

112TH CONGRESS  
2D SESSION

# H. R. 6043

To amend the Public Health Service Act and the Social Security Act to extend health information technology assistance eligibility to behavioral health, mental health, and substance abuse professionals and facilities, and for other purposes.

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## IN THE HOUSE OF REPRESENTATIVES

JUNE 27, 2012

Mr. MURPHY of Pennsylvania (for himself, Mr. RYAN of Ohio, Mr. MARINO, Mr. SULLIVAN, Mrs. BLACKBURN, and Mr. TIBERI) 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

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## A BILL

To amend the Public Health Service Act and the Social Security Act to extend health information technology assistance eligibility to behavioral health, mental health, and substance abuse professionals and facilities, 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 “Behavioral Health In-  
5       formation Technology Act of 2012”.

**1 SEC. 2. EXTENSION OF HEALTH INFORMATION TECHNOLOGY ASSISTANCE FOR BEHAVIORAL AND**

**2 MENTAL HEALTH AND SUBSTANCE ABUSE.**

4 Section 3000(3) of the Public Health Service Act (42  
5 U.S.C. 300jj(3)) is amended by inserting before “and any  
6 other category” the following: “behavioral and mental  
7 health professionals (as defined in section  
8 331(a)(3)(E)(i)), a substance abuse professional, a psy-  
9 chiatric hospital (as defined in section 1861(f) of the So-  
10 cial Security Act (42 U.S.C. 1395x(f))), a community  
11 mental health center meeting the criteria specified in sec-  
12 tion 1913(c), a residential or outpatient mental health or  
13 substance abuse treatment facility.”.

14 SEC. 3. EXTENSION OF ELIGIBILITY FOR MEDICARE AND  
15 MEDICAID HEALTH INFORMATION TECH-  
16 NOLOGY IMPLEMENTATION ASSISTANCE.

17 (a) PAYMENT INCENTIVES FOR ELIGIBLE PROFESSIONALS UNDER MEDICARE.—Section 1848 of the Social  
18 Security Act (42 U.S.C. 1395w-4) is amended—  
19

20 (1) by amending clause (iii) of subsection  
21 (a)(7)(E) to read as follows:

22                             “(iii) ELIGIBLE PROFESSIONAL.—The  
23                             term ‘eligible professional’ means any of  
24                             the following:

1                         “(II) A clinical psychologist pro-  
2                         viding qualified psychologist services  
3                         (as defined in section 1861(ii)).”; and  
4                         (2) by amending subparagraph (C) of sub-  
5                         section (o)(5) to read as follows:

6                         “(C) ELIGIBLE PROFESSIONAL.—The term  
7                         ‘eligible professional’ means any of the fol-  
8                         lowing:

9                         “(i) A physician (as defined in section  
10                         1861(r)).

11                         “(ii) A clinical psychologist providing  
12                         qualified psychologist services (as defined  
13                         in section 1861(ii)).”.

14                         (b) ELIGIBLE HOSPITALS.—Section 1886(n)(6)(B)  
15                         of the Social Security Act (42 U.S.C. 1395ww(n)(6)(B))  
16                         is amended by inserting before the period at the end the  
17                         following: “or an inpatient hospital that is a psychiatric  
18                         hospital (as defined in section 1861(f))”.

19                         (c) MEDICAID PROVIDERS.—Section 1903(t) of the  
20                         Social Security Act (42 U.S.C. 1396b(t)) is amended as  
21                         follows:

22                         (1) Paragraph (2)(B) is amended—  
23                         (A) in clause (i), by striking “, or” and in-  
24                         serting a semicolon;

1                             (B) in clause (ii), by striking the period  
2                             and inserting a semicolon; and

3                             (C) by adding after clause (ii) the following  
4                             new clauses:

5                                 “(iii) a public hospital that is prin-  
6                                 cipally a psychiatric hospital (as defined in  
7                                 section 1861(f));

8                                 “(iv) a private hospital that is prin-  
9                                 cipally a psychiatric hospital (as defined in  
10                                 section 1861(f)) and that has at least 10  
11                                 percent of its patient volume (as estimated  
12                                 in accordance with a methodology estab-  
13                                 lished by the Secretary) attributable to in-  
14                                 dividuals receiving medical assistance  
15                                 under this title;

