

112TH CONGRESS  
1ST SESSION

# H. R. 891

To amend part D of title XVIII of the Social Security Act to promote medication therapy management under the Medicare part D prescription drug program.

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## IN THE HOUSE OF REPRESENTATIVES

MARCH 3, 2011

Mrs. McMORRIS RODGERS (for herself and Mr. ROSS of Arkansas) introduced the following bill; which was referred to the Committee on Energy and Commerce, and in addition to the Committee on Ways and Means, for a period to be subsequently determined by the Speaker, in each case for consideration of such provisions as fall within the jurisdiction of the committee concerned

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## A BILL

To amend part D of title XVIII of the Social Security Act to promote medication therapy management under the Medicare part D prescription drug program.

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

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Medication Therapy  
5 Management Benefits Act of 2011”.

6 **SEC. 2. FINDINGS.**

7 Congress finds the following:

1           (1) Medications are important to the manage-  
2           ment of chronic diseases that require long-term or  
3           lifelong therapy. Pharmacists are uniquely qualified  
4           as medication experts to work with patients to man-  
5           age their medications and chronic conditions and  
6           play a key role in helping patients take their medica-  
7           tions as prescribed.

8           (2) Nonadherence with medications is a signifi-  
9           cant problem. According to a report by the World  
10          Health Organization, in developed countries, only 50  
11          percent of patients with chronic diseases adhere to  
12          medication therapies. For example, in the United  
13          States only 51 percent of patients taking blood pres-  
14          sure medications are adherent; similarly, only 40 to  
15          70 percent of patients taking antidepressant medica-  
16          tions adhere to prescribed therapies.

17          (3) Failure to take medications as prescribed  
18          costs over \$177 billion annually. The problem of  
19          nonadherence is particularly important for patients  
20          with chronic diseases that require use of medica-  
21          tions; poor adherence leads to unnecessary disease  
22          progression, reduced functional status, lower quality  
23          of life, and premature death.

24          (4) When patients adhere to, or comply with,  
25          their medication therapy, it is possible to reduce

1 higher-cost medical attention, such as emergency de-  
2 partment visits and catastrophic care, and avoid the  
3 preventable human costs that impact patients and  
4 those who care for them.

5 (5) Studies have clearly demonstrated that com-  
6 munity-based medication therapy management  
7 (MTM) services provided by pharmacists improve  
8 health care outcomes and reduce spending. For ex-  
9 ample, the Asheville Project—a diabetes program  
10 designed for city employees in Asheville, North Caro-  
11 lina, and delivered by community pharmacists—re-  
12 sulted over a 5-year period in a decrease in total di-  
13 rect medical costs ranging from \$1,622 to \$3,356  
14 per patient per year, a 50 percent decrease in the  
15 use of sick days, and an increase in productivity ac-  
16 counting for an estimated savings of \$18,000 annu-  
17 ally. Another project involving pharmacist-provided  
18 care to patients with high cholesterol increased com-  
19 pliance with medication to 90 percent from a na-  
20 tional average of 40 percent. In North Carolina, the  
21 ChecKmeds NC program, which offers eligible sen-  
22 iors one-on-one MTM consultations with phar-  
23 macists, saved an estimated \$10,000,000 in  
24 healthcare costs and avoided numerous health prob-  
25 lems in the first year of the program for the more

1 than 15,000 seniors receiving MTM. Similar results  
2 have been achieved in several other demonstrations  
3 using community pharmacists.

4 (6) Therefore, enhancement of the MTM ben-  
5 efit under part D of the Medicare program should  
6 be a key component of the national health care re-  
7 form agenda.

8 **SEC. 3. IMPROVEMENT IN PART D MEDICATION THERAPY**  
9 **MANAGEMENT (MTM) PROGRAMS.**

10 (a) IMPROVEMENTS TO REQUIRED INTERVEN-  
11 TIONS.—Section 1860D–4(c)(2)(C) of the Social Security  
12 Act (42 U.S.C. 1395w–104(c)(2)(C)) is amended—

13 (1) by amending clause (i)(I) to read as follows:

14 “(I) shall include a review of the  
15 individual’s medications, creation of a  
16 personal medication record, and a rec-  
17 ommended medication action plan in  
18 consultation with the individual and  
19 the prescriber; and”;

20 (2) by redesignating clause (ii) as clause (iii)  
21 and inserting after clause (i) the following new  
22 clause:

23 “(ii) Targeted medication reviews fur-  
24 nished person-to-person by a licensed phar-  
25 macist offered no less frequently than once

1 every quarter to assess medication use  
2 since the last annual comprehensive medi-  
3 cation review, to monitor unresolved issues,  
4 to identify problems with new drug thera-  
5 pies or if the individual has experienced a  
6 transition in care.”.

