115TH CONGRESS  
1ST SESSION  
H. R.  

To amend the Public Health Service Act to foster more effective implementation and coordination of clinical care for people with a complex metabolic or autoimmune disease, a disease resulting from insulin deficiency or insulin resistance, or complications caused by such a disease, and for other purposes.

IN THE HOUSE OF REPRESENTATIVES

Mr. Olson introduced the following bill; which was referred to the Committee on 

A BILL

To amend the Public Health Service Act to foster more effective implementation and coordination of clinical care for people with a complex metabolic or autoimmune disease, a disease resulting from insulin deficiency or insulin resistance, or complications caused by such a disease, 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,
SECTION 1. SHORT TITLE.

This Act may be cited as the “National Clinical Care Commission Act”.

SEC. 2. ESTABLISHMENT OF A NATIONAL CLINICAL CARE COMMISSION.

Part P of title III of the Public Health Service Act (42 U.S.C. 280g et seq.) is amended by adding at the end the following new section:

“SEC. 399V–7. NATIONAL CLINICAL CARE COMMISSION.

“(a) ESTABLISHMENT.—There is hereby established, within the Department of Health and Human Services, a National Clinical Care Commission (in this section referred to as the ‘Commission’) to evaluate, and recommend solutions regarding better coordination and leveraging of, programs within the Department and other Federal agencies that relate in any way to supporting appropriate clinical care (such as any interactions between physicians and other health care providers and their patients related to treatment and care management) for individuals with—

“(1) one or more complex metabolic or autoimmune diseases;

“(2) one or more diseases resulting from insulin deficiency or insulin resistance; or

“(3) complications caused by one or more of any of such diseases.
“(b) Membership.—

“(1) IN GENERAL.—The Commission shall be composed of the following voting members:

“(A) The heads (or their designees) of the following Federal agencies and departments:


“(iii) The Centers for Disease Control and Prevention.

“(iv) The Indian Health Service.

“(v) The Department of Veterans Affairs.

“(vi) The National Institutes of Health.

“(vii) The Food and Drug Administration.

“(viii) The Health Resources and Services Administration.

“(ix) The Department of Defense.

“(B) Twelve additional voting members appointed under paragraph (2).

“(C) Such additional voting members as may be appointed by the Secretary, at the Sec-
Secretary’s discretion, from among the heads (or their designees) of governmental or nongovernmental entities that impact clinical care of individuals with any of the diseases and complications described in subsection (a).

“(2) ADDITIONAL MEMBERS.—The Commission shall include additional voting members appointed by the Secretary, in consultation with national medical societies and patient advocacy organizations with expertise in the care and epidemiology of any of the diseases and complications described in subsection (a), including one or more such members from each of the following categories:

“(A) Clinical endocrinologists.

“(B) Physician specialties (other than as described in subparagraph (A)) that play a role in diseases and complications described in subsection (a), such as cardiologists, nephrologists, and eye care professionals.

“(C) Primary care physicians.

“(D) Non-physician health care professionals, such as certified diabetes educators, registered dieticians and nutrition professionals, nurses, nurse practitioners, physician assistants.
“(E) Patient advocates.

“(F) National experts in the duties listed under subsection (c).

“(G) Health care providers furnishing services to a patient population that consists of a high percentage (as specified by the Secretary) of individuals who are enrolled in a State plan under title XIX of the Social Security Act or who are not covered under a health plan or health insurance coverage.

“(3) CHAIRPERSON.—The voting members of the Commission shall select a chairperson from the members appointed under paragraph (2) from the category under paragraph (2)(A).

“(4) MEETINGS.—The Commission shall meet at least twice, and not more than 4 times, a year.

“(5) BOARD TERMS.—Members of the Commission appointed pursuant to subparagraph (B) or (C) of paragraph (1), including the chairperson, shall serve for a 3-year term. A vacancy on the Commission shall be filled in the same manner as the original appointments.

“(e) DUTIES.—The Commission shall—

“(1) evaluate programs of the Department of Health and Human Services regarding the utiliza-
tion of diabetes screening benefits, annual wellness
visits, and other preventive health benefits that may
reduce the incidence of the diseases and complica-
tions described in subsection (a), including identi-
fying problems regarding such utilization and related
data collection mechanisms and make recommenda-
tions;

“(2) identify current activities and critical gaps
in Federal efforts to support clinicians in providing
integrated, high-quality care to individuals with any
of the diseases and complications described in sub-
section (a);

“(3) make recommendations regarding the co-
ordination of clinically-based activities that are being
supported by the Federal Government with respect
to the diseases and complications described in sub-
section (a);

“(4) make recommendations regarding the de-
development