

110TH CONGRESS
2D SESSION

S. 3164

To amend title XVIII of the Social Security Act to reduce fraud under the Medicare program.

IN THE SENATE OF THE UNITED STATES

JUNE 19, 2008

Mr. MARTINEZ (for himself and Mr. CORNYN) introduced the following bill;
which was read twice and referred to the Committee on Finance

A BILL

To amend title XVIII of the Social Security Act to reduce fraud 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 “Seniors and Taxpayers
5 Obligation Protection Act of 2008”.

1 **SEC. 2. REQUIRING THE SECRETARY OF HEALTH AND**
2 **HUMAN SERVICES TO CHANGE THE MEDI-**
3 **CARE BENEFICIARY IDENTIFIER USED TO**
4 **IDENTIFY MEDICARE BENEFICIARIES UNDER**
5 **THE MEDICARE PROGRAM.**

6 (a) PROCEDURES.—

7 (1) IN GENERAL.—Not later than 1 year after
8 the date of enactment of this Act, the Secretary of
9 Health and Human Services (in this section referred
10 to as the “Secretary”) shall establish and implement
11 procedures to change the Medicare beneficiary iden-
12 tifier used to identify individuals entitled to benefits
13 under part A of title XVIII of the Social Security
14 Act or enrolled under part B of such title so that
15 such an individual’s social security account number
16 is not used.

17 (2) MAINTAINING EXISTING HICN STRUC-
18 TURE.—In order to minimize the impact of the
19 change under paragraph (1) on systems that com-
20 municate with Medicare beneficiary eligibility sys-
21 tems, the procedures under paragraph (1) shall pro-
22 vide that the new Medicare beneficiary identifier
23 maintain the existing Health Insurance Claim Num-
24 ber structure.

25 (3) PROTECTION AGAINST FRAUD.—The proce-
26 dures under paragraph (1) shall provide for a proc-

ess for changing the Medicare beneficiary identifier for an individual to a different identifier in the case of the discovery of fraud, including identity theft.

(4) PHASE-IN AUTHORITY.—

(A) IN GENERAL.—Subject to subparagraphs (B) and (C), the Secretary may phase in the change under paragraph (1) in such manner as the Secretary determines appropriate.

(B) LIMIT.—The phase-in period under subparagraph (A) shall not exceed 10 years.

(C) NEWLY ENTITLED AND ENROLLED INDIVIDUALS.—The Secretary shall ensure that the change under paragraph (1) is implemented not later than January 1, 2010 with respect to any individual who first becomes entitled to benefits under part A of title XVIII of the Social Security Act or enrolled under part B of such title on or after such date.

(b) EDUCATION AND OUTREACH.—The Secretary shall establish a program of education and outreach for individuals entitled to benefits under part A of title XVIII of the Social Security Act or enrolled under part B of such title, providers of services (as defined in subsection (u) of section 1861 of such Act (42 U.S.C. 1395x)), and sup-

pliers (as defined in subsection (d) of such section) on the change under paragraph (1).

(c) AUTHORIZATION OF APPROPRIATIONS.—There are authorized to be appropriated such sums as may be necessary to carry out this section.

SEC. 3. MONTHLY VERIFICATION OF ACCURACY OF CHARGES FOR PHYSICIANS' SERVICES.

(a) IN GENERAL.—Section 1893 of the Social Security Act (42 U.S.C. 1395ddd) is amended—

(1) in subsection (b), by adding at the end the following new paragraph:

“(7) The monthly verification of the accuracy of charges for physicians’ services under the system under subsection (i).”;

(2) in subsection (c), by adding at the end of the flush matter following paragraph (4), the following new sentence: “In the case of the activity described in subsection (b)(7), an entity shall only be eligible to enter into a contract under the Program to carry out the activity if the entity is a medicare administrative contractor with a contract under section 1874A.”; and

(3) by adding at the end the following new subsection:

