June 27, 2019

The Honorable Chuck Grassley
Chairman
United States Senate
Committee on Finance
Washington, DC 20510

The Honorable Ron Wyden
Ranking Member
United States Senate
Committee on Finance
Washington, DC 20510

Dear Chairman Grassley and Ranking Member Wyden:

The undersigned organizations share a commitment to improving the nation’s drug pricing system in ways that will strengthen the health and economic security of current and future Medicare beneficiaries. We appreciate your bipartisan attention to this matter and urge you to include needed reforms to the Part D benefit in any forthcoming prescription drug legislation.

Based on our work with people with Medicare and their families, we know that prescription drug affordability is an ongoing challenge. Our organizations frequently hear from older adults and people with disabilities who are struggling to cover their drug costs and obtain needed medications.

Many people with Medicare live on fixed or limited incomes and cannot keep paying ever-rising health and prescription drug costs. Currently, half of all Medicare beneficiaries—nearly 30 million older adults and people with disabilities—live on $26,200 or less per year, while one quarter have incomes below $15,250 and less than $14,550 in savings.\(^1\) At the same time, health care costs are taking up a larger and more disproportionate share of beneficiaries’ limited budgets. In 2016, nearly 30% of Medicare households spent 20% or more of their income on health care, while only 6% of non-Medicare households did so.\(^2\) Out-of-pocket costs for prescription drugs represent a significant share of this amount, accounting for nearly one out of every five beneficiary health care dollars.\(^3\)

Accordingly, we respectfully ask that in any legislation to address the problem of high and rising prescription drug prices, you include updates to the Medicare Part D benefit structure, appeals process, and low-income programs, as outlined below. These long-overdue improvements would help older adults and people with disabilities better access and afford needed medications, as well as build and maintain their health and financial well-being.

**Medicare Part D Benefit Structure**

The Medicare Part D benefit structure is contributing to the affordability challenges facing many people with Medicare. In particular, the absence of a meaningful cap on beneficiary expenses and lack of an effective reinsurance scheme mean the program is unable to lower drug prices or protect enrollees from these rising costs. We urge you to revisit and improve the benefit’s design. To that end, we recommend the following changes:


Establish an Out-of-Pocket Cap. While the Part D drug benefit does include an out-of-pocket (OOP) limit, it is not a hard cap. Instead, beneficiaries not receiving low-income subsidies (LIS) that help pay Part D premiums and cost sharing must pay 5% of their drug costs indefinitely once they reach the catastrophic coverage level—at which point they’ve already spent thousands of dollars out-of-pocket. In 2017, over 1 million non-LIS Part D enrollees had OOP spending above the catastrophic threshold, with average annual out-of-pocket costs exceeding $3,200. A hard OOP cap would help reduce costs and enhance predictability for enrollees. We oppose increasing the current $5,100 threshold amount by $1,250 up to $6,350 at the end of this year, as prescribed in current law.

Ensure Proportional Liability Among Payers. When contemplating the appropriate liability for each payer in the post-cap period, we encourage you to ensure that both prescription drug plans and drug manufacturers have meaningful and proportional liability for each phase of the Part D benefit. Equitable, balanced incentives would help better control costs for beneficiaries and the program.

Maintain Manufacturer Coverage Gap Discounts in the TrOOP Calculation. Some proposals to restructure the Part D benefit suggest that, in addition to changing the reinsurance liabilities above the catastrophic threshold and establishing an OOP cap, manufacturer coverage gap discounts should no longer count towards true out-of-pocket costs (TrOOP). We strongly oppose excluding manufacturer discounts from TrOOP costs. Even when combined with an OOP cap, this policy would increase costs for many beneficiaries by keeping them in the coverage gap longer.

Simultaneously Address the Root of the Problem. While we support capping OOP costs for Part D enrollees, there must also be efforts to address the underlying problem of high and rising prescription drug prices, such as exploring opportunities for bipartisan agreement on ways the Medicare program can negotiate prices with manufacturers. Consumers should not pay sky-high prices and neither should the Medicare program.

Medicare Part D Appeals Process

The Medicare Part D appeals process is an essential safety valve, for example, allowing access to prescription medications that are not on the plan’s formulary, or are subject to high cost sharing, when formulary or lower-cost alternatives are not appropriate. However, we consistently observe that many Part D enrollees struggle to navigate this overly complex system, resulting in delays in access to needed prescription drugs, abandonment of prescribed medications, reduced adherence to treatment protocols, worse health outcomes, and higher costs. To alleviate these challenges, we encourage you to make the following improvements:

Streamline Medicare Part D Coverage Determinations and Appeals. The multi-step, prolonged Part D exceptions and appeals process proves burdensome and time-consuming for Medicare beneficiaries, pharmacists, plans, and prescribing physicians and can significantly delay access to necessary medications. To begin with, many Part D enrollees are unaware of their right to appeal and do not know

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how to go about initiating the appeals process. Furthermore, Part D enrollees are not provided individualized information or adequate education when refused a medication at the pharmacy counter, resulting in the individual and their physician spending hours trying to obtain this information in order to make a proper exceptions request.