7 (b) INCREASE AVAILABILITY OF MTM SERVICES TO  
8 BENEFICIARIES AND INCREASE COMMUNITY PHARMACY  
9 INVOLVEMENT IN PROVISION OF MTM SERVICES.—

10 (1) INCREASED BENEFICIARY ACCESS TO MTM  
11 SERVICES.—Section 1860D–4(c)(2) of such Act (42  
12 U.S.C. 1395w–104(c)(2)) is further amended—

13 (A) in subparagraph (A)(ii)(I), by inserting  
14 before the semicolon at the end the following:  
15 “or any chronic disease that accounts for high  
16 spending in the Medicare program including di-  
17 abetes, hypertension, heart failure,  
18 dyslipidemia, respiratory disease (such as asth-  
19 ma, chronic obstructive pulmonary disease or  
20 chronic lung disorders), bone disease-arthritis  
21 (such as osteoporosis and osteoarthritis), rheu-  
22 matoid arthritis, and mental health (such as de-  
23 pression, schizophrenia, or bipolar disorder)”;

24 (B) by adding at the end of subparagraph  
25 (A) the following new clause:

1                   “(iii) IDENTIFICATION OF INDIVID-  
2                   UALS WHO MAY BENEFIT FROM MEDICA-  
3                   TION THERAPY MANAGEMENT.—The pre-  
4                   scription drug plan sponsor shall identify a  
5                   process subject to the Secretary’s approval  
6                   that allows licensed pharmacists or other  
7                   qualified providers to identify for medica-  
8                   tion therapy management interventions po-  
9                   tential enrollees who are not described as  
10                  targeted beneficiaries under clause (ii) or  
11                  are not otherwise offered services described  
12                  in subparagraph (C).”;

13                  (C) by redesignating subparagraphs (F)  
14                  and (G) as subparagraphs (I) and (J), respec-  
15                  tively;

16                  (D) by redesignating the subparagraph  
17                  (E), relating to development of program in co-  
18                  operation with licensed pharmacists, as sub-  
19                  paragraph (H);

20                  (E) by redesignating subparagraph (D)  
21                  and the subparagraph (E), relating to auto-  
22                  matic enrollment with ability to opt-out, as sub-  
23                  paragraphs (F) through (G), respectively; and

24                  (F) by inserting after subparagraph (C)  
25                  the following new subparagraph:

1           “(D) MEDICATION REVIEWS FOR DUAL  
2           ELIGIBLES AND ENROLLEES IN TRANSITION OF  
3           CARE.—Without regard to whether an enrollee  
4           is a targeted beneficiary described in subpara-  
5           graph (A)(ii), the medication therapy manage-  
6           ment program under this program shall offer—

7                   “(i) a comprehensive medication re-  
8                   view described in subparagraph (C)(i) at  
9                   the time of initial enrollment under the  
10                  plan for an enrollee who is a full-benefit  
11                  dual eligible individual (as defined in sec-  
12                  tion 1935(c)(6)); and

13                  “(ii) a targeted medication review de-  
14                  scribed in subparagraph (C)(ii) for any en-  
15                  rollee at the time of transition of care  
16                  (such as being discharged from a hospital  
17                  or another institutional setting) where new  
18                  medications have been introduced to the  
19                  individual’s therapy.”.

20           (2) COMMUNITY PHARMACY ACCESS.—Section  
21           1840D–4(c)(2) of such Act, as amended by para-  
22           graph (1), is further amended by inserting after sub-  
23           paragraph (D) the following new subparagraph:

24                   “(E) PHARMACY ACCESS REQUIRE-  
25                   MENTS.—A prescription drug plan sponsor shall

1 offer any willing pharmacy in its network the  
2 ability to provide medication therapy manage-  
3 ment services to assure that enrollees have the  
4 option of obtaining services under the medica-  
5 tion therapy management program from com-  
6 munity-based retail pharmacies.”.