1 “(i) MONTHLY VERIFICATION OF ACCURACY OF
2 CHARGES FOR PHYSICIANS’ SERVICES.—

3 “(1) SYSTEM.—

4 “(A) IN GENERAL.—Not later than 1 year
5 after the date of the enactment of this sub-
6 section, the Secretary shall establish and imple-
7 ment a system to verify (electronically or other-
8 wise, taking into consideration the administra-
9 tive burden of such verification on physicians
10 and group practices) on a monthly basis that
11 the claims for reimbursement under part B for
12 physicians’ services furnished in high risk areas
13 are—

14 “(i) for physicians’ services actually
15 furnished by the physician (or the physi-
16 cian’s group practice); and

17 “(ii) otherwise accurate.

18 “(B) NO DETERMINATION OF MEDICAL
19 NECESSITY.—In no case shall any verification
20 conducted under the system established under
21 subparagraph (A) include a determination of
22 the medical necessity of the physicians’ service.

23 “(2) VERIFICATION.—Under the system, the
24 Secretary, at the end of each month, shall provide
25 the physician (or the group practice) with a detailed

1 list of such claims for reimbursement that were sub-
2 mitted during the month in order for the physician
3 (or the group practice) to review and verify the list.
4 In providing the detailed list, the Secretary shall use
5 the provider number of the physician (or the group
6 practice).

7 “(3) AUDITS.—The Secretary shall conduct au-
8 dits of the review and verification by physicians and
9 group practices of the detailed list provided under
10 paragraph (2). Such audits shall assess whether the
11 physician or group practice conducted such review
12 and verification in a fraudulent manner. In the case
13 where the Secretary determines such review and
14 verification was conducted in a fraudulent manner,
15 the Secretary shall recoup any payments resulting
16 from the fraudulent review and verification and im-
17 pose a civil money penalty in an amount determined
18 appropriate by the Secretary on the physician or
19 group practice who conducted the fraudulent review
20 and verification. The provisions of section 1128A
21 (other than subsections (a) and (b)) shall apply to
22 a civil money penalty under the previous sentence in
23 the same manner as such provisions apply to a pen-
24 alty or proceeding under section 1128A(a).

1 “(4) HIGH RISK AREAS DEFINED.—In this sub-
 2 section, the term ‘high risk area’ means a county
 3 designated as a high risk area under subsection
 4 (j)(1).

5 “(5) ACTIONS THROUGH MEDICARE ADMINIS-
 6 TRATIVE CONTRACTORS.—In carrying out this sub-
 7 section, the Secretary shall act through medicare ad-
 8 ministrative contractors with a contract under sec-
 9 tion 1874A.

10 “(6) REPORT BY THE SECRETARY.—Not later
 11 than 1 year after implementation of the system es-
 12 tablished under paragraph (1), the Secretary shall
 13 submit a report to Congress on the progress of such
 14 implementation. Such report shall include rec-
 15 ommendations—

16 “(A) on how to improve such implementa-
 17 tion, including whether the system should be ex-
 18 panded to include verification of claims for re-
 19 imbursement under part B for physicians’ serv-
 20 ices furnished in additional areas; and

21 “(B) for such legislation and administra-
 22 tive action as the Secretary determines appro-
 23 priate.”.

24 (b) AUTHORIZATION OF APPROPRIATIONS.—To carry
 25 out the amendments made by this section, there are au-

1 thorized to be appropriated such sums as may be nec-
 2 essary for each of fiscal years 2009 through 2013.