Additionally, rather than being able to appeal the denial at the pharmacy counter, a beneficiary must formally make an exception request, which can require significant time and effort for both the patient and their physician. Only upon receipt of a written denial in response to this request, known as the coverage determination, may the beneficiary request a formal appeal, termed a redetermination. Therefore, we recommend requiring a coverage determination to be provided automatically to a beneficiary at a point-of-sale refusal. Requiring the pharmacy counter refusal to serve as the coverage determination serves the dual purpose of removing a burdensome step for beneficiaries and their doctors while also expediting the appeals process for those who need it. We strongly support the bipartisan Streamlining Part D Appeals Process Act (S. 1861), which would accomplish these goals.

We note with interest a February 4, 2014 letter to CMS Administrator Marilyn Tavenner, signed by every then-member of the Senate Finance Committee. It illustrates clear bipartisan support for this important reform, stating, in part: “We recommend improving the Part D appeals process... For instance, we encourage CMS to explore ways to allow the beneficiary to initiate the appeals process at the pharmacy counter when he/she is first notified the drug is not covered by the part D plan.”6 We urge you to address this long-standing need without delay.

In the interim, we recommend requiring that the existing pharmacy counter refusal notice explain the reason (i.e., prior authorization, step therapy, quantity limits, off-formulary, non-covered, etc.) why the beneficiary is being turned away. This simple, straight-forward, much-needed information would better equip Part D enrollees and their providers to determine the appropriate next steps, whether requesting a coverage determination or pursuing an alternative medication.

**Expand Tiering Exceptions.** We also strongly support the establishment of a cost-sharing exception and appeal process for drugs included on the specialty tier. For all other plan formulary tiers, beneficiaries may request an exception from the plan, asking that the drug be placed on a lower cost-sharing tier—provided that the lower-tier medication is inappropriate for their specific medical needs. No such request is available for specialty tier drugs.

At the same time, spending on these drugs has increased substantially in recent years. According to the Congressional Budget Office (CBO), Part D spending on specialty drugs rose from $8.7 billion in 2010 to $32.8 billion in 2015, accounting for about 30% of Part D spending on prescription drugs that year, up from 13% in 2010.7

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Drugs on the specialty tier represent an increasing cost burden for beneficiaries. We encourage you to work with CMS on implementing an exception and appeal process for the specialty drug tier at the earliest possible time.

**Apply Medicare Part B and Medicaid Standards to Coverage Rules for Off-Label Prescription Drugs in Medicare Part D.** When determining whether to cover an off-label use of an FDA-approved prescription drug, the Medicare Part B program, as well as many private insurance carriers and state Medicaid programs, include in their analysis a consideration of peer-reviewed literature, such as respected medical journals. The Medicare Part D program, however, does not permit reliance on peer-reviewed support at all. Instead, the program automatically denies coverage unless there is a supportive listing in one of three commercially-produced compendia. This inconsistency with standard practice of many other insurers has created serious barriers to accessing effective and sometimes life-saving prescription drugs and has been a source of frustration for providers and beneficiaries alike.

We urge you to correct this misalignment and allow use of peer reviewed literature in Medicare Part D. Specifically, we support amending Section 1927(k)(6) of the Social Security Act (42 U.S.C. 1396r-8(k)(6)), the Part D definition of a “medically accepted indication,” to align the definition with that used in the Medicaid portion of the Social Security Act.

**Medicare Part D Low Income Programs**

The Part D Low-Income Subsidy (LIS), or Extra Help, was designed to address the needs of low-income Medicare beneficiaries, but the program has significant flaws that should be addressed. Congress should consider making the following changes to Extra Help and other low-income assistance programs in order to increase access and affordability:

**Eliminate the Asset Test & Expand Eligibility for LIS.** We support eliminating the asset test and extending the standard LIS benefit to all people under 200% of the federal poverty limit, as outlined in the Medicare Extra Rx HELP Act (S. 691). We also support interim steps to lessen the burden of the asset test, such as treating retirement savings accounts in the same manner as pensions are currently—with distributions counting as income but discounting the savings from assets.

**Eliminate Cost-Sharing on Generics for LIS Beneficiaries.** We also recommend eliminating cost sharing on generics for LIS beneficiaries. Even a minimal amount of cost sharing can be a barrier to access. While some plans do offer $0 copay for some generics, applying this policy consistently to all generics would more effectively reduce financial burdens for low-income beneficiaries and increase medication adherence. Encouraging the use of generics should never come at a cost of limiting access to the full range of medications, however. It is important that reducing generic copays to $0 not be accompanied by an increase in LIS cost sharing for branded drugs.

