To respond to a Medicare funding warning.

IN THE SENATE OF THE UNITED STATES

FEBRUARY 25, 2008

Mr. BAUCUS (for himself and Mr. GREGG) (by request) introduced the following bill; which was referred to the Committee on Finance

A BILL

To respond to a Medicare funding warning.

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

SECTION 1. SHORT TITLE; REFERENCES; PURPOSE OF LEG-
ISLATION.

(a) Short Title.—This Act may be cited as the “Medicare Funding Warning Response Act of 2008”.

(b) References.—In this Act:

(1) Except where otherwise specifically pro-
vided, references in this Act shall be considered to
be made to the Social Security Act, or to a section
or other provision thereof.
(2) The term “Secretary” shall be deemed a reference to the Secretary of Health and Human Services.

(3) The terms “Medicare” and “Medicare program” mean the program under title XVIII of the Social Security Act (42 U.S.C. 1395 et seq.).

(4) The Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (Public Law 108–173) shall be referred to as the “MMA”.

(5) The term “excess general revenue medicare funding” has the meaning given such term by section 801(c) of the MMA.

(6) The term “Trustees Report” means the annual report submitted under subsection (b)(2) of sections 1817 and 1841 of the Social Security Act (42 U.S.C. 1395i(b)(2) and 1395t(b)(2), respectively).

(c) PURPOSE.—It is the purpose of this Act to respond to the medicare funding warning currently in effect under section 801(a)(2) of the MMA.
TITLE I—INTRODUCING PRINCIPLES OF VALUE-BASED HEALTH CARE INTO THE MEDICARE PROGRAM

SEC. 101. INTRODUCING PRINCIPLES OF VALUE-BASED HEALTH CARE INTO THE MEDICARE PROGRAM.

(a) ELECTRONIC HEALTH RECORDS.—The Secretary shall develop and implement a system for encouraging nationwide adoption and use of interoperable electronic health records and to make available personal health records for Medicare beneficiaries.

(b) PRICING TRANSPARENCY.—The Secretary shall make publicly available information on prices and payments under the Medicare program for treatments (including episodes of care), items, and services to assist Medicare beneficiaries in making choices among providers, plans, and treatment options.

(c) QUALITY TRANSPARENCY.—The Secretary shall make publicly available information on the quality of care provided to Medicare beneficiaries to assist them in making choices among providers, plans, and treatments. To ensure the continued development and evolution of quality measures, the Secretary shall develop and implement a plan for ensuring that, by the year 2013, quality measures
are available and reported with respect to at least 50 percent of the care provided under the Medicare program (determined according to the amount of payment made under such program for items and services with respect to which such measures are available). The Secretary shall report to the Committees on Ways and Means and Energy and Commerce in the House of Representatives and the Committee on Finance in the Senate annually on the progress of the goal specified in the preceding sentence.

(d) INCENTIVES FOR VALUE.—

(1) INCENTIVES FOR PROVIDERS AND SUPPLIERS.—

(A) IN GENERAL.—The Secretary shall design and implement a system for use in the Medicare program under which a portion of the payments that would otherwise be made under such program to some or all classes of individuals and entities furnishing items or services to beneficiaries of such program would be based on the quality and efficiency of their performance.

(B) IMPLEMENTATION.—The Secretary shall first implement such system in settings where measures are well-accepted and already collected, including hospitals, physicians’ of-
fices, home health agencies, skilled nursing facilities, and renal dialysis facilities. The initial focus of such efforts shall be on quality, but the Secretary shall add measures of efficiency as they are identified. The system shall also include incentives for reducing unwarranted geographic variations in quality and efficiency.

(C) SECRETARY’S AUTHORITY.—The Secretary may implement the system described in this paragraph without regard to any provision of title XVIII of the Social Security Act that would, in the absence of subparagraphs (A) and (B), apply with respect to payment to an individual or entity furnishing items or services for which payment may be made under the Medicare program.

(2) BENEFICIARY INCENTIVES.—

(A) IN GENERAL.—The Secretary shall implement incentives for Medicare beneficiaries to use more efficient providers and preventive services known to reduce costs.

(B) ACCESS TO HEALTH SAVINGS ACCOUNTS.—The Secretary shall assure a transition into the Medicare program for individuals who are not yet enrolled in such program who
own health savings accounts, and shall provide
for the availability of high deductible health
plan options in the Medicare program.

(e) Broadly Transforming the Private Health
Care Marketplace.—The Secretary shall use and re-
lease Medicare data for quality improvement, performance
measurement, public reporting, and treatment-related pur-
poses. In implementing the preceding sentence, the Sec-
retary shall apply risk adjustment techniques where ap-
propriate and shall determine the circumstances under
which it is appropriate to release such data.

(f) Protecting Individually Identifiable
Health Information.—In implementing this title, the
Secretary shall ensure that individually identifiable bene-
ficiary health information is protected (in accordance with
the regulations adopted under section 264(e) of the Health
Insurance Portability and Accountability Act of 1996 and
such other laws and regulations as may apply).