3 **SEC. 4. DETECTION OF MEDICARE FRAUD IN HIGH RISK**
 4 **AREAS.**

5 (a) IN GENERAL.—Section 1893 of the Social Secu-
 6 rity Act (42 U.S.C. 1395ddd), as amended by section 3,
 7 is amended—

8 (1) in subsection (b), by adding at the end the
 9 following new paragraph:

10 “(8) Implementation of prepayment fraud de-
 11 tection methods under subsection (j).”;

12 (2) in subsection (c), in the second sentence of
 13 the flush matter following paragraph (4), by striking
 14 “activity described in subsection (b)(7)” and insert-
 15 ing “activities described in paragraphs (7) and (8)
 16 of subsection (b)”;

17 (3) by adding at the end the following new sub-
 18 section:

19 “(j) DETECTION OF MEDICARE FRAUD IN HIGH
 20 RISK AREAS.—

21 “(1) ESTABLISHMENT OF SYSTEM TO IDENTIFY
 22 COUNTIES MOST VULNERABLE TO FRAUD.—Not
 23 later than 6 months after the date of the enactment
 24 of this subsection, the Secretary shall establish a
 25 system to identify the 50 counties most vulnerable to

1 fraud with respect to items and services furnished by
2 providers of services (other than hospitals and crit-
3 ical access hospitals) and suppliers based on the de-
4 gree of county-specific reimbursement and analysis
5 of payment trends under this title. The Secretary
6 shall designate the counties identified under the pre-
7 ceding sentence as ‘high risk areas’.

8 “(2) PREPAYMENT FRAUD DETECTION.—The
9 Secretary shall establish procedures for the imple-
10 mentation of prepayment fraud detection methods
11 under this title with respect to items and services
12 furnished by such providers of services and suppliers
13 in high risk areas designated under paragraph (1),
14 including the following:

15 “(A) Pre-enrollment site visits for such
16 providers of services and suppliers which have
17 the highest probability of committing fraud
18 under this title.

19 “(B) Data analysis to establish prepay-
20 ment claim edits designed to target the claims
21 for reimbursement under this title for such
22 items and services that are most likely to be
23 fraudulent.

24 “(C) Prepayment benefit integrity reviews
25 for claims for reimbursement under this title

1 for such items and services that are suspended
2 as a result of such edits.

3 “(3) ACTIONS THROUGH MEDICARE ADMINIS-
4 TRATIVE CONTRACTORS.—In carrying out this sub-
5 section, the Secretary shall act through medicare ad-
6 ministrative contractors with a contract under sec-
7 tion 1874A.

8 “(4) REPORT TO CONGRESS.—The Secretary
9 shall, upon request, appear and testify before Con-
10 gress regarding the status of the implementation of
11 prepayment fraud detection methods under this sub-
12 section.”.

13 (b) AUTHORIZATION OF APPROPRIATIONS.—To carry
14 out the amendments made by this section, there are au-
15 thorized to be appropriated such sums as may be nec-
16 essary, not to exceed \$50,000,000, for each of fiscal years
17 2009 through 2013.

18 **SEC. 5. STUDY ON THE USE OF TECHNOLOGY FOR REAL-**
19 **TIME DATA REVIEW.**

20 (a) STUDY ON THE USE OF TECHNOLOGY FOR REAL-
21 TIME DATA REVIEW.—The Secretary of Health and
22 Human Services shall conduct a study on the use of tech-
23 nology (similar to that used with respect to the analysis
24 of credit card charging patterns) to provide real-time data
25 analysis of claims for reimbursement under the Medicare

1 program under title XVIII of the Social Security Act to
2 identify and investigate unusual billing or order practices
3 under the Medicare program that could indicate fraud or
4 abuse. Such study shall include an analysis of the fol-
5 lowing:

6 (1) Whether such technology could be used to
7 identify unusual billing or order practices under the
8 Medicare program by an individual provider of serv-
9 ices or for a certain HCPCS code in a particular
10 area of the country without alerting potentially
11 fraudulent providers of services and allowing them to
12 escape or go unnoticed.

13 (2) How such technology can be implemented
14 under the Medicare program to provide for the effec-
15 tive review of claim logs in an accurate and timely
16 manner.

17 (b) REPORT.—Not later than 1 year after the date
18 of enactment of this Act, the Secretary shall submit a re-
19 port to Congress on the study conducted under subsection
20 (a), together with recommendations for such legislation
21 and administrative action as the Secretary determines ap-
22 propriate.