7 (c) REIMBURSEMENT AND INCENTIVES BASED ON  
8 PERFORMANCE.—

9 (1) APPROPRIATE REIMBURSEMENT FOR THE  
10 PROVISION OF MTM SERVICES.—Section 1860D-  
11 4(c)(2)(J) of such Act (42 U.S.C. 1395w-  
12 104(c)(2)(J)), as redesignated by subsection  
13 (b)(1)(C), is amended by striking the first sentence  
14 and inserting the following: “The PDP sponsor shall  
15 reimburse pharmacists and other entities furnishing  
16 medication therapy management services under this  
17 paragraph based on the resources used and the time  
18 required to provide such services.”.

19 (2) EVALUATION OF PERFORMANCE FOR PAY-  
20 MENT INCENTIVES.—Section 1860D-4(c)(2) of such  
21 Act (42 U.S.C. 1395w-104(c)(2)), as amended by  
22 subsection (b), is further amended by adding at the  
23 end the following new subparagraph:

24 “(K) EVALUATION OF PERFORMANCE.—



1                   “(i) DATA COLLECTION AND PRO-  
2                   VIDER MEASURES.—The Secretary shall  
3                   establish measures and standards for data  
4                   collection by prescription drug plan spon-  
5                   sors to evaluate performance of pharmacies  
6                   and other entities in furnishing medication  
7                   therapy management services. Such meas-  
8                   ures and standards shall be developed by  
9                   such date as to allow the application of  
10                  such measures under this subparagraph  
11                  beginning with the first plan year begin-  
12                  ning after the date of the enactment of the  
13                  Medication Therapy Management Benefits  
14                  Act of 2011. Such measures shall be de-  
15                  signed to help assess and improve overall  
16                  quality of care, including a reduction in  
17                  adverse medication reactions, improve-  
18                  ments in adherence and persistence in  
19                  chronic medication use, and a reduction in  
20                  drug spending, where appropriate. Pre-  
21                  scription drug plan sponsors shall use such  
22                  measures to compare outcomes based on  
23                  the type of entity offering such services  
24                  and shall ensure broader participation of  
25                  entities that achieve better outcomes with

1 respect to such services. The measures es-  
2 tablished under this clause shall include  
3 measures developed by the Pharmacy  
4 Quality Alliance (PQA) in the case of  
5 pharmacist providers.

6 “(ii) CONTINUAL DEVELOPMENT AND  
7 INCORPORATION OF MEDICATION THERAPY  
8 MANAGEMENT MEASURES IN BROADER  
9 HEALTH CARE OUTCOMES MEASURES.—

10 The Secretary shall support the continual  
11 development and refinement of perform-  
12 ance measures described in clause (i), in-  
13 cluding the incorporation of medication use  
14 measures as part of broader health care  
15 outcomes measures. The Secretary shall  
16 work with State Medicaid programs to in-  
17 corporate similar performance-based meas-  
18 ures into State drug use review programs  
19 provided pursuant to section 1927(g).

20 “(iii) INCENTIVE PAYMENTS.—

21 “(I) IN GENERAL.—Subject to  
22 subclause (II), for plan years begin-  
23 ning on or after the date that is 1  
24 year after the date the establishment  
25 of measures and standards under

1 clause (i), pharmacies and other enti-  
2 ties that furnish medication therapy  
3 management services under this part  
4 shall be provided (in a manner speci-  
5 fied by the Secretary) with additional  
6 incentive payments based on the per-  
7 formance of such pharmacies and en-  
8 tities in meeting the such measures  
9 and standards. Such payments shall  
10 be made from the Medicare Prescrip-  
11 tion Drug Account except that such  
12 payments may be made from the Fed-  
13 eral Hospital Insurance Trust Fund  
14 or the Federal Supplemental Medical  
15 Insurance Trust Fund if the Sec-  
16 retary determines, based on data  
17 under this part and parts A and B,  
18 that such services have resulted in a  
19 reduction in expenditures under part  
20 A or part B, respectively.

21 “(II) LIMITATION.—The total  
22 amount of additional incentive pay-  
23 ments made under subclause (I) for a  
24 plan year may not exceed the amount  
25 by which the Secretary determines

1                   there are reductions in expenditures  
2                   under this title during such plan year  
3                   resulting from medication therapy  
4                   management services furnished under  
5                   this part.”.

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