State Senator Hearcel F. Craig
Ohio's 15th Senate District

Senate Health, Human Services and Medicaid Committee
Senate Bill 283 Sponsor Testimony
Designate Cardiac Amyloidosis Recognition Day
December 16, 2020

Chairman Burke, Vice Chair Huffman, Ranking Member Antonio, and members of the Senate Health, Human Services and Medicaid Committee, thank you for the opportunity to testify today in support of Senate Bill 283. Senate Bill 283 would designate November 14 as Cardiac Amyloidosis Recognition Day.

Amyloidosis is a group of diseases where abnormal proteins, or amyloids, form on the body's organs and body tissues.[i] This can lead to progressive organ dysfunction.

Symptoms of cardiac amyloidosis include shortness of breath, swelling of the legs and abdomen, angina, tingling and/or numbness in the hands and feet, carpal tunnel syndrome, and unexplained bruising.[ii] The disease is rare and not widely understood. There are two types of the disease, and both are life-threatening because they can lead to heart failure.[iii] One strain of cardiac amyloidosis almost exclusively impacts people of African descent. In the United States, the most common type of the disease affects African American men in their 60s.[iv]

Currently, cardiac amyloidosis has no established protocol for treatment.[v] Less than 50 percent of cardiologists are familiar with the type of testing used to diagnose the disease.[vi] As a result, many individuals living with this disease go misdiagnosed or undiagnosed.[vii] The average person with the disease lives just two to three more years after receiving a diagnosis.

The purpose of the resolution is to raise awareness for this rare and deadly disease. By providing more information to patients, medical professionals and caregivers, there could be earlier detections, improved treatments and better outcomes. Thank you, Chairman

and Members of the Senate Health committee, for your time. I welcome any questions from the committee at this time.

---

[i] Siddiqi OK, Ruberg FL. Cardiac amyloidosis: an update on pathophysiology, diagnosis and treatment. Trends Cardiovasc Med. 2017;1050-1738.

[ii] https://my.clevelandclinic.org/health/diseases/17855-amyloidosis-attr

[iii] Siddiqi OK, Ruberg FL. Cardiac amyloidosis: an update on pathophysiology, diagnosis and treatment. Trends Cardiovasc Med. 2017;1050-1738.

[iv] Jacobson DR, Alexander AA, Tagoe C, Buxbaum JN. Prevalence of the amyloidogenic transthyretin (TTR) V122I allele in 14 333 African-Americans. Amyloid. 2015;22(3):171-174

[v] Siddiqi OK, Ruberg FL. Cardiac amyloidosis: an update on pathophysiology, diagnosis and treatment. Trends Cardiovasc Med. 2017;1050-1738.

[vi] Pfizer Data on File. Harris Poll Consumer Data. 2018.

[vii]Connors LH, Sam F, Skinner M, et al. Heart failure due to age-related cardiac amyloid disease associated with wild-type transthyretin: a prospective, observational cohort study. Circulation. 2016;133(3):282-290.