

MEMORANDUM OF SUPPORT

SB560 (Sara Gelser & Tim Knopp)

May 2, 2021

On behalf of the Lupus and Allied Diseases Association and the millions of Oregon residents both directly and indirectly affected by diseases of unmet need, I am expressing our strong support for SB560, An act relating to health insurance cost sharing; requiring each health insurer issuing, delivering, or renewing a policy in this state which provides prescription drug coverage or each pharmacy benefits manager on behalf of such health insurer must apply any amount paid by an insured or by another person on behalf of the insured toward the insured's total contribution to an out-of-pocket maximum or any cost-sharing requirement and applying the Act to all policies, contracts, and health benefit plans issued, delivered, or renewed in the State on or after January 1, 2022.

As an individual who struggles daily to manage multiple autoimmune diseases, I passionately urge you and the members of the House Committee on Health Care to support this important legislation as SB560 provides vital patient access and affordability at a time when our healthcare system is overburdened by the COVID-19 pandemic.

Copay accumulators are a relatively new insurance benefit design being adopted by health insurance plans that prevent patients from using copay cards or coupons to cover their out-of-pocket expenses. This cost-shifting mechanism changes the way an insured individual's out-of-pocket contributions for prescription drugs are calculated. Copay accumulators do not consider any discounts or coupons that the insured person receives from the drug manufacturer when calculating the insured's out-of-pocket expenses and therefore do not count them toward the patient's maximum out-of-pocket limit.

This practice shifts the cost towards patients as the insurance plans are essentially "double dipping" by requiring the individual to pay their normal copay while still receiving a discount or coupon from the drug manufacturer, in turn causing the individual to take longer to satisfy their deductible. This unfair design is especially challenging for individuals with high deductible or high copayment requirements in their health insurance plans.

Given the current global healthcare crisis in which Oregon residents and others throughout the world are experiencing hardship in dealing with the COVID-19 pandemic, individuals already struggling to manage their chronic and rare medical conditions are now facing additional challenges of covering the cost of their medication and accessing appropriate healthcare. Copay coupons and cards provide financial relief to patients through manufacturer programs created to offset the cost sharing associated with certain treatments while assisting patients in meeting their maximum out-of-pocket limits. Copay assistance programs provide patient access to life-sustaining and lifesaving medication that would otherwise be unaffordable while promoting treatment adherence.

Oregon has a longstanding record of providing access to affordable healthcare and developing some of the strongest patient protections in the country. The proposed legislation discontinues the use of copay accumulators statewide by requiring insurance companies and pharmacy benefit managers to apply

price reduction instruments for out-of-pocket expenses when calculating an insured individual's cost-sharing requirement. We urge you to make Oregon the next state to protect its residents from insurmountable out-of-pocket costs and join the seven other states who have passed similar legislation to ensure all copays count toward the deductible and out-of-pocket maximum.

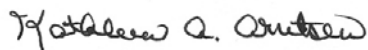
The Lupus and Allied Diseases Association was founded in 1978 and is a national non-profit organization led by people with lupus and allied diseases and their loved ones who are dedicated to ensuring that the patient perspective is included and recognized as an equal stakeholder in the healthcare, regulatory and public policy arenas and across the research continuum. It is our goal to improve access to care and quality of life by fostering collaboration among stakeholders and by wielding the patient voice as a catalyst to advance innovative advocacy, education, awareness and biomedical research initiatives that will identify causes, advance better diagnostics, and discover superior treatments, and cures.

As patient stakeholders who represent patients and loved ones dealing with serious medical conditions on a daily basis who strongly support establishing essential patient protections that improve access to vital therapies and as members of the Oregon All Co-Pays Count Coalition, we passionately urge your support and passage of **SB560** to stop this unfair practice and to help individuals afford the medication they require to remain active. The crucial need for this legislation has only been intensified during the COVID-19 pandemic as countless Oregon families face financial uncertainty. We request that the Senate Committee on Health Care and state legislature support this legislation and stand with patients by prohibiting copay accumulators and requiring health insurance plans to count all dollars towards an insured individual's deductible or out-of-pocket maximum.

We request that you vote YES to support this legislation and stand with patients by prohibiting copay accumulators and requiring health insurance plans to count all dollars towards an insured individual's deductible or out-of-pocket maximum. We must remain vigilant in protecting Oregon residents while promoting unfettered access to vital life-enhancing and lifesaving treatments.

We thank you for the opportunity to provide our unique patient viewpoint and for your support of strong patient safeguards. Please contact me at 315-264-9101 or kathleen@ladainc.org if you have any questions.

Respectfully Submitted-



Kathleen A. Arntsen
President & CEO