(g) Regulations.—The Secretary may implement a
system described in this section by regulation, but only
if such regulation is issued after public notice and an op-
portunity for public comment.

(h) Definitions.—As used in this section:

(1) The term “efficiency” means the delivery of
health care in a manner that reduces the costs of
providing care for Medicare beneficiaries while maintaining or improving the quality of such care.

(2) The term “information on quality of care” means such measures of—

(A) the use of clinical processes and structures known to improve care;

(B) health outcomes; and

(C) patient perceptions of their care,
as the Secretary may select with preference given to those measures that have been recognized through a consensus-based process.

(i) Savings Requirement.—

(1) In General.—The Secretary may implement the provisions of subsections (a) through (e) of section 101 and section 102 for a year only to the extent that the Secretary determines (and the Chief Actuary of the Centers for Medicare & Medicaid Services certifies) that—

(A) the total amount of payment made under title XVIII of the Social Security Act over the five and ten year periods that begin with January 1 of such year as a result of the implementation of such subsections (a) through (e) and section 102 is less than the amount
that would have been made over such periods if such implementation had not occurred; and

(B) the total amount of payment made under each of titles XIX and XXI of such Act over such periods as a result of such implementation is no greater than the amount that would have been made under each such title over such periods if such implementation had not occurred.

(2) AVAILABILITY OF APPROPRIATIONS.—The Secretary shall carry out the provisions of this section subject to the availability of appropriations and to the extent permitted consistent with paragraph (1).

SEC. 102. RELEASE OF PHYSICIAN PERFORMANCE MEASUREMENTS.

Section 1848(k) (42 U.S.C. 1395w–4(k)) is amended by adding at the end the following new paragraph:

“(9) RELEASE OF QUALITY MEASUREMENTS.—

“(A) IN GENERAL.—Notwithstanding section 552a of title 5, United States Code, the Secretary may—

“(i) release to the public physician-specific measurements of the quality or efficiency of physician performance against a
standard (reflecting measurements that have been recognized through a consensus-based process) that has been endorsed by the Secretary; and

“(ii) release, to an entity that will generate or calculate such measurements, data that the entity may use to perform such task.

“(B) ENDORSEMENT OF STANDARDS.—The Secretary may make an endorsement under subparagraph (A) by publication of a notice in the Federal Register.”.

TITLE II—REDUCING THE EXCESSIVE BURDEN THE LIABILITY SYSTEM PLACES ON THE HEALTH CARE DELIVERY SYSTEM

SEC. 201. SHORT TITLE.
This title may be cited as the “Help Efficient, Accessible, Low-cost, Timely Healthcare (HEALTH) Act of 2008”.

SEC. 202. FINDINGS AND PURPOSE.

(a) FINDINGS.—

(1) EFFECT ON HEALTH CARE ACCESS AND COSTS.—Congress finds that our current civil justice
system is adversely affecting patient access to health
care services, better patient care, and cost-efficient
health care, in that the health care liability system
is a costly and ineffective mechanism for resolving
claims of health care liability and compensating in-
jured patients, and is a deterrent to the sharing of
information among health care professionals which
impedes efforts to improve patient safety and quality
of care.

(2) EFFECT ON INTERSTATE COMMERCE.—
Congress finds that the health care and insurance
industries are industries affecting interstate com-
merce and the health care liability litigation systems
existing throughout the United States are activities
that affect interstate commerce by contributing to
the high costs of health care and premiums for
health care liability insurance purchased by health
care system providers.

(3) EFFECT ON FEDERAL SPENDING.—Con-
gress finds that the health care liability litigation
systems existing throughout the United States have
a significant effect on the amount, distribution, and
use of Federal funds because of—
(A) the large number of individuals who receive health care benefits under programs operated or financed by the Federal Government;

(B) the large number of individuals who benefit because of the exclusion from Federal taxes of the amounts spent to provide them with health insurance benefits; and

(C) the large number of health care providers who provide items or services for which the Federal Government makes payments.

(b) PURPOSE.—It is the purpose of this title to implement reasonable, comprehensive, and effective health care liability reforms designed to—

(1) improve the availability of health care services in cases in which health care liability actions have been shown to be a factor in the decreased availability of services;

(2) reduce the incidence of “defensive medicine” and lower the cost of health care liability insurance, all of which contribute to the escalation of health care costs;

(3) ensure that persons with meritorious health care injury claims receive fair and adequate compensation, including reasonable noneconomic damages;
(4) improve the fairness and cost-effectiveness of our current health care liability system to resolve disputes over, and provide compensation for, health care liability by reducing uncertainty in the amount of compensation provided to injured individuals; and

(5) provide an increased sharing of information in the health care system which will reduce unintended injury and improve patient care.

SEC. 203. ENCOURAGING SPEEDY RESOLUTION OF CLAIMS.

