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OHIO LEGISLATIVE SERVICE COMMISSION

Office of Research
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Office

H.B. 108
133rd General Assembly

Bill Analysis

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Version: As Reported by House Health

Primary Sponsor: Rep. Patterson

Elizabeth Molnar, Attorney

DETAILED ANALYSIS

Designation	Arthrogryposis Multiplex Congenita Awareness Day ¹
Date	June 30
Other Information	According to the National Institutes of Health's Gene and Rare Diseases Information Center, arthrogryposis multiplex congenita, or AMC, refers to the development – prior to birth – of multiple joint contractures affecting two or more areas of the body. A contracture occurs when a joint becomes permanently fixed in a bent or straightened position, which may impact the joint's function and range of motion and may lead to muscle atrophy. AMC is not a specific diagnosis, but rather a physical symptom associated with several different medical conditions. It is thought that AMC is related to decreased fetal movement during development, which can have many causes, including environmental factors, single gene changes, chromosomal abnormalities, and various syndromes. Treatment depends on the signs and symptoms found in each person, but may include physical therapy, removable splints, exercise, or surgery. ²

¹ R.C. 5.2514.

² U.S. Department of Health and Human Services, National Institutes of Health, National Center for Advancing Translational Sciences, Genetic and Rare Diseases Information Center, *Arthrogryposis Multiplex Congenita*, available at <https://rarediseases.info.nih.gov/diseases/777/arthrogryposis-multiplex-congenita>.

HISTORY

Action	Date
Introduced	02-27-19
Reported, H. Health	10-23-19
