# 114TH CONGRESS 1ST SESSION S.

To establish a demonstration program requiring the utilization of Value-Based Insurance Design in order to demonstrate that reducing the copayments or coinsurance charged to Medicare beneficiaries for selected high-value prescription medications and clinical services can increase their utilization and ultimately improve clinical outcomes, enhance beneficiary satisfaction, and lower health care expenditures.

### IN THE SENATE OF THE UNITED STATES

Mr. Thune introduced the following bill; which was read twice and referred to the Committee on

# A BILL

To establish a demonstration program requiring the utilization of Value-Based Insurance Design in order to demonstrate that reducing the copayments or coinsurance charged to Medicare beneficiaries for selected high-value prescription medications and clinical services can increase their utilization and ultimately improve clinical outcomes, enhance beneficiary satisfaction, and lower health care expenditures.

- 1 Be it enacted by the Senate and House of Representa-
- 2 tives of the United States of America in Congress assembled,

#### 1 SECTION 1. SHORT TITLE.

This Act may be cited as the "Value-Based Insurance

3 Design Seniors Copayment Reduction Act of 2015".

## 4 SEC. 2. FINDINGS.

5 Congress makes the following findings:

- (1) A growing body of evidence demonstrates that increases in patient-level financial barriers (including deductibles, copayments, and coinsurance) for high-value medical services (such as prescription medications, clinician visits, diagnostic tests, and procedures) systematically reduces the use of such services. Savings attributable to cost-related, decreased utilization of specific services may lead to an increase in total medical expenditures due to increased use of other related clinical services, such as hospitalizations and emergency room visits.
  - (2) Empirical research studies demonstrate that reductions in beneficiary out-of-pocket expenses for high-value prescription medications and clinical services can mitigate the adverse health and financial consequences attributable to cost-related decreased utilization of high-value services.
  - (3) Financial barriers to prescription medications and clinical services that are deemed to be high-value should be reduced or eliminated to increase their use.

1	(4) Value-Based Insurance Design is a method-
2	ology that adjusts patient out-of-pocket costs for
3	prescription medications and clinical services accord-
4	ing to the clinical value, not exclusively the cost
5	Value-Based Insurance Design is based on the con-
6	cept of clinical nuance that recognizes—
7	(A) prescription medications and clinical
8	services differ in the clinical benefit provided
9	and
10	(B) the clinical benefit derived from a spe-
11	cific prescription medication or clinical service
12	depends on the clinical situation, the provider
13	and where the care is delivered.
14	(5) The current "one-size-fits-all" copayment or
15	coinsurance design for prescription medications and
16	clinical services provided under the Medicare pro-
17	gram does not recognize the well-established value
18	differences in health outcomes produced by various
19	medical interventions.
20	(6) The establishment by Medicare of copay-
21	ment and coinsurance requirements using Value-
22	Based Insurance Design methodologies will improve
23	patient-centered health outcomes, enhance personal
24	responsibility, and afford a more efficient use of tax-
25	payer dollars.

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1	SEC	3	DEMONSTRATION PROGRAM	

2	(a) In General.—Not later than 1 year after the
3	date of enactment of this Act, the Secretary of Health and
4	Human Services (in this section referred to as the "Sec-
5	retary") shall establish a demonstration program to test
6	Value-Based Insurance Design methodologies in Medicare
7	Advantage plans under part C of title XVIII of the Social
8	Security Act for beneficiaries with chronic clinical condi-
9	tions.
10	(b) Demonstration Program Design.—
11	(1) In general.—The Secretary shall select
12	not less than 2 Medicare Advantage plans to partici-
13	pate in the demonstration program under this sec-
14	tion.
15	(2) REQUIREMENTS.—A Medicare Advantage
16	plan selected to participate in the demonstration
17	program under paragraph (1) shall meet the fol-
18	lowing requirements:
19	(A) The plan offers a coordinated Medicare
20	Part D drug benefit.
21	(B) The plan and the Medicare Advantage
22	organization offering the plan meet such other
23	criteria as the Secretary determines appro-
24	priate.
25	(e) Expansion of Demonstration Program.—
26	The Secretary shall expand the demonstration program by

