115TH CONGRESS 1ST SESSION

H. R. 4724

To provide for a demonstration project to further examine the benefits of providing coverage and payment for items and services necessary to administer intravenous immune globulin (IVIG) in the home, and for other purposes.

IN THE HOUSE OF REPRESENTATIVES

DECEMBER 21, 2017

Mr. Holding (for himself, Mr. Blumenauer, Mr. Lance, Mr. Butterfield, and Mr. Meehan) 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

A BILL

To provide for a demonstration project to further examine the benefits of providing coverage and payment for items and services necessary to administer intravenous immune globulin (IVIG) in the home, and for other purposes.

- 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 "Medicare IVIG Access
- 5 Enhancement Act".

1	SEC. 2. MEDICARE PATIENT IVIG ACCESS DEMONSTRATION
2	PROJECT.
3	(a) Establishment.—The Secretary of Health and
4	Human Services (in this section referred to as the "Sec
5	retary") shall establish and implement a demonstration
6	project under part B of title XVIII of the Social Security
7	Act to evaluate the benefits of providing payment for items
8	and services needed for the in-home administration of in-
9	travenous immune globulin for the treatment of chronic
10	inflammatory demyelinating polyneuropathy or multifoca
11	motor neuropathy.
12	(b) Duration and Scope.—
13	(1) Duration.—Beginning not later than 1
14	year after the date of enactment of this Act, the
15	Secretary shall conduct the demonstration project
16	for a period of 3 years.
17	(2) Scope.—The Secretary shall enroll not
18	greater than 3,000 Medicare beneficiaries who have
19	been diagnosed with chronic inflammatory demyelin-
20	ating polyneuropathy or multifocal motor neurope
21	athy for participation in the demonstration project
22	A Medicare beneficiary may participate in the dem-
23	onstration project on a voluntary basis and may ter-
24	minate participation at any time.
25	(c) COVERAGE —Except as otherwise provided in this

26 section, items and services for which payment may be

- 1 made under the demonstration program shall be treated
- 2 and covered under part B of title XVIII of the Social Se-
- 3 curity Act in the same manner as similar items and serv-
- 4 ices covered under such part.

(d) Payment.—

(1) Intravenous immune globulin furnished under this section, the Secretary shall make payment using the payment methodology under section 1847A of the Social Security Act (42 U.S.C. 1395w–3a).

(2) Other items and services.—

- (A) IN GENERAL.—The Secretary shall establish, subject to subparagraph (B), a per-visit payment amount for items and services (other than intravenous immune globulin) needed for the in-home infusion of intravenous immune globulin for the treatment of chronic inflammatory demyelinating polyneuropathy or multifocal motor neuropathy based on the national per visit low-utilization payment amount under the prospective payment system for home health services established under section 1895 of the Social Security Act (42 U.S.C. 1395fff).
- (B) LIMITATION.—The per-visit payment amount established under subparagraph (A) for

1 items and services described in such subpara-2 graph shall not be less than the payment 3 applied under the demonstration amount 4 project established under section 101 of the Medicare IVIG Access and Strengthening Medi-6 care and Repaying Taxpayers Act of 2012 7 (Public Law 112–242) for comparable items 8 and services needed for the in-home administra-9 tion of intravenous immune globulin for the treatment of primary immune deficiency dis-10 11 eases.

12 (e) WAIVER AUTHORITY.—The Secretary may waive 13 such requirements of title XVIII of the Social Security Act 14 as may be necessary to carry out the demonstration 15 project.

(f) Reports to Congress.—

- (1) Interim evaluation and report.—Not later than 3 years after the date of enactment of this Act, the Secretary shall submit to Congress a report that contains—
- 21 (A) an evaluation of the impact of the 22 demonstration project on access for Medicare 23 beneficiaries with chronic inflammatory demye-24 linating polyneuropathy and Medicare bene-25 ficiaries with multifocal motor neuropathy to

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1	items and services needed for the in-home ad-
2	ministration of intravenous immune globin; and
3	(B) an analysis of the appropriateness of
4	expanding or extending the demonstration
5	project or implementing a new methodology for
6	payment for intravenous immune globulins in
7	all care settings under part B of title XVIII of
8	the Social Security Act (42 U.S.C. 1395k et
9	seq.) and, to the extent such analysis deter-
10	mines such an expansion, extension, or method-
11	ology appropriate, recommendations for such
12	expansion, extension, or methodology, respec-
13	tively.
14	(2) Final evaluation and report.—Not
15	later than one year after the date of completion of
16	the demonstration project, the Secretary shall sub-
17	mit to Congress a report that contains—
18	(A) a final evaluation of the impact de-
19	scribed in paragraph (1)(A); and
20	(B) a final analysis and recommendations
21	described in paragraph (1)(B).
22	(g) Definitions.—In this section:
23	(1) Demonstration project.—The term
24	"demonstration project" means the demonstration
25	project conducted under this Act.

1 (2) MEDICARE BENEFICIARY.—The term
2 "Medicare beneficiary" means an individual who is
3 enrolled for benefits under part B of title XVIII of
4 the Social Security Act.

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