16                                 “(v) a community mental health cen-  
17                                 ter meeting the criteria specified in section  
18                                 1913(c) of the Public Health Service Act;  
19                                 or

20                                 “(vi) a residential or outpatient men-  
21                                 tal health or substance abuse treatment fa-  
22                                 cility that—

23                                 “(I) is accredited by the Joint  
24                                 Commission on Accreditation of  
25                                 Healthcare Organizations, the Com-

1 mission on Accreditation of Rehabili-  
2 tation Facilities, the Council on Ac-  
3 creditation, or any other national ac-  
4 crediting agency recognized by the  
5 Secretary; and

6 “(II) has at least 10 percent of  
7 its patient volume (as estimated in ac-  
8 cordance with a methodology estab-  
9 lished by the Secretary) attributable  
10 to individuals receiving medical assist-  
11 ance under this title.”.

12 (2) Paragraph (3)(B) is amended—

13 (A) in clause (iv), by striking “and” after  
14 the semicolon;

15 (B) in clause (v), by striking the period  
16 and inserting “; and”; and

17 (C) by adding at the end the following new  
18 clause:

19 “(vi) clinical psychologist providing  
20 qualified psychologist services (as defined  
21 in section 1861(ii)), if such clinical psy-  
22 chologist is practicing in an outpatient  
23 clinic that—

24 “(I) is led by a clinical psycholo-  
25 gist; and

1                         “(II) is not otherwise receiving  
2                         payment under paragraph (1) as a  
3                         Medicaid provider described in para-  
4                         graph (2)(B).”.

5 **SEC. 4. PROVIDING PROTECTIONS FOR CERTAIN PRO-**  
6 **VIDERS, VENDORS, AND USERS OF CERTIFIED**  
7 **EHR TECHNOLOGY.**

8 (a) COVERED ENTITIES.—

9                         (1) COVERED ENTITIES.—For purposes of this  
10          subsection, a covered entity means, with respect to  
11          certified EHR technology (as defined in section  
12          1848(o)(4) of the Social Security Act) and a year,  
13          any of the following:

14                         (A) MEANINGFUL EHR USERS.—Any of the  
15          following, with respect to such year:

16                         (i) An eligible professional (as defined  
17          in paragraph (5)(C) of section 1848(o) of  
18          the Social Security Act) determined to be  
19          a meaningful EHR user under paragraph  
20          (2) of such section for the EHR reporting  
21          period (as defined in paragraph (5)(B) of  
22          such section) during such year.

23                         (ii) In the case of a qualifying MA or-  
24          ganization (as defined in paragraph (5) of  
25          section 1853(l) of such Act), an eligible

1 professional described in paragraph (2) of  
2 such section of the organization who the  
3 organization attests under paragraph (6)  
4 of such section to be a meaningful EHR  
5 user for such year.

6 (iii) In the case of a qualifying MA  
7 organization (as defined in paragraph (5)  
8 of section 1853(l) of such Act), an eligible  
9 hospital described in section 1853(m)(2) of  
10 such Act of the organization which attests  
11 under section 1853(l)(6) of such Act to be  
12 a meaningful EHR user for the applicable  
13 period with respect to such year.

14 (iv) An eligible hospital (as defined in  
15 paragraph (6)(B) of section 1886(n) of  
16 such Act) determined to be a meaningful  
17 EHR user under paragraph (3) of such  
18 section for the EHR reporting period (as  
19 defined in paragraph (6)(A) of such sec-  
20 tion) with respect to such year.

21 (v) A critical access hospital deter-  
22 mined pursuant to section 1814(l)(3) of  
23 such Act to be a meaningful EHR user (as  
24 would be determined under paragraph (3)  
25 of section 1886(n) of such Act) for an

1                   EHR reporting period (as defined in para-  
2                   graph (6)(A) of such section) for a cost re-  
3                   porting period beginning during such year.

4                   (vi) A Medicaid provider (as defined  
5                   in paragraph (2) of section 1903(t) of such  
6                   Act) eligible for payments described in  
7                   paragraph (1) of such section for such  
8                   year.