The time for the commencement of a health care lawsuit shall be 3 years after the date of manifestation of injury or 1 year after the claimant discovers, or through the use of reasonable diligence should have discovered, the injury, whichever occurs first. In no event shall the time for commencement of a health care lawsuit exceed 3 years after the date of manifestation of injury unless tolled for any of the following—

(1) upon proof of fraud;

(2) intentional concealment; or

(3) the presence of a foreign body, which has no therapeutic or diagnostic purpose or effect, in the person of the injured person.

Actions by a minor shall be commenced within 3 years from the date of the alleged manifestation of injury except that actions by a minor under the full age of 6 years shall
be commenced within 3 years of manifestation of injury or prior to the minor’s 8th birthday, whichever provides a longer period. Such time limitation shall be tolled for minors for any period during which a parent or guardian and a health care provider or health care organization have committed fraud or collusion in the failure to bring an action on behalf of the injured minor.

SEC. 204. COMPENSATING PATIENT INJURY.

(a) UNLIMITED AMOUNT OF DAMAGES FOR ACTUAL ECONOMIC LOSSES IN HEALTH CARE LAWSUITS.—In any health care lawsuit, nothing in this title shall limit a claimant’s recovery of the full amount of the available economic damages, notwithstanding the limitation in subsection (b).

(b) ADDITIONAL NONECONOMIC DAMAGES.—In any health care lawsuit, the amount of noneconomic damages, if available, may be as much as $250,000, regardless of the number of parties against whom the action is brought or the number of separate claims or actions brought with respect to the same injury.

(c) NO DISCOUNT OF AWARD FOR NONECONOMIC DAMAGES.—For purposes of applying the limitation in subsection (b), future noneconomic damages shall not be discounted to present value. The jury shall not be informed about the maximum award for noneconomic damages. An award for noneconomic damages in excess of
$250,000 shall be reduced either before the entry of judgment, or by amendment of the judgment after entry of judgment, and such reduction shall be made before accounting for any other reduction in damages required by law. If separate awards are rendered for past and future noneconomic damages and the combined awards exceed $250,000, the future noneconomic damages shall be reduced first.

(d) Fair Share Rule.—In any health care lawsuit, each party shall be liable for that party’s several share of any damages only and not for the share of any other person. Each party shall be liable only for the amount of damages allocated to such party in direct proportion to such party’s percentage of responsibility. Whenever a judgment of liability is rendered as to any party, a separate judgment shall be rendered against each such party for the amount allocated to such party. For purposes of this section, the trier of fact shall determine the proportion of responsibility of each party for the claimant’s harm.

SEC. 205. MAXIMIZING PATIENT RECOVERY.

(a) Court Supervision of Share of Damages Actually Paid to Claimants.—In any health care lawsuit, the court shall supervise the arrangements for payment of damages to protect against conflicts of interest
that may have the effect of reducing the amount of dam-
ages awarded that are actually paid to claimants. In par-
ticular, in any health care lawsuit in which the attorney
for a party claims a financial stake in the outcome by vir-
tue of a contingent fee, the court shall have the power
to restrict the payment of a claimant’s damage recovery
to such attorney, and to redirect such damages to the
claimant based upon the interests of justice and principles
of equity. In no event shall the total of all contingent fees
for representing all claimants in a health care lawsuit ex-
ceed the following limits:

(1) 40 percent of the first $50,000 recovered by
the claimant(s).

(2) 33\(\frac{1}{3}\) percent of the next $50,000 recovered
by the claimant(s).

(3) 25 percent of the next $500,000 recovered
by the claimant(s).

(4) 15 percent of any amount by which the re-
covery by the claimant(s) is in excess of $600,000.

(b) APPLICABILITY.—The limitations in this section
shall apply whether the recovery is by judgment, settle-
ment, mediation, arbitration, or any other form of alter-
native dispute resolution. In a health care lawsuit involv-
ing a minor or incompetent person, a court retains the
authority to authorize or approve a fee that is less than
the maximum permitted under this section. The requirement for court supervision in the first two sentences of subsection (a) applies only in civil actions.

SEC. 206. ADDITIONAL HEALTH BENEFITS.

In any health care lawsuit involving injury or wrongful death, any party may introduce evidence of collateral source benefits. If a party elects to introduce such evidence, any opposing party may introduce evidence of any amount paid or contributed or reasonably likely to be paid or contributed in the future by or on behalf of the opposing party to secure the right to such collateral source benefits. No provider of collateral source benefits shall recover any amount against the claimant or receive any lien or credit against the claimant’s recovery or be equitably or legally subrogated to the right of the claimant in a health care lawsuit involving injury or wrongful death. This section shall apply to any health care lawsuit that is settled as well as a health care lawsuit that is resolved by a fact finder. This section shall not apply to section 1862(b) (42 U.S.C. 1395y(b)) or section 1902(a)(25) (42 U.S.C. 1396a(a)(25)) of the Social Security Act, or to section 8131 or section 8132 of title 5, United States Code. This section shall not apply to section 1862(b) (42 U.S.C. 1395y(b)) or section 1902(a)(25) (42 U.S.C. 1396a(a)(25)) of the Social Security Act, or to section
8131 or section 8132 of title 5, United States Code, or to a collateral source provider that is an employee benefit plan under section 3(3) of the Employee Retirement Income Security Act of 1974 (29 U.S.C. 1002(3)).