1 **SEC. 6. EDITS ON 855S MEDICARE ENROLLMENT APPLICA-**
 2 **TION.**

3 Section 1834(a) of the Social Security Act (42 U.S.C.
 4 1395m(a)) is amended by adding at the end the following
 5 new paragraph:

6 “(22) CONFIRMATION WITH NATIONAL SUP-
 7 PLIER CLEARINGHOUSE PRIOR TO REIMBURSE-
 8 MENT.—

9 “(A) IN GENERAL.—Not later than 1 year
 10 after the date of enactment of this paragraph,
 11 the Secretary shall establish procedures to re-
 12 quire carriers, prior to paying a claim for reim-
 13 bursement for durable medical equipment, pros-
 14 thetics, orthotics, and supplies under this title,
 15 to confirm with the National Supplier Clearing-
 16 house—

17 “(i) that the Medicare identification
 18 number of the supplier is active; and

19 “(ii) that the item or service for which
 20 the claim for reimbursement is submitted
 21 was properly identified on the CMS–855S
 22 Medicare enrollment application.

23 “(B) ONLINE DATABASE FOR IMPLEMEN-
 24 TATION.—Not later than 18 months after the
 25 date of enactment of this paragraph, the Sec-
 26 retary shall establish an online database similar

1 to that used for the National Provider Identifier
 2 to enable providers of services, accreditors, car-
 3 riers, and the National Supplier Clearinghouse
 4 to view information on specialties and the types
 5 of items and services each supplier has indi-
 6 cated on the CMS–855S Medicare enrollment
 7 application submitted by the supplier.

8 “(C) NOTIFICATION OF CLAIM DENIAL
 9 AND RESUBMISSION.—In the case where a claim
 10 for reimbursement for durable medical equip-
 11 ment, prosthetics, orthotics, and supplies under
 12 this title is denied because the item or service
 13 furnished does not correctly match up with the
 14 information on file with the National Supplier
 15 Clearinghouse—

16 “(i) the National Supplier Clearing-
 17 house shall—

18 “(I) provide the supplier written
 19 notification of the reason for such de-
 20 nial; and

21 “(II) allow the supplier 60 days
 22 to provide the National Supplier
 23 Clearinghouse with appropriate certifi-
 24 cation, licensing, or accreditation; and

1 “(ii) the Secretary shall waive applica-
 2 ble requirements relating to the time frame
 3 for the submission of claims for payment
 4 under this title in order to permit the re-
 5 submission of such claim if payment of
 6 such claim would otherwise be allowed
 7 under this title.”.

8 **SEC. 7. SERIAL NUMBER TRACKING SYSTEM FOR DURABLE**
 9 **MEDICAL EQUIPMENT.**

10 (a) IN GENERAL.—Section 1834(a) of the Social Se-
 11 curity Act (42 U.S.C. 1395m(a)), as amended by section
 12 6(a), is amended by adding at the end the following new
 13 paragraph:

14 “(23) SERIAL NUMBER TRACKING SYSTEM FOR
 15 DURABLE MEDICAL EQUIPMENT.—

16 “(A) ESTABLISHMENT.—In the case of
 17 any item of durable medical equipment which
 18 has not been issued a unique identifier under
 19 the unique device identification system estab-
 20 lished under section 519(f) of the Federal
 21 Food, Drug, and Cosmetic Act, the Secretary
 22 shall promulgate regulations establishing a sys-
 23 tem for such durable medical equipment requir-
 24 ing the label of such equipment to bear a
 25 unique identifier, unless the Secretary requires

1 an alternative placement or provides an excep-
2 tion for a particular item or type of durable
3 medical equipment under such section 519(f).