9                   (B) HEALTH INFORMATION EXCHANGE  
10                  ENTITIES.—Individuals and entities (other than  
11                  States or State designated entities) which dur-  
12                  ing such year are health information exchange  
13                  contractors (consisting of technology providers),  
14                  health information exchange participants (con-  
15                  sisting of organizations providing supportive  
16                  technology to a health information exchange),  
17                  and other users of health information exchanges  
18                  (consisting of other entities that may be ex-  
19                  changing clinical or administrative data). Man-  
20                  ufacturers of EHR Software and other health  
21                  information technologies who participate in the  
22                  reporting of adverse events or who otherwise  
23                  contribute relevant patient safety work product  
24                  under subsection (c)(1) of this Act.

(C) CERTAIN OTHER EHR USERS.—A health care professional who, during such year—

(i) is a user of such certified EHR technology;

(ii) is not eligible for incentive payments based on meaningful use of such technology under title XVIII or XIX of the Social Security Act solely because the professional is not—

(I) an eligible professional (as defined in paragraph (5)(C) of section 1848(o) of such Act);

(II) an eligible professional described in paragraph (2) of section 1853(l) of such Act, with respect to a qualifying MA organization (as defined in paragraph (5) of such section);

(III) an eligible hospital described in section 1853(m)(2) of such Act, with respect to such a qualifying MA organization;

14 (b) IMPROVING PATIENT SAFETY THROUGH ERROR  
15 REPORTING AND REMEDIATION, AND CLARIFICATION OF  
16 AUTHORITY.—

17                             (1) IN GENERAL.—A covered entity may submit  
18                             to a Patient Safety Organization as defined in sec-  
19                             tion 921. Title IX of the Public Health Service Act  
20                             (42 U.S.C. 299 et seq.) information on EHR-related  
21                             adverse events with respect to certified EHR tech-  
22                             nology as defined in section 3001 of the Public  
23                             Health Service Act (42 U.S.C. 300jj–11) used or  
24                             provided by such entity, as applicable. The utiliza-  
25                             tion of patient safety work product shall be for the

1       purpose of providing direct feedback and assistance  
2       to covered entities to effectively minimize patient  
3       risk. Patient Safety Organizations may furnish the  
4       Office of the National Coordinator de-identified re-  
5       ports of their findings for the purposes of tracking  
6       the number and nature of such adverse events.

7                     (2) APPLICATION OF SAFETY ORGANIZATION  
8                     PRIVILEGE AND CONFIDENTIALITY PROTECTIONS.—  
9       In the case of a covered entity that submits to such  
10      a body information on such an adverse event and in  
11      the case of the collection and maintenance of such  
12      information by such a body, the provisions of section  
13      922 of the Public Health Service Act shall apply to  
14      such information and to the body and the entity in  
15      the same manner such provisions apply to patient  
16      safety work product and a patient safety organiza-  
17      tion and provider under part C of title IX of such  
18      Act.

19                     (3) CLARIFICATION OF AUTHORITY.—Certified  
20      EHR's shall not be considered a device for purposes  
21      of the Federal Food, Drug, and Cosmetic Act (21  
22      U.S.C. 301 et seq.).

23                     (c) RULES RELATING TO E-DISCOVERY.—In any  
24      health care lawsuit against a covered entity that is related  
25      to an EHR-related adverse event, with respect to certified

1 EHR technology used or provided by the covered entity,  
2 electronic discovery shall be limited to—

3 (1) information that is related to such EHR-re-  
4 lated adverse event; and  
5 (2) information from the period in which such  
6 EHR-related adverse event occurred.

7 (d) LEGAL PROTECTIONS FOR COVERED ENTI-  
8 TIES.—

9 (1) GENERAL.—For a covered entity described  
10 in subsection (b), the following protections apply:

11 (A) ENCOURAGING SPEEDY RESOLUTION  
12 OF CLAIMS.—

13 (i) GENERAL.—A claimant may not  
14 commence a health care lawsuit against a  
15 covered entity on any date that is 3 years  
16 after the date of manifestation of injury or  
17 1 year after the claimant discovers, or  
18 through the use of reasonable diligence  
19 should have discovered, the injury, whichever  
20 occurs first. This limitation shall be  
21 tolled to the extent that the claimant is  
22 able to prove—

23 (I) fraud;  
24 (II) intentional concealment; or

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

(B) EQUITABLE ASSIGNMENT OF RESPONSIBILITY.—In any health care lawsuit against a covered entity—

