[~118H5397]

Bur ftyplick
(Original Signature of Member)

119TH CONGRESS 1ST SESSION

H.R.

To amend title XVIII of the Social Security Act to provide coverage of external infusion pumps and non-self-administrable home infusion drugs under the Medicare program.

IN THE HOUSE OF REPRESENTATIVES

Mr. FITZPATRICK introduced the following bill; which was referred to the Committee on _____

A BILL

- To amend title XVIII of the Social Security Act to provide coverage of external infusion pumps and non-self-administrable home infusion drugs under the Medicare 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 "Joe Fiandra Access
 - 5 to Home Infusion Act of 2025".

| 1 | SEC. 2. MEDICARE COVERAGE OF EXTERNAL INFUSION |
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| 2 | PUMPS AND NON-SELF-ADMINISTRABLE |
| 3 | HOME INFUSION DRUGS. |
| 4 | (a) IN GENERAL.—Section 1861(n) of the Social Se- |
| 5 | curity Act (42 U.S.C. 1395x(n)) is amended by adding |
| 6 | at the end the following new sentence: "Beginning with |
| 7 | the first calendar quarter beginning on or after the date |
| 8 | that is 1 year after the date of the enactment of this sen- |
| 9 | tence, an external infusion pump and associated home in- |
| 10 | fusion drug (as defined in subsection (iii)(3)(C)) or other |
| 11 | associated supplies that do not meet the appropriate for |
| 12 | use in the home requirement applied to the definition of |
| 13 | durable medical equipment under section 414.202 of title |
| 14 | 42, Code of Federal Regulations (or any successor to such |
| 15 | regulation) shall be treated as meeting such requirement |
| 16 | if each of the following criteria is satisfied: |
| 17 | "(1) The prescribing information approved by |
| 18 | the Food and Drug Administration for the home in- |
| 19 | fusion drug associated with the pump instructs that |
| 20 | the drug should be administered by or under the su- |
| 21 | pervision of a health care professional. |
| 22 | "(2) A qualified home infusion therapy supplier |
| 23 | (as defined in subsection (iii)(3)(D)) administers or |
| 24 | supervises the administration of the drug or biologi- |
| 25 | cal in a safe and effective manner in the patient's |
| 26 | home (as defined in subsection (iii)(3)(B)). |

| 1 | "(3) The prescribing information described in |
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| 2 | paragraph (1) instructs that the drug should be in- |
| 3 | fused at least 12 times per year— |
| 4 | "(A) intravenously or subcutaneously; or |
| 5 | "(B) at infusion rates that the Secretary |
| 6 | determines would require the use of an external |
| 7 | infusion pump.". |
| 8 | (b) Cost Sharing Notification.—The Secretary |
| 9 | of Health and Human Services shall ensure that patients |
| 10 | are notified of the cost sharing for electing home infusion |
| 11 | therapy compared to other applicable settings of care for |
| 12 | the furnishing of infusion drugs under the Medicare pro- |
| 13 | gram. |