4 “(B) PROVISION OF UNIQUE IDENTIFIER
5 TO THE SECRETARY.—A manufacturer of an
6 item of durable medical equipment shall submit
7 to the Secretary the unique identifier issued
8 under subparagraph (A) or such section 519(f)
9 with respect to such item (in accordance with
10 procedures established by the Secretary). The
11 Secretary shall provide for the storage of such
12 unique identifier in accordance with subpara-
13 graph (D)(i).

14 “(C) REQUIREMENTS FOR MANUFACTUR-
15 ERS AND WHOLESALERS.—A manufacturer of
16 an item of durable medical equipment, or, in
17 the case where a wholesaler provides an item of
18 durable medical equipment to a supplier, the
19 wholesaler, shall—

20 “(i) upon issuing an item to a sup-
21 plier, develop a product description for the
22 item which includes—

23 “(I) the unique identifier of the
24 item;

1 “(II) the specific Healthcare
 2 Common Procedure Coding System
 3 (HCPCS) code for the item;

4 “(III) the name of the supplier
 5 the item was shipped to; and

6 “(IV) the supplier’s Medicare
 7 identification number; and

8 “(ii) submit the product description
 9 developed under clause (i) to the Secretary
 10 for storage in the unique identifier data-
 11 base in accordance with subparagraph
 12 (E)(i).

13 “(D) REQUIREMENTS FOR SUPPLIERS.—A
 14 supplier of an item of durable medical equip-
 15 ment shall—

16 “(i) upon issuing the item to a bene-
 17 ficiary, note the unique identifier of such
 18 item on—

19 “(I) the claim form submitted for
 20 such item; and

21 “(II) when appropriate or other-
 22 wise required, the detailed product de-
 23 scription of the item;

24 “(ii) in the case where the item is
 25 issued to a beneficiary on a rental basis,

1 designate the unique identifier with an ‘R’
2 after the number to indicate that the item
3 was rented, and not purchased, by the ben-
4 eficiary; and

5 “(iii) upon return of the item to the
6 supplier, notify the Secretary—

7 “(I) before reissuing that item
8 and resubmitting that number on
9 such a claim form; or

10 “(II) upon resubmitting that
11 number on such a claim form.

12 “(E) REQUIREMENTS FOR THE SEC-
13 RETARY.—

14 “(i) MAINTENANCE OF DATABASE OF
15 SERIAL NUMBERS.—The Secretary shall
16 establish and maintain a database con-
17 taining the unique identifiers submitted by
18 manufacturers of items of durable medical
19 equipment under subparagraph (B).

20 “(ii) PAYMENT.—

21 “(I) LIMITATION.—Subject to
22 subclause (II), payment may only be
23 made for an item of durable medical
24 equipment under this part if the
25 unique identifier on the claim form

1 submitted for such item matches the
2 unique identifier submitted by the
3 manufacturer of such item under sub-
4 paragraph (B).

5 “(II) EXCEPTION TO LIMITATION
6 AFTER VERIFICATION OF RECEIPT.—
7 In the case where the unique identi-
8 fier is not on the claim form sub-
9 mitted for such item or does not
10 match the unique identifier submitted
11 by the manufacturer of such item
12 under subparagraph (B), no payment
13 shall be made under this part for the
14 item of durable medical equipment
15 until the Secretary has verified that
16 the beneficiary has received such item
17 in accordance with subclause (IV).

18 “(III) DUPLICATIVE UNIQUE
19 IDENTIFIERS.—In the case where a
20 unique identifier is submitted on more
21 than 1 claim form submitted for such
22 an item and there is no indication
23 from the supplier that the item of du-
24 rable medical equipment has been re-
25 turned by 1 beneficiary and is now

1 being used by another beneficiary, no
2 payment shall be made under this
3 part for such item of durable medical
4 equipment unless the Secretary has
5 verified that the beneficiary has re-
6 ceived such item in accordance with
7 subclause (IV).

