 17 WHEREAS, Patients with MECP2 duplication syndrome have
- 18 multiple neurological deficits, but, thankfully, their brains
- 19 are not static and are highly responsive to proper stimulation
- 20 such as schooling, exercise, music, age-appropriate social
- 21 interactions and related activities; and
- 22 WHEREAS, MECP2 duplication syndrome research is currently
- 23 taking place with efforts to reduce symptoms in mice, in hope
- 24 that there will be effective treatments for humans with this
- 25 disorder soon; therefore be it
- 26 RESOLVED, That the House of Representatives designate the
- 27 month of July 2010 as "MECP2 Duplication Syndrome Awareness
- 28 Month" in Pennsylvania.