**Notify All LIS Enrollees Who Have Premium Liability about $0 Premium Plans.** Currently, CMS sends the LIS “Chooser’s Notice” only to LIS enrollees with new or increased premium liability relative to the previous year. We are concerned that many LIS enrollees who will have reduced or identical premium liability year-over-year do not receive the notice, regardless of how high their new premium would be. We recommend requiring that CMS send the Chooser’s Notice to all LIS enrollees who have premium liability. This small change would help the LIS program work more efficiently and give LIS enrollees the tools they need to choose the lowest cost plans—thereby decreasing the financial burden for all stakeholders involved.
Index LIS Copayments and Deductibles for LIS Enrollees to the Social Security Cost of Living Adjustment (COLA). Under current law, LIS enrollees with incomes below 100% of the federal poverty level (FPL) have their prescription drug cost sharing increased according to the Consumer Price Index. For LIS enrollees with incomes between 100 and 150% of poverty, their cost sharing is increased according to the percentage increase in average per capita aggregate expenditures for covered Part D drugs. The Social Security COLA is the most accurate reflection of annual income increases for these fixed-income populations and is therefore likely the best index to use for each of these groups. The current indexing methodology for the 100-150% FPL group is particularly problematic since it increases out-of-pocket costs at a higher rate than increases in ability to pay and erodes the value of the LIS benefit over time.

Replace LIS Random Assignment with Intelligent Assignment. Rather than randomly assigning LIS enrollees into new plans, assigning them based on their individual prescription drug needs would reduce their out-of-pocket costs and Medicare program spending. A 2007 House bill, H.R. 3162, included a provision on intelligent assignment that CBO scored at $1.2 billion in savings over 10 years. Factoring in prior medication use, pharmacy preferences, and cost savings into the assignment process can improve access to needed drugs and save money both for beneficiaries and the Medicare program. According to a June 14, 2014 Health Affairs article: “We used an intelligent assignment algorithm and 2008-09 Part D drug use and spending data to match enrollees to available plans according to their medication needs. We found that such a reassignment approach could have saved the federal government over $5 billion in 2009, for government savings of $710 (median: $368) per enrollee with a low-income subsidy.”

Make LI NET Permanent. We are also supportive of proposals to make the Limited Income Newly Eligible Transition (LI NET) Program permanent, including as provided for in S. 1999/H.R. 3029, and as advanced by the House Committee on Ways and Means earlier this week (H.R. 3417). This program acts as an important safety net for beneficiaries who are eligible for LIS but not yet receiving Part D coverage. It allows beneficiaries immediate access to covered Part D drugs at the point-of-sale during the period that begins on the first day of the month a person is determined to be eligible for LIS. This prevents beneficiaries from experiencing a lapse in access to their prescription drugs.

Improve Language Access for Part D Beneficiaries. U.S. Census data estimates that in 2017, over 10 million older adults over age 60 speak a language other than English at home and 6 million speak English less than “very well.” Furthermore, the 2017 Medicare Beneficiary Survey Early Look Data Brief shows that 12% of Medicare beneficiaries living in the community report that English is not their primary language. Other reports from the Office of Minority Health estimate that almost 2 million beneficiaries speak languages other than English or Spanish, including over 200,000 beneficiaries who speak Chinese, over 150,000 who speak Vietnamese, and over 140,000 who speak Tagalog.

Currently, important Part D notices and other Medicare information are sent to most of these limited English proficient (LEP) beneficiaries in English (or Spanish). Moreover, beneficiaries can only submit the

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LIS application in English or Spanish. While the Social Security Administration has made translations of the LIS application available in multiple languages, those translations only serve as guides. LEP beneficiaries must transcribe their information to the English or Spanish version of the application in order to submit it.

We recommend requiring that CMS translate important Part D notices, including notices of disenrollment and coverage denials, into all languages spoken by either 1,000 individuals or 5% of the population in the service area, whichever is lower. CMS’s current 5% threshold, used alone, results in most documents only being translated into Spanish. Similarly, we recommend directing the Social Security Commissioner to allow individuals to directly submit LIS applications in the 18 languages already translated.  

Finally we recommend requiring CMS to translate the Medicare & You Handbook into additional languages beyond Spanish and English. This document provides vital information on all aspects of Medicare, including Part D. All of these translations are necessary to address health disparities, empower beneficiaries with limited English proficiency to access this financial assistance, and are consistent with Title VI of the Civil Rights Act and Section 1557 of the Affordable Care Act.

Thank you for your consideration. We look forward to continued collaboration on efforts to improve prescription drug access and affordability for all people with Medicare.

Sincerely,

Center for Medicare Advocacy  
Justice in Aging  
Medicare Rights Center  
National Committee to Preserve Social Security and Medicare  
National Council on Aging

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11 Recently, Congressman Jimmy Gomez led a letter signed by 62 Members of Congress to CMS Administrator Verma encouraging Medicare to translate key materials such as the Medicare & You Handbook into additional languages. See: https://gomez.house.gov/uploadedfiles/medicare_language_access_final.pdf.