4 (i) each party to the lawsuit other  
5 than the claimant that is such a covered  
6 entity shall be liable for that party's sev-  
7 eral share of any damages only and not for  
8 the share of any other person and such  
9 several share shall be in direct proportion  
10 to that party's proportion of responsibility  
11 for the injury, as determined under clause  
12 (iii);

22 (C) SUBSEQUENT REMEDIAL MEASURES.—  
23 Evidence of subsequent remedial measures to  
24 an EHR-related adverse event with respect to  
25 certified EHR technology used or provided by

1           the covered entity (including changes to the cer-  
2           tified EHR system, additional training require-  
3           ments, or changes to standard operating proce-  
4           dures) by a covered entity shall not be admis-  
5           sible in health care lawsuits.

6           (D) INCREASED BURDEN OF PROOF PRO-  
7           TECTION FOR COVERED ENTITIES.—Punitive  
8           damages may, if otherwise permitted by appli-  
9           cable State or Federal law, be awarded against  
10          any covered entity in a health care lawsuit only  
11          if it is proven by clear and convincing evidence  
12          that such entity acted with reckless disregard  
13          for the health or safety of the claimant. In any  
14          such health care lawsuit where no judgment for  
15          compensatory damages is rendered against such  
16          entity, no punitive damages may be awarded  
17          with respect to the claim in such lawsuit.

18           (E) PROTECTION FROM LIBEL OR SLAN-  
19           DER.—Covered entities and employees, agents  
20          and representatives of covered entities are im-  
21          mune from civil action for libel or slander arising  
22          from information or entries made in cer-  
23          tified EHR technology and for the transfer of  
24          such information to another eligible provider,  
25          hospital or health information exchange, if the

1           information, transfer of information, or entries  
2           were made in good faith and without malice.

3           (e) DEFINITIONS.—

4           (1) CLAIMANT.—The term “claimant” means  
5           any person who brings a health care lawsuit, includ-  
6           ing a person who asserts or claims a right to legal  
7           or equitable contribution, indemnity, or subrogation,  
8           arising out of a health care liability claim or action,  
9           and any person on whose behalf such a claim is as-  
10          serted or such an action is brought, whether de-  
11          ceased, incompetent, or a minor.

12          (2) COMPENSATORY DAMAGES.—The term  
13          “compensatory damages” means objectively verifi-  
14          able monetary losses incurred as a result of the pro-  
15          visions of, use of, or payment for (or failure to pro-  
16          vide, use, or pay for) health care services or medical  
17          products, such as past and future medical expenses,  
18          loss of past and future earnings, cost of obtaining  
19          domestic services, loss of employment, and loss of  
20          business or employment opportunities, damages for  
21          physical and emotional pain, suffering, inconven-  
22          ience, physical impairment, mental anguish, dis-  
23          figurement, loss of enjoyment in life, loss of society  
24          and companionship, loss of consortium (other than  
25          loss of domestic service), hedonic damages, injury to

1       reputation, and all other nonpecuniary losses of any  
2       kind or nature. Such term includes economic dam-  
3       ages and noneconomic damages, as such terms as  
4       defined in this subsection.

5                     (3) ECONOMIC DAMAGES.—The term “economic  
6       damages” means objectively verifiable monetary  
7       losses incurred as a result of the provisions of, use  
8       of, or payment for (or failure to provide, use, or pay  
9       for) health care services or medical products, such as  
10      past and future medical expenses, loss of past and  
11      future earnings, cost of obtaining domestic services,  
12      loss of employment, and loss of business or employ-  
13      ment opportunities.

14                    (4) CERTIFIED EHR TECHNOLOGY.—The term  
15      “certified EHR technology” has the meaning given  
16      such term in section 1848(o)(4) of the Social Secu-  
17      rity Act.

18                   (5) EHR-RELATED ADVERSE EVENT.—The  
19      term “EHR-related adverse event” means, with re-  
20      spect to a provider, a defect, malfunction, or error  
21      in the certified health information technology or  
22      electronic health record used by the provider, or in  
23      the input or output of data maintained through such  
24      technology or record, that results or could reason-  
25      ably result in harm to a patient.