SEC. 207. PUNITIVE DAMAGES.

(a) IN GENERAL.—Punitive damages may, if otherwise permitted by applicable State or Federal law, be awarded against any person in a health care lawsuit only if it is proven by clear and convincing evidence that such person acted with malicious intent to injure the claimant, or that such person deliberately failed to avoid unnecessary injury that such person knew the claimant was substantially certain to suffer. In any health care lawsuit where no judgment for compensatory damages is rendered against such person, no punitive damages may be awarded with respect to the claim in such lawsuit. No demand for punitive damages shall be included in a health care lawsuit as initially filed. A court may allow a claimant to file an amended pleading for punitive damages only upon a motion by the claimant and after a finding by the court, upon review of supporting and opposing affidavits or after a hearing, after weighing the evidence, that the claimant has established by a substantial probability that the claimant will prevail on the claim for punitive damages. At the re-
quest of any party in a health care lawsuit, the trier of fact shall consider in a separate proceeding—

(1) whether punitive damages are to be awarded and the amount of such award; and

(2) the amount of punitive damages following a determination of punitive liability.

If a separate proceeding is requested, evidence relevant only to the claim for punitive damages, as determined by applicable State law, shall be inadmissible in any proceeding to determine whether compensatory damages are to be awarded.

(b) Determining Amount of Punitive Damages.—

(1) Factors Considered.—In determining the amount of punitive damages, if awarded, in a health care lawsuit, the trier of fact shall consider only the following—

(A) the severity of the harm caused by the conduct of such party;

(B) the duration of the conduct or any concealment of it by such party;

(C) the profitability of the conduct to such party;

(D) the number of products sold or medical procedures rendered for compensation, as
the case may be, by such party, of the kind
causing the harm complained of by the claim-
ant;

(E) any criminal penalties imposed on such
party, as a result of the conduct complained of
by the claimant; and

(F) the amount of any civil fines assessed
against such party as a result of the conduct
complained of by the claimant.

(2) MAXIMUM AWARD.—The amount of punitive
damages, if awarded, in a health care lawsuit may
be as much as $250,000 or as much as two times
the amount of economic damages awarded, which-
ever is greater. The jury shall not be informed of
this limitation.

(e) NO PUNITIVE DAMAGES FOR PRODUCTS THAT
COMPLY WITH FDA STANDARDS.—

(1) IN GENERAL.—

(A) No punitive damages may be awarded
against the manufacturer or distributor of a
medical product, or a supplier of any compo-
nent or raw material of such medical product,
based on a claim that such product caused the
claimant’s harm where—
(i)(I) such medical product was subject to premarket approval, clearance, or licensure by the Food and Drug Administration with respect to the safety of the formulation or performance of the aspect of such medical product which caused the claimant’s harm or the adequacy of the packaging or labeling of such medical product; and

(II) such medical product was so approved, cleared, or licensed; or

(ii) such medical product is generally recognized among qualified experts as safe and effective pursuant to conditions established by the Food and Drug Administration and applicable Food and Drug Administration regulations, including without limitation those related to packaging and labeling, unless the Food and Drug Administration has determined that such medical product was not manufactured or distributed in substantial compliance with applicable Food and Drug Administration statutes and regulations.
(B) RULE OF CONSTRUCTION.—Subparagraph (A) may not be construed as establishing
the obligation of the Food and Drug Adminis-
tration to demonstrate affirmatively that a
manufacturer, distributor, or supplier referred
to in such subparagraph meets any of the con-
ditions described in such subparagraph.

(2) LIABILITY OF HEALTH CARE PROVIDERS.—
A health care provider who prescribes, or who dis-
penses pursuant to a prescription, a medical product
approved, licensed, or cleared by the Food and Drug
Administration shall not be named as a party to a
product liability lawsuit involving such product and
shall not be liable to a claimant in a class action
lawsuit against the manufacturer, distributor, or
seller of such product. Nothing in this paragraph
prevents a court from consolidating cases involving
health care providers and cases involving products li-
ability claims against the manufacturer, distributor,
or product seller of such medical product.

(3) PACKAGING.—In a health care lawsuit for
harm which is alleged to relate to the adequacy of
the packaging or labeling of a drug which is required
to have tamper-resistant packaging under regula-
tions of the Secretary of Health and Human Serv-
ices (including labeling regulations related to such packaging), the manufacturer or product seller of the drug shall not be held liable for punitive damages unless such packaging or labeling is found by the trier of fact by clear and convincing evidence to be substantially out of compliance with such regulations.