1	issuing regulations to implement, on a permanent basis,
2	the components of the demonstration program that are
3	beneficial to Medicare beneficiaries and the Medicare pro-
4	gram, unless the report under subsection (e) or (f)(3) con-
5	tains an evaluation that the demonstration program under
6	this section—
7	(1) increases Medicare program expenditures
8	for beneficiaries participating in the demonstration
9	program; or
10	(2) decreases the quality of health care services
11	provided to Medicare beneficiaries participating in
12	the demonstration program.
13	(d) Value-Based Insurance Design Method-
14	OLOGY.—
15	(1) Value-based insurance design.—For
16	purposes of this section, "Value-Based Insurance
17	Design" is a methodology for identifying specific
18	prescription medications and clinical services for
19	which copayments or coinsurance should be reduced
20	or eliminated due to the high-value and effectiveness
21	of such medications and services for specific clinical
22	conditions.
23	(2) Reduction of copayments and coinsur-
24	ANCE.—Under the demonstration program, a Medi-
25	care Advantage organization, using Value-Based In-

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surance Design methodologies, shall identify each prescription medication and clinical service for which the amount of the copayment or coinsurance payable should be reduced or eliminated.

(3) Reduction of Copayments and Coinsurance to encourage use of specific clinical services.—Under the demonstration program, the Medicare Advantage organization, using Value-Based Insurance Design, may lower cost-sharing under the plan for the purpose of encouraging enrollees to use prescription medications and clinical services (such as preventive care, primary care, specialty visits, diagnostic tests, procedures, and durable medical equipment) that such organization has identified as high-value for the management of specified clinical conditions in paragraph (5). Any such variation on copayment or coinsurance by a Medicare Advantage organization must occur on an annual basis and be evidence-based.

(4) Reduction of copayments and coinsurance to encourage use of specific high-performing providers.—Under the demonstration program, the Medicare Advantage organization, using Value-Based Insurance Design, may lower cost-sharing under the plan for the purpose of en-

1	couraging enrollees to use providers that such orga-
2	nization has identified as high-performing based on
3	quality metrics. Any such variation on copayment or
4	coinsurance by a Medicare Advantage organization
5	must occur on an annual basis.
6	(5) Specific clinical conditions.—In iden-
7	tifying clinical conditions for purposes of paragraph
8	(3), the Medicare Advantage organization shall, at a
9	minimum, consider the services utilized across the
10	spectrum of care in the management of the following
11	clinical conditions:
12	(A) Asthma.
13	(B) Atrial fibrillation.
14	(C) Deep venous thrombosis.
15	(D) Cancer.
16	(E) Chronic obstructive pulmonary disease.
17	(F) Chronic renal failure/End stage renal
18	disease.
19	(G) Congestive heart failure.
20	(H) Ischemic heart disease/Myocardial in-
21	farction.
22	(I) Depression.
23	(J) Diabetes mellitus.
24	(K) Hyperlipidemia.
25	(L) Hypertension.

1	(M) Osteoporosis.
2	(N) Stroke.
3	(O) Tobacco abuse disorder.
4	(6) Prohibition of increases of copay-
5	MENTS AND COINSURANCE.—A Medicare Advantage
6	plan selected to participate in the demonstration
7	program under paragraph (1) may not raise cost-
8	sharing on any item or service to discourage its use.
9	(e) Report on Implementation.—
10	(1) In general.—Not later than 1 year after
11	the date of the enactment of this Act, the Secretary
12	shall submit to Congress a report on the implemen-
13	tation by the Secretary of the demonstration pro-
14	gram under this section.
15	(2) Elements.—The report required by para-
16	graph (1) shall include the following:
17	(A) A statement setting forth each medica-
18	tion and clinical service identified pursuant to
19	subsection $(d)(3)$ .
20	(B) For each such medication or clinical
21	service identified pursuant to subsection (d)(3),
22	a statement of the amount of the copayment or
23	coinsurance required to be paid for such service
24	and the amount of the reduction from previous
25	cost-sharing levels.