1                             (6) HEALTH CARE LAWSUIT.—The term  
2        “health care lawsuit” means any health care liability  
3        claim concerning the provision of health care items  
4        or services or any medical product affecting inter-  
5        state commerce, or any health care liability action  
6        concerning the provision of health care items or  
7        services or any medical product affecting interstate  
8        commerce, brought in a State or Federal court or  
9        pursuant to an alternative dispute resolution system,  
10      against a health care provider, a health care organi-  
11      zation, or the manufacturer, distributor, supplier,  
12      marketer, promoter, or seller of a medical product,  
13      regardless of the theory of liability on which the  
14      claim is based, or the number of claimants, plain-  
15      tiffs, defendants, or other parties, or the number of  
16      claims or causes of action, in which the claimant al-  
17      leges a health care liability claim. Such term does  
18      not include a claim or action which is based on  
19      criminal liability; which seeks civil fines or penalties  
20      paid to Federal, State, or local government; or which  
21      is grounded in antitrust.

22                             (7) HEALTH CARE LIABILITY ACTION.—The  
23        term “health care liability action” means a civil ac-  
24        tion brought in a State or Federal court or pursuant  
25        to an alternative dispute resolution system, against

1       a health care provider, a health care organization, or  
2       the manufacturer, distributor, supplier, marketer,  
3       promoter, or seller of a medical product, regardless  
4       of the theory of liability on which the claim is based,  
5       or the number of plaintiffs, defendants, or other par-  
6       ties, or the number of causes of action, in which the  
7       claimant alleges a health care liability claim.

8                     (8) HEALTH CARE LIABILITY CLAIM.—The  
9       term “health care liability claim” means a demand  
10      by any person, whether or not pursuant to alter-  
11      native dispute resolution, against a health care pro-  
12      vider, health care organization, or the manufacturer,  
13      distributor, supplier, marketer, promoter, or seller of  
14      a medical product, including third-party claims,  
15      cross-claims, counter-claims, or contribution claims,  
16      which are based upon the provision of, use of, or  
17      payment for (or the failure to provide, use or pay  
18      for) health care services or medical products, regard-  
19      less of the theory of liability on which the claim is  
20      based, or the number of plaintiffs, defendants, or  
21      other parties, or the number of causes of action.

22                     (9) HEALTH CARE ORGANIZATION.—The term  
23      “health care organization” means any person or en-  
24      tity which is obligated to provide or pay for health  
25      benefits under any health plan, including any person

1       or entity acting under a contract or arrangement  
2       with a health care organization to provide or admin-  
3       ister any health benefit.

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

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

19                  (12) MALICIOUS INTENT TO INJURE.—The  
20       term “malicious intent to injure” means inten-  
21       tionally causing or attempting to cause physical in-  
22       jury other than providing health care items or serv-  
23       ices.

24                  (13) MEDICAL PRODUCT.—The term “medical  
25       product” means a drug, device, or biological product

1 intended for humans, and the terms “drug”, “de-  
2 vice”, and “biological product” have the meanings  
3 given such terms in sections 201(g)(1) and 201(h)  
4 of the Federal Food, Drug, and Cosmetic Act (21  
5 U.S.C. 321(g)(1) and (h)) and section 351(a) of the  
6 Public Health Service Act (42 U.S.C. 262(a)), re-  
7 spectively, including any component or raw material  
8 used therein, but excluding health care services.

9                     (14) NONECONOMIC DAMAGES.—The term  
10       “noneconomic damages” means damages for phys-  
11       ical impairment, mental anguish, disfigurement, loss  
12       of enjoyment of life, loss of society and companion-  
13       ship, loss of consortium (other than loss of domestic  
14       service), hedonic damages, injury to reputation, and  
15       all other nonpecuniary losses of any kind of nature.

16                     (15) PUNITIVE DAMAGES.—The term “punitive  
17       damages” means damages awarded, for the purpose  
18       of punishment or deterrence, and not solely for com-  
19       pensatory purposes, against a health care provider,  
20       health care organization, or a manufacturer, dis-  
21       tributor, or supplier of a medical product. Punitive  
22       damages are neither economic nor noneconomic dam-  
23       ages.

24                     (16) STATE.—The term “State” means each of  
25       the several States, District of Columbia, the Com-

1 monwealth of Puerto Rico, the Virgin Islands,  
2 Guam, American Samoa, the Northern Mariana Is-  
3 lands, the Trust Territory of the Pacific Islands, and  
4 any other territory or possession of the United  
5 States, or any political subdivision thereof.