(4) EXCEPTION.—Paragraph (1) shall not apply in any health care lawsuit in which—

(A) a person, before or after premarket approval, clearance, or licensure of such medical product, knowingly misrepresented to or withheld from the Food and Drug Administration information that is required to be submitted under the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.) or section 351 of the Public Health Service Act (42 U.S.C. 262) that is material and is causally related to the harm which the claimant allegedly suffered; or

(B) a person made an illegal payment to an official of the Food and Drug Administration for the purpose of either securing or maintaining approval, clearance, or licensure of such medical product.
SEC. 208. AUTHORIZATION OF PAYMENT OF FUTURE DAMAGES TO CLAIMANTS IN HEALTH CARE LAWSUITS.

(a) In General.—In any health care lawsuit, if an award of future damages, without reduction to present value, equaling or exceeding $50,000 is made against a party with sufficient insurance or other assets to fund a periodic payment of such a judgment, the court shall, at the request of any party, enter a judgment ordering that the future damages be paid by periodic payments. In any health care lawsuit, the court may be guided by the Uniform Periodic Payment of Judgments Act promulgated by the National Conference of Commissioners on Uniform State Laws.

(b) Applicability.—This section applies to all actions which have not been first set for trial or retrial before the effective date of this Act.

SEC. 209. DEFINITIONS.

In this title:

(1) Alternative dispute resolution system; ADR.—The term “alternative dispute resolution system” or “ADR” means a system that provides for the resolution of health care lawsuits in a manner other than through a civil action brought in a State or Federal court.
(2) CLAIMANT.—The term “claimant” means any person who brings a health care lawsuit, including a person who asserts or claims a right to legal or equitable contribution, indemnity or subrogation, arising out of a health care liability claim or action, and any person on whose behalf such a claim is asserted or such an action is brought, whether deceased, incompetent, or a minor.

(3) COLLATERAL SOURCE BENEFITS.—The term “collateral source benefits” means any amount paid or reasonably likely to be paid in the future to or on behalf of the claimant, or any service, product or other benefit provided or reasonably likely to be provided in the future to or on behalf of the claimant, as a result of the injury or wrongful death, pursuant to—

(A) any State or Federal health, sickness, income-disability, accident, or workers’ compensation law (except the Federal Employees’ Compensation Act (5 U.S.C. 8101 et seq.));

(B) any health, sickness, income-disability, or accident insurance that provides health benefits or income-disability coverage;

(C) any contract or agreement of any group, organization, partnership, or corporation
to provide, pay for, or reimburse the cost of medical, hospital, dental, or income disability benefits; and

(D) any other publicly or privately funded program.

(4) COMPENSATORY DAMAGES.—The term “compensatory damages” means objectively verifiable monetary losses incurred as a result of the provision of, use of, or payment for (or failure to provide, use, or pay for) health care services or medical products, such as past and future medical expenses, loss of past and future earnings, cost of obtaining domestic services, loss of employment, and loss of business or employment opportunities, damages for physical and emotional pain, suffering, inconvenience, physical impairment, mental anguish, disfigurement, loss of enjoyment of life, loss of society and companionship, loss of consortium (other than loss of domestic service), hedonic damages, injury to reputation, and all other nonpecuniary losses of any kind or nature. The term “compensatory damages” includes economic damages and non-economic damages, as such terms are defined in this section.
(5) **CONTINGENT FEE.**—The term “contingent fee” includes all compensation to any person or persons which is payable only if a recovery is effected on behalf of one or more claimants.

(6) **ECONOMIC DAMAGES.**—The term “economic damages” means objectively verifiable monetary losses incurred as a result of the provision of, use of, or payment for (or failure to provide, use, or pay for) health care services or medical products, such as past and future medical expenses, loss of past and future earnings, cost of obtaining domestic services, loss of employment, and loss of business or employment opportunities.

(7) **HEALTH CARE LAWSUIT.**—The term “health care lawsuit” means any health care liability claim concerning the provision of health care goods or services or any medical product affecting interstate commerce, or any health care liability action concerning the provision of health care goods or services or any medical product affecting interstate commerce, brought in a State or Federal court or pursuant to an alternative dispute resolution system, against a health care provider, a health care organization, or the manufacturer, distributor, supplier, marketer, promoter, or seller of a medical product,
regardless of the theory of liability on which the claim is based, or the number of claimants, plaintiffs, defendants, or other parties, or the number of claims or causes of action, in which the claimant alleges a health care liability claim. Such term does not include a claim brought by the United States Government or a relator under the False Claims Act (31 U.S.C. 3729 et seq.) or a claim or action which is based on criminal liability; which seeks civil fines or penalties paid to Federal, State, or local government; or which is grounded in antitrust.

(8) Health care liability action.—The term “health care liability action” means a civil action brought in a State or Federal Court or pursuant to an alternative dispute resolution system, against a health care provider, a health care organization, or the manufacturer, distributor, supplier, marketer, promoter, or seller of a medical product, regardless of the theory of liability on which the claim is based, or the number of plaintiffs, defendants, or other parties, or the number of causes of action, in which the claimant alleges a health care liability claim.

