

111TH CONGRESS  
2D SESSION

# H. R. 5742

To encourage the use of medical checklists through research, and for other purposes.

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## IN THE HOUSE OF REPRESENTATIVES

JULY 15, 2010

Mr. HOLT (for himself and Mrs. CAPPS) introduced the following bill; which was referred to the Committee on Energy and Commerce

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## A BILL

To encourage the use of medical checklists through research, 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 “Medical Checklist Act  
5       of 2010”.

6       **SEC. 2. RESEARCH INTO MEDICAL CHECKLIST DEVELOP-**  
7       **MENT AND EFFICACY.**

8       (a) STUDY.—The Director of the Agency for  
9       Healthcare Research and Quality, acting through the Cen-  
10      ter for Quality Improvement and Patient Safety, shall con-

1 duct research and a study, in accordance with the require-  
2 ments of this section, regarding the development and effi-  
3 cacy of medical checklists.

4 (b) CONTENTS.—In carrying out subsection (a), the  
5 Director shall conduct research and a study regarding the  
6 following:

7 (1) Testing of different models of medical  
8 checklists to measure the effect of checklist format,  
9 length, and design for different clinical tasks on—

10 (A) adoption of checklists by health care  
11 professionals;

12 (B) time spent by health care professionals  
13 on the clinical task of interest; and

14 (C) reliable completion of medical proce-  
15 dures.

16 (2) Examination of checklist development and  
17 use in other industries, such as commercial aviation  
18 and nuclear power, and the feasibility of applying  
19 and adapting methodology developed in those indus-  
20 tries to the health care industry in a way that would  
21 result in health care quality improvement.

22 (3) Identification of organizational characteris-  
23 tics needed to effectively implement the use of med-  
24 ical checklists in health care settings.

1           (4) Measurement of the effects of the use of  
2           medical checklists on patient safety and health out-  
3           comes.

4           (5) Identification of medical procedures for  
5           which the development and use of medical checklists  
6           would be beneficial.

7           (6) Investigation of the development, implemen-  
8           tation, and use of available medical checklists, in-  
9           cluding checklists for safe surgery and central line  
10          insertion and maintenance, to inform further med-  
11          ical checklist development.

12          (c) SCOPE.—The Director shall ensure that each as-  
13          pect of the research and study conducted under subsection  
14          (a) is examined across a variety of health care provider  
15          characteristics, medical procedures, patient populations,  
16          and other factors that could affect the use of medical  
17          checklists.

18          (d) DISSEMINATION.—The Director shall make avail-  
19          able to the public the results of the study conducted under  
20          subsection (a) and shall disseminate such results to pa-  
21          tient safety organizations listed pursuant to section 924(d)  
22          of the Public Health Service Act (42 U.S.C. 299b–24(d)).

23          (e) AUTHORIZATION OF APPROPRIATIONS.—There  
24          are authorized to be appropriated to carry out this section

1 such sums as may be necessary for each of fiscal years  
2 2011 through 2014.

3 **SEC. 3. COORDINATING MEDICAL CHECKLISTS AND**  
4 **HEALTH INFORMATION TECHNOLOGY SYS-**  
5 **TEMS.**

6 (a) IN GENERAL.—The HIT Policy Committee in the  
7 Office of the National Coordinator for Health Information  
8 Technology (as established in section 3002 of the Public  
9 Health Service Act (42 U.S.C. 300jj–12) shall develop pol-  
10 icy recommendations regarding—

11 (1) the extent to which the use of medical  
12 checklists should be incorporated into health infor-  
13 mation technology systems; and

14 (2) measures to determine the effectiveness of  
15 such use.

16 (b) AREAS OF CONSIDERATION.—In making rec-  
17 ommendations under subsection (a), the HIT Policy Com-  
18 mittee may consider the following areas:

19 (1) The ease with which medical checklists in  
20 electronic formats can be used by health care profes-  
21 sionals.

22 (2) The effect of the availability of medical  
23 checklists in electronic formats on the adoption and  
24 use of medical checklists by health care profes-  
25 sionals.

1           (3) The effect of the use of medical checklists  
2           in electronic formats on the time spent by health  
3           care professionals on medical procedures.

4           (4) The ability of the health information tech-  
5           nology system to collect data on patient safety and  
6           health outcomes that could be analyzed to aid in the  
7           design and update of medical checklists.

8           (5) The ease with which medical checklists in  
9           electronic formats can be updated on an ongoing  
10          basis based on evidence from medical research and  
11          local experience.