8 “(IV) VERIFICATION.—The Sec-
9 retary shall conduct any verification
10 required under subclause (II) or (III)
11 within 30 days after receipt by the
12 Secretary of the relevant claim form.
13 In the case where such verification is
14 not completed within such time pe-
15 riod, the Secretary shall pay such
16 claim, complete the verification, and,
17 in the case where the Secretary has
18 entered into a contract with an entity
19 for the conduct of such verification,
20 recover any payments that would not
21 have been made if the verification had
22 been completed within such time pe-
23 riod from such entity.

24 “(iii) QUALITY CONTROL AUDITS.—
25 The Secretary shall conduct quality control

audits to identify unusual billing patterns with respect to items of durable medical equipment for which payment is made under this part and may conduct unannounced site visits or commission other agencies to conduct such site visits as part of such quality control audits.

“(iv) NO USE AS A PRECERTIFICATION MECHANISM.—In no case shall a unique identifier issued under subparagraph (A) or section 519(f) of the Federal Food, Drug, and Cosmetic Act be used as a precertification mechanism for the supply of an item of durable medical equipment or the payment of a claim for such an item under this part.”.

(b) EFFECTIVE DATE.—The amendment made by subsection (a) shall take effect 3 years after the date of enactment of this Act.

SEC. 8. SENSE OF THE SENATE REGARDING SURETY BOND REQUIREMENTS FOR SUPPLIERS OF DURABLE MEDICAL EQUIPMENT.

(a) FINDINGS.—The Senate finds the following:

(1) Documented fraud in the Medicare Durable Medical Equipment, Prosthetics, Orthotics, and Sup-

1 plies Competitive Bidding Program under section
2 1847 of the Social Security Act (42 U.S.C. 1395w–
3 3) has potentially cost taxpayers in the United
4 States billions of dollars.

5 (2) Congress, having previously recognized
6 fraudulent practices with respect to durable medical
7 equipment under the Medicare program under title
8 XVIII of the Social Security Act, directed the Sec-
9 retary of Health and Human Services to take action
10 against such fraudulent practices through the imple-
11 mentation of a surety bond requirement under sec-
12 tion 1834(a)(16) of the Social Security Act (42
13 U.S.C. 1395m(a)(16)), as added by section 4312 of
14 the Balanced Budget Act of 1997 (Public Law 105–
15 33).

16 (3) Such surety bond requirement is necessary
17 to—

18 (A) limit the risk to the Medicare program
19 of fraudulent suppliers of durable medical
20 equipment;

21 (B) enhance the enrollment process under
22 the Medicare program to ensure that only legiti-
23 mate suppliers of durable medical equipment
24 are enrolled or are allowed to remain enrolled in

1 any programs established or implemented under
2 the Medicare program;

3 (C) ensure that the Medicare program re-
4 coups erroneous payments that result from
5 fraudulent or abusive billing practices by allow-
6 ing the Centers for Medicare & Medicaid Serv-
7 ices, or entities under a contract with the Cen-
8 ters for Medicare & Medicaid Services, to seek
9 payments from a surety up to the penal sum;
10 and

11 (D) help ensure that beneficiaries under
12 the Medicare program receive items and serv-
13 ices that are considered reasonable and nec-
14 essary from legitimate suppliers of durable
15 medical equipment.

16 (4) To date, more than a decade after the en-
17 actment of the Balanced Budget Act of 1997 (Public
18 Law 105–33), such section 1834(a)(16) has yet to
19 be implemented by the Secretary of Health and
20 Human Services, potentially costing taxpayers and
21 Medicare beneficiaries billions of additional dollars
22 and negatively impacting responsible suppliers of du-
23 rable medical equipment under the Medicare pro-
24 gram.

1 (b) SENSE OF THE SENATE.—It is the Sense of the
2 Senate that the Secretary of Health and Human Services
3 must put in place the surety bond requirement under sec-
4 tion 1834(a)(16) of the Social Security Act (42 U.S.C.
5 1395m(a)(16)) within 6 months of the date of enactment
6 of this Act in order to maintain integrity under the Medi-
7 care program.

○