(9) Health care liability claim.—The term “health care liability claim” means a demand
by any person, whether or not pursuant to ADR, against a health care provider, health care organization, or the manufacturer, distributor, supplier, marketer, promoter, or seller of a medical product, including, but not limited to, third-party claims, cross-claims, counter-claims, or contribution claims, which are based upon the provision of, use of, or payment for (or the failure to provide, use, or pay for) health care services or medical products, regardless of the theory of liability on which the claim is based, or the number of plaintiffs, defendants, or other parties, or the number of causes of action.

(10) **HEALTH CARE ORGANIZATION.**—The term “health care organization” means any person or entity which is obligated to provide or pay for health benefits under any health plan, including any person or entity acting under a contract or arrangement with a health care organization to provide or administer any health benefit.

(11) **HEALTH CARE PROVIDER.**—The term “health care provider” means any person or entity required by State or Federal laws or regulations to be licensed, registered, or certified to provide health care services, and being either so licensed, reg-
istered, or certified, or exempted from such require-
ment by other statute or regulation.

(12) HEALTH CARE GOODS OR SERVICES.—The
term “health care goods or services” means any
goods or services provided by a health care organiza-
tion, provider, or by any individual working under
the supervision of a health care provider, that relates
to the diagnosis, prevention, or treatment of any
human disease or impairment, or the assessment or
care of the health of human beings.

(13) MALICIOUS INTENT TO INJURE.—The
term “malicious intent to injure” means inten-
tionally causing or attempting to cause physical in-
jury other than providing health care goods or serv-
ices.

(14) MEDICAL PRODUCT.—The term “medical
product” means a drug, device, or biological product
intended for humans, and the terms “drug”, “de-
vice”, and “biological product” have the meanings
given such terms in sections 201(g)(1) and 201(h)
of the Federal Food, Drug and Cosmetic Act (21
U.S.C. 321) and section 351(a) of the Public Health
Service Act (42 U.S.C. 262(a)), respectively, includ-
ing any component or raw material used therein, but
excluding health care services.
(15) NONECONOMIC DAMAGES.—The term “noneconomic damages” means damages for physical and emotional pain, suffering, inconvenience, physical impairment, mental anguish, disfigurement, loss of enjoyment of life, loss of society and companionship, loss of consortium (other than loss of domestic service), hedonic damages, injury to reputation, and all other nonpecuniary losses of any kind or nature.

(16) PUNITIVE DAMAGES.—The term “punitive damages” means damages awarded, for the purpose of punishment or deterrence, and not solely for compensatory purposes, against a health care provider, health care organization, or a manufacturer, distributor, or supplier of a medical product. Punitive damages are neither economic nor noneconomic damages.

(17) RECOVERY.—The term “recovery” means the net sum recovered after deducting any disbursements or costs incurred in connection with prosecution or settlement of the claim, including all costs paid or advanced by any person. Costs of health care incurred by the plaintiff and the attorneys’ office overhead costs or charges for legal services are not deductible disbursements or costs for such purpose.
(18) STATE.—The term “State” means each of the several States, the District of Columbia, the Commonwealth of Puerto Rico, the Virgin Islands, Guam, American Samoa, the Northern Mariana Islands, the Trust Territory of the Pacific Islands, and any other territory or possession of the United States, or any political subdivision thereof.

SEC. 210. EFFECT ON OTHER LAWS.

(a) VACCINE INJURY.—

(1) To the extent that title XXI of the Public Health Service Act establishes a Federal rule of law applicable to a civil action brought for a vaccine-related injury or death—

(A) this title does not affect the application of the rule of law to such an action; and

(B) any rule of law prescribed by this title in conflict with a rule of law of such title XXI shall not apply to such action.

(2) If there is an aspect of a civil action brought for a vaccine-related injury or death to which a Federal rule of law under title XXI of the Public Health Service Act does not apply, then this title or otherwise applicable law (as determined under this title) will apply to such aspect of such action.
(b) **Other Federal Law.**—Except as provided in this section, nothing in this title shall be deemed to affect any defense available to a defendant in a health care lawsuit or action under any other provision of Federal law.

**Sec. 211. State Flexibility and Protection of States' Rights.**

(a) **Health Care Lawsuits.**—The provisions governing health care lawsuits set forth in this title preempt, subject to subsections (b) and (c), State law to the extent that State law prevents the application of any provisions of law established by or under this title. The provisions governing health care lawsuits set forth in this title supersede chapter 171 of title 28, United States Code, to the extent that such chapter—

1. provides or allows for a greater amount of damages or contingent fees, or a longer period in which a health care lawsuit may be commenced, than provided in this title;

2. precludes or reduces the applicability or scope of periodic payment of future damages as provided in this title; or

3. through application of State law, conflicts with provisions of this title concerning joint liability, collateral source benefits, subrogation, or liens.
(b) Protection of States’ Rights and Other Laws.—

(1) Any issue that is not governed by any provision of law established by or under this title (including State standards of negligence) shall be governed by otherwise applicable State or Federal law.

(2) This title shall not preempt or supersede any State or Federal law that imposes greater procedural or substantive protections for health care providers and health care organizations from liability, loss, or damages than those provided by this title or create a cause of action.