1	(C) For each such high-performing pro-
2	vider identified pursuant to subsection (d)(4), a
3	statement of the amount of the copayment or
4	coinsurance required to be paid for such clini-
5	cian visit and the amount of the reduction from
6	previous cost-sharing levels.
7	(f) REVIEW AND ASSESSMENT OF UTILIZATION OF
8	Value-based Insurance Design Methodologies.—
9	(1) IN GENERAL.—The Secretary shall enter
10	into a contract or agreement with an independent,
11	nonbiased entity having expertise in Value-Based In-
12	surance Design to review and assess the implemen-
13	tation of the demonstration program under this sec-
14	tion. The review and assessment shall include the
15	following:
16	(A) An assessment of the utilization of
17	Value-Based Insurance Design methodologies
18	referred to in subsection (d).
19	(B) An analysis of whether reducing or
20	eliminating the copayment or coinsurance for
21	each medication and clinical service identified
22	pursuant to subsection (d)(3) resulted in in-
23	creased adherence to medication regimens, in-
24	creased service utilization, improvement in qual-

1	ity metrics, better health outcomes, or enhanced
2	beneficiary experience.
3	(C) An analysis of the cost-savings result-
4	ing from reducing or eliminating the copayment
5	or coinsurance for each medication or clinical
6	service so identified.
7	(D) An analysis of whether reducing or
8	eliminating the copayment or coinsurance for
9	each high-performing provider identified pursu-
10	ant to subsection (d)(4) resulted in improve-
11	ment in quality metrics, better health outcomes,
12	or enhanced beneficiary experience.
13	(E) An analysis of the cost-savings result-
14	ing from reducing or eliminating the copayment
15	or coinsurance for each high-performing pro-
16	vider so identified.
17	(F) Such other matters as the Secretary
18	considers appropriate.
19	(2) Report.—The contract or agreement en-
20	tered into under paragraph (1) shall require the en-
21	tity concerned to submit to the Secretary a report on
22	the review and assessment conducted by the entity
23	under that paragraph in time for the inclusion of the
24	results of such report in the report required by para-
25	graph (3).

1	(3) REPORT TO CONGRESS.—Not later than 3
2	years after the date of the enactment of this Act, the
3	Secretary shall submit to Congress a report on the
4	review and assessment conducted under this sub-
5	section. The report shall include the following:
6	(A) A description of the results of the re-
7	view and assessment.
8	(B) Such recommendations as the Sec-
9	retary considers appropriate for enhancing the
10	utilization of the methodologies referred to in
11	subsection (d)(1) so as to reduce copayments
12	and coinsurance paid by Medicare beneficiaries
13	for high-value prescription medications and
14	clinical services furnished under the Medicare
15	program and to otherwise improve the quality
16	of health care provided under such Medicare
17	program.
18	(g) Waiver.—The Secretary may waive such provi-
19	sions of titles XI and XVIII of the Social Security Act
20	as may be necessary to carry out the demonstration pro-
21	gram under this section.
22	(h) Implementation Funding.—For purposes of
23	carrying out the demonstration program under this sec-
24	tion, the Secretary shall provide for the transfer from the
25	Federal Hospital Insurance Trust Fund under section

- 1 1817 of the Social Security Act (42 U.S.C. 1395i) and
- 2 the Federal Supplementary Insurance Trust Fund under
- 3 section 1841 of the Social Security Act (42 U.S.C. 1395t),
- 4 including the Medicare Prescription Drug Account in such
- 5 Trust Fund, in such proportion as determined appropriate
- 6 by the Secretary, of such sums as may be necessary.