12          (6) The capability of health information tech-  
13          nology systems to collect data, where applicable, re-  
14          garding the use of medical checklists by health care  
15          providers, and any relation between that use and pa-  
16          tient safety and health outcomes.

17 **SEC. 4. INSTITUTE OF MEDICINE STUDY ON FURTHER MED-**  
18 **ICAL CHECKLIST RESEARCH.**

19          (a) IN GENERAL.—The Secretary of Health and  
20          Human Services shall enter into an agreement with the  
21          Institute of Medicine and the National Academy of Engi-  
22          neering of the National Academies to conduct a study in  
23          accordance with this section.

24          (b) STUDY.—The Secretary shall ensure that the  
25          study conducted under this section—

1           (1) reviews available medical checklists and  
2           similar quality improvement techniques, data on the  
3           adoption and use of such techniques by health care  
4           professionals, and evidence of the efficacy of such  
5           techniques in relation to patient safety and health  
6           outcomes;

7           (2) identifies areas of research needed to im-  
8           prove medical checklists in order to increase the  
9           adoption and efficacy of medical checklists;

10          (3) analyzes organizational impediments to the  
11          adoption and use of medical checklists;

12          (4) reviews the degree to which there is suffi-  
13          cient evidence with which to develop new medical  
14          checklists and, if such evidence is insufficient, identi-  
15          fies areas requiring further study in order to develop  
16          such evidence; and

17          (5) determines whether the availability of an in-  
18          creased number of medical checklists would improve  
19          patient safety and health outcomes and, if so, identi-  
20          fies methods for using recent medical research to de-  
21          velop new medical checklists.

22          (c) METHODOLOGY OF STUDY.—

23               (1) SCOPE.—The Secretary shall ensure that  
24               the agreement entered into under subsection (a) pro-

vides that the study conducted under such subsection will consider the perspectives of—

(A) various types of health care professionals in various types of health care settings;

(B) individuals conducting academic research in health care quality; and

(C) patients.

(2) CONSULTATION WITH RELEVANT ORGANIZATIONS.—The Secretary shall ensure that the agreement entered into under subsection (a) provides that relevant agencies and organizations with expertise on medical checklists will be consulted during the study conducted under such subsection, including the following:

(A) The Agency for Healthcare Research and Quality.

(B) The American Nurses Association.

(C) The Institute for Healthcare Improvement.

(D) The American Hospital Association.

(E) The American Medical Association.

(F) The World Health Organization.

(G) The National Committee for Quality Assurance.

(H) The Joint Commission.

1 (I) The American Academy of Physician  
2 Assistants.

3 (d) REPORT.—The Secretary shall ensure that the  
4 agreement entered into under subsection (a) provides that  
5 not later than 18 months after the date of the enactment  
6 of this Act, a report providing the findings and rec-  
7 ommendations made in the study conducted under such  
8 subsection will be submitted to the Secretary, the Com-  
9 mittee on Energy and Commerce of the House of Rep-  
10 resentatives, and the Committee on Health, Education,  
11 Labor, and Pensions of the Senate.

12 **SEC. 5. DEFINITIONS.**

13 In this Act, the following definitions apply:

14 (1) HEALTH CARE PROFESSIONAL.—The term  
15 “health care professional” means an individual who  
16 provides health care services, including a physician,  
17 physician assistant, nurse practitioner, clinical nurse  
18 specialist (as those terms are defined in section  
19 1861 of the Social Security Act (42 U.S.C. 1395x)),  
20 and such other individuals as the Secretary of  
21 Health and Human Services determines appropriate.

22 (2) HEALTH CARE SETTING.—The term “health  
23 care setting” means a facility at which health care  
24 services are provided, including a hospital providing  
25 inpatient hospital services (as that term is defined in



1 section 1861 of the Social Security Act (42 U.S.C.  
2 1395x)), an ambulatory surgical center (meeting the  
3 standards specified under section 1832(a)(2)(F)(i)  
4 of the Social Security Act (42 U.S.C. 1395k)), and  
5 such other facilities as the Secretary of Health and  
6 Human Services determines appropriate.

7 (3) HEALTH CARE PROVIDER.—The term  
8 “health care provider” means a health care profes-  
9 sional or a health care setting.

10 (4) MEDICAL CHECKLIST.—The term “medical  
11 checklist” means a predetermined, evidence-based,  
12 well-defined set of steps that should be completed  
13 during a designated medical clinical encounter or  
14 medical procedure, as further defined by the Direc-  
15 tor of the Agency for Healthcare Research and  
16 Quality in consultation with the Institute of Medi-  
17 cine and the National Academy of Engineering of  
18 the National Academies.

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