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(Original Signature of Member)

118TH CONGRESS  
1ST SESSION

**H. R.** \_\_\_\_\_

To amend the American Taxpayer Relief Act of 2012 to delay implementation of the inclusion of oral-only ESRD-related drugs in the Medicare ESRD prospective payment system.

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IN THE HOUSE OF REPRESENTATIVES

Mr. CARTER of Georgia introduced the following bill; which was referred to the Committee on \_\_\_\_\_

\_\_\_\_\_  
**A BILL**

To amend the American Taxpayer Relief Act of 2012 to delay implementation of the inclusion of oral-only ESRD-related drugs in the Medicare ESRD prospective payment system.

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 “Kidney Patient Access  
5 to Technologically Innovative and Essential Nephrology  
6 Treatments Act of 2023” or the “Kidney PATIENT Act  
7 of 2023”.

1 **SEC. 2. DELAY OF IMPLEMENTATION OF ORAL-ONLY POL-**  
2 **ICY UNDER MEDICARE ESRD PROSPECTIVE**  
3 **PAYMENT SYSTEM.**

4 Section 632(b) of the American Taxpayer Relief Act  
5 of 2012 (42 U.S.C. 1395rr note) is amended—

6 (1) in the heading, by striking “TWO-YEAR”;

7 and

8 (2) in the first sentence of paragraph (1), by  
9 striking “may not implement” and all that follows  
10 through “January 1, 2025.” and inserting “shall not  
11 implement the policy under section 413.174(f)(6) of  
12 title 42, Code of Federal Regulations (relating to  
13 oral-only ESRD-related drugs in the ESRD prospec-  
14 tive payment system) to incorporate the payment for  
15 oral drugs indicated for the reduction, management,  
16 or control of the serum phosphate of an individual,  
17 until the earlier of January 1, 2033, or such time  
18 as an intravenous drug indicated for the reduction,  
19 management, or control of the serum phosphate of  
20 an individual has been approved by the Food and  
21 Drug Administration.”.