March 4, 2020

Representative Bart Rowland
Chair
House Banking and Insurance Committee
702 Capital Ave
Annex Room 416D
Frankfort, KY 40601

Representative Joseph M. Fischer
Vice Chair
House Banking and Insurance Committee
702 Capital Ave
Annex Room 329D
Frankfort, KY 40601

Re: Support for House Bill 72 – a bill related to prescription drug copay assistance

Dear Chairman Rowland and Vice-Chair Fischer:

On behalf of the 1-in-10 individuals in Kentucky with one of the approximately 7,000 known rare diseases, the National Organization for Rare Disorders (NORD) urges you to support House Bill 72, a bill related to prescription drug copay assistance. We urge you to move this bill forward when it comes up in your committee this week.

NORD is a unique federation of voluntary health organizations dedicated to helping people with rare “orphan” diseases and assisting the organizations that serve them. Since 1983, we have been committed to the identification, treatment, and cure of rare disorders through programs of education, advocacy, research, and patient services.

A rare disease is defined as any disorder that affects 200,000 or fewer people in the United States. There are over 7,000 such disorders that have been identified, and over 90 percent of them do not have a treatment approved by the Food and Drug Administration. Even when a treatment exists, however, access is not assured. Due to the innovative nature of many of the treatments for rare diseases and the small populations they are intended to treat, these therapies can be expensive. Thankfully, both charitable and manufacturer-based programs exist to help rare disease patients defray the cost.

As you know, House Bill 72 must apply cost-sharing amounts paid by either the insured or on behalf of the insured by another person for prescription drugs towards a patient’s insurance policy. Cost-sharing includes the coverage limit, copayment, coinsurance, deductible or other out-of-pocket expense requirements associated with an individual’s health plan.

NORD recognizes that the high cost of drugs has a direct impact on patient access. Addressing this and other barriers to care is a priority for NORD. Cost is a primary reason why patients decline to fill their

prescriptions. Further, non-adherence to prescribed medication is responsible for an estimated 125,000 deaths, ten percent of hospitalizations, and hundreds of billions of dollars in costs to the healthcare system per year.\(^2\)\(^3\)\(^4\)

With this in mind, it is critical that payments made by patients, or on behalf of patients, apply toward their deductible and yearly maximum out-of-pocket costs. Without applying payments in this manner, patients will be less likely to be able to meet their deductible and, thus, may quickly exhaust any charitable or manufacturer-based assistance they may have by repeatedly paying for the full cost of the drug. If that happens, patients will be left having to pay the bulk of their deductible, as well as the entirety of their copay or coinsurance, despite having already spent enough to meet their deductible. Patients could be forced to decide between forgoing their critical therapy or facing medical bankruptcy. House Bill 72 would greatly help rare patients and families in Kentucky avoid that tough choice.

Thank you for the opportunity to comment on this legislation. We look forward to continuing to work with you on House Bill 72, to ensure that rare disease patients in Kentucky are supported in their daily struggle to afford their medications. If you have any questions, do not hesitate to contact me at hross@rarediseases.org. Thank you again for your consideration.

Sincerely,

Heidi Ross
Director of State Policy

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Patrick Dunegan
Kentucky Rare Action Network
Volunteer State Ambassador

Jennifer Dunegan
Rare Action Network
Volunteer State Ambassador

CC: House Committee on Banking and Insurance Members