(e) State Flexibility.—No provision of this title shall be construed to preempt—

(1) any State law (whether effective before, on, or after the date of the enactment of this title) that specifies a particular monetary amount of compensatory or punitive damages (or the total amount of damages) that may be awarded in a health care lawsuit, regardless of whether such monetary amount is greater or lesser than is provided for under this title, notwithstanding section 204(a); or

(2) any defense available to a party in a health care lawsuit under any other provision of State or Federal law.
SEC. 212. APPLICABILITY; EFFECTIVE DATE.

This title shall apply to any health care lawsuit brought in a Federal or State court, or subject to an alternative dispute resolution system, that is initiated on or after the date of the enactment of this title, except that any health care lawsuit arising from an injury occurring prior to the date of the enactment of this title shall be governed by the applicable statute of limitations provisions in effect at the time the injury occurred.

TITLE III—INCREASING HIGH-INCOME BENEFICIARY AWARENESS AND RESPONSIBILITY FOR HEALTH CARE COSTS

SEC. 301. INCOME-RELATED REDUCTION IN PART D PREMIUM SUBSIDY.

(a) Income-Related Reduction in Part D Premium Subsidy.—

(1) In general.—Section 1860D–13(a) (42 U.S.C. 1395w–113(a)) is amended by adding at the end the following new paragraph:

“(7) Reduction in premium subsidy based on income.—

“(A) In general.—In the case of an individual whose modified adjusted gross income exceeds the threshold amount applicable under subparagraph (B) for the calendar year, the
monthly amount of the premium subsidy applicable to the premium under this section for a month after December 2008 shall be reduced (and the monthly beneficiary premium shall be increased) by the monthly adjustment amount specified in subparagraph (C).

“(B) Threshold amount.—For purposes of this paragraph, the threshold amount is—

“(i) except as provided in clause (ii), $82,000; and

“(ii) in the case of a joint return, twice the amount applicable under clause (i) for the calendar year.

“(C) Monthly adjustment amount.—

“(i) In general.—The monthly adjustment amount specified in this subparagraph for an individual for a month in a year is equal to the product of—

“(I) the quotient obtained by dividing—

“(aa) the applicable percentage specified in the table in clause (ii) for the individual for the calendar year reduced by

25.5 percent; by
“(bb) 25.5 percent; and

“(II) the base beneficiary premium (as computed under paragraph (2)).

“(ii) APPLICABLE PERCENTAGE.—

“(I) IN GENERAL.—

<table>
<thead>
<tr>
<th>If the modified adjusted gross income is:</th>
<th>The applicable percentage is:</th>
</tr>
</thead>
<tbody>
<tr>
<td>More than $82,000 but not more than $102,000</td>
<td>35 percent</td>
</tr>
<tr>
<td>More than $102,000 but not more than $153,000</td>
<td>50 percent</td>
</tr>
<tr>
<td>More than $153,000 but not more than $205,000</td>
<td>65 percent</td>
</tr>
<tr>
<td>More than $205,000</td>
<td>80 percent</td>
</tr>
</tbody>
</table>

“(II) JOINT RETURNS.—In the case of a joint return, subclause (I) shall be applied by substituting dollar amounts which are twice the dollar amounts otherwise applicable under subclause (I) for the calendar year.

“(III) MARRIED INDIVIDUALS FILING SEPARATE RETURNS.—In the case of an individual who—

“(aa) is married as of the close of the taxable year (within the meaning of section 7703 of the Internal Revenue Code of 1986) but does not file a joint return for such year, and
“(bb) does not live apart
from such individual’s spouse at
all times during the taxable year,
subclause (I) shall be applied by re-
ducing each of the dollar amounts
otherwise applicable under such sub-
clause for the calendar year by the
threshold amount for such year appli-
cable to an unmarried individual.

“(D) Determination by Commissioner
of Social Security.—The Commissioner of
Social Security shall have the authority to make
initial and reconsideration determinations nec-
essary to carry out the income-related reduction
in premium subsidy under this paragraph.

“(E) Modified Adjusted Gross In-
come.—For purposes of this paragraph, the
term ‘modified adjusted gross income’ has the
meaning given such term in subparagraph (A)
of section 1839(i)(4), determined for the tax-
able year applicable under subparagraphs (B)
and (C) of such section.

“(F) Joint Return Defined.—For pur-
poses of this paragraph, the term ‘joint return’
has the meaning given to such term by section

“(G) PROCEDURES TO ASSURE CORRECT INCOME-RELATED REDUCTION IN PREMIUM SUBSIDY.—

“(i) DISCLOSURE OF BASE BENEFICIARY PREMIUM.—Not later than September 15 of each year beginning with 2008, the Secretary shall disclose to the Commissioner of Social Security the amount of the base beneficiary premium (as computed under paragraph (2)) for the purpose of carrying out the income-related reduction in premium subsidy under this paragraph with respect to the following year.

“(ii) ADDITIONAL DISCLOSURE.—Not later than October 15 of each year beginning with 2008, the Secretary shall disclose to the Commissioner of Social Security the following information for the purpose of carrying out the income-related reduction in premium subsidy under this paragraph with respect to the following year:
“(I) The monthly adjustment amount specified in subparagraph (C).

“(II) Any other information the Commissioner of Social Security determines necessary to carry out the income-related reduction in premium subsidy under this paragraph.

“(H) RULE OF CONSTRUCTION.—The formula used to determine the monthly adjustment amount specified under subparagraph (C) shall only be used for the purpose of determining such monthly adjustment amount under such subparagraph.”.

(2) COLLECTION OF MONTHLY ADJUSTMENT AMOUNT.—Section 1860D–13(c) (42 U.S.C. 1395w–113(c)) is amended—

(A) in paragraph (1), by striking “(2) and (3)” and inserting “(2), (3), and (4)”; and

(B) by adding at the end the following new paragraph:

“(4) COLLECTION OF MONTHLY ADJUSTMENT AMOUNT.—

“(A) IN GENERAL.—Notwithstanding any provision of this subsection or section 1854(d)(2), subject to subparagraph (B), the
amount of the income-related reduction in pre-
mium subsidy for an individual for a month (as
determined under subsection (a)(7)) shall be
paid through withholding from benefit pay-
ments in the manner provided under section
1840.

“(B) AGREEMENTS.—In the case where
the monthly benefit payments of an individual
that are withheld under subparagraph (A) are
insufficient to pay the amount described in such
subparagraph, the Commissioner of Social Se-
curity shall enter into agreements with the Sec-
retary, the Director of the Office of Personnel
Management, and the Railroad Retirement
Board as necessary in order to allow other
agencies to collect the amount described in sub-
paragraph (A) that was not withheld under
such subparagraph.”.

(b) CONFORMING AMENDMENTS.—

(1) MEDICARE.—Part D of title XVIII (42
U.S.C. 1395w–101 et seq.) is amended—

(A) in section 1860D–13(a)(1)—

(i) by redesignating subparagraph (F)
as subparagraph (G);
(ii) in subparagraph (G), as redesignated by subparagraph (A), by striking “(D) and (E)” and inserting “(D), (E), and (F)”;

(iii) by inserting after subparagraph (E) the following new subparagraph:

“(F) INCREASE BASED ON INCOME.—The monthly beneficiary premium shall be increased pursuant to paragraph (7).”; and

(B) in section 1860D–15(a)(1)(B), by striking “paragraph (1)(B)” and inserting “paragraphs (1)(B) and (1)(F)”.

(2) INTERNAL REVENUE CODE.—Section 6103(l)(20) of the Internal Revenue Code of 1986 (relating to disclosure of return information to carry out Medicare part B premium subsidy adjustment) is amended—

(A) in the heading, by striking “PART B PREMIUM SUBSIDY ADJUSTMENT” and inserting “PARTS B AND D PREMIUM SUBSIDY ADJUSTMENTS”; and

(B) in subparagraph (A)—

(i) in the matter preceding clause (i), by inserting “or 1860D–13(a)(7)” after “1839(i)”;}
(ii) in clause (vii), by inserting after
“subsection (i) of such section” the fol-
lowing: “or under section 1860D–13(a)(7)
of such Act”; and

(C) in subparagraph (B)—

(i) by inserting “or such section
1860D–13(a)(7)” before the period at the
end;

(ii) as amended by clause (i), by add-
ing at the end the following new sentence:
“Such return information may be disclosed
to officers and employees of the Depart-
ments of Health and Human Services and
Justice, to the extent necessary, and solely
for their use, in any administrative or judi-
cial proceeding ensuing from an adjust-
ment to any such premium.”; and

(D) by adding at the end the following new
subparagraph:

“(C) TIMING OF DISCLOSURE.—Return in-
formation shall be disclosed to officers, employ-
ees, and contractors of the Social Security Ad-
ministration under subparagraph (A):

“(i) for taxpayers currently entitled to
benefits under title II of the Social Secu-
rity Act, or as qualified railroad retirement
beneficiaries within the meaning of section
7(d) of the Railroad Retirement Act of
1974, within 4 months preceding the
month in which the taxpayer first becomes
entitled to benefits under part A or is eligi-
ble to enroll in part B or part D of title
XVIII of the Social Security Act; and
“(ii) for taxpayers not currently re-
ceiving benefits under title II of the Social
Security Act, or as qualified railroad re-
tirement beneficiaries within the meaning
of section 7(d) of the Railroad Retirement
Act of 1974, or who have participated in
Medicare qualified government employment
as defined in section 210(p) of the Social
Security Act, after the taxpayer applies for
a benefit under part A or part B and is eli-
gible to enroll in part D of title XVIII of
the Social Security Act.”.

(c) IMPLEMENTATION.—Notwithstanding any other
provision of law, the Secretary, in consultation with the
Commissioner of Social Security may implement this sec-
tion, and the amendments made by this section, by pro-
gram instruction or otherwise.