

March 28, 2023

The Honorable Joseline A. Pena-Melnyk Chair, House Health and Government Operations Committee Room 241 House Office Building Annapolis, Maryland 21401

RE: Support for SB 515 – Step Therapy Protocol Reform

Dear Chair Pena-Melnyk,

On behalf of the Movement Disorders Policy Coalition, I am writing in support of SB 515. This legislation is critical in ensuring patients – including those with movement disorders – can access the therapies their health care provider prescribes, in a timely and appropriate manner by providing clear exemptions and approval timelines when step therapy is required.

The <u>Movement Disorders Policy Coalition</u> (MDPC) serves as a platform from which stakeholders, including health care providers and patients, can provide input on policy decisions impacting patient-centered care for those living with movement disorders. As a coalition of stakeholder groups across the movement disorders space, MDPC advocates at the federal, state, and health plan levels for key health reforms that increase access to personalized care for patients with movement disorders including tardive dyskinesia, Parkinson's disease, essential tremor, Tourette Syndrome, dystonia, ataxia and Huntington's disease.

Step therapy is a utilization management tool used by health insurance plans to contain health care costs. Sometimes called "fail first," step therapy protocols require a patient to try and fail one or more medications, often at lower cost to the insurer, before he or she can access the medication prescribed by their healthcare provider. These step therapy protocols interfere with the physician-patient relationship, delaying accessing appropriate care, and in turn leading to increased burden on both patients and their health care providers.

People living with movement disorders manage complex conditions and treatment regimens, and already experience significant challenges in everyday life. Timely and effective therapy is paramount to ensuring these patients can successfully manage their disease and have quality of life. Therefore, treatment decisions for these patients should be considered on an individual basis, based on the patient's health care status and clinician's expertise, rather than based on insurer step therapy protocols. Patients with movement disorders need direct, continuous access to the medications that have been demonstrated to treat both the physical and mental health symptoms of their condition.

The course of care prescribed by physicians and other healthcare providers is the foundation of patient-centric care. When health care plans interfere with that process and limit the course of treatment, it jeopardizes the patient's overall health.

Movement Disorders Policy Coalition 2020 K Street NW, Suite 505 Washington, DC 20006 SB 515 would improve patient access by requiring insurance companies to establish a clear and convenient process for patients and providers to request exceptions from step therapy protocols. While the bill does not prohibit the use of step therapy or limit the number of steps that an insurer can require, it does outline specific instances where an exception to step therapy protocols would be granted. Clear guidelines, improved accessibility, and more efficient response times to step therapy appeals will aid in improving access to medications and protecting the provider-patient relationship that is critical to successful care.

On behalf of the Movement Disorders Policy Coalition and our membership, we ask that you support SB 515 and thank you for your leadership on this important issue. If we can provide further details or answer any questions, please reach out to Josie Cooper at jcooper@allianceforpatientaccess.org.

Sincerely,

Josie Cooper

Executive Director

Movement Disorders Policy Coalition

Co-Signing Organizations:

Alliance for Patient Access
Caregiver Action Network
Clinical Neurological Society of America
Depression and Bipolar Support Alliance
HD Reach
National Ataxia Foundation
Parkinson & Movement Disorder Alliance
Parkinson's Foundation

The Michael J. Fox Foundation for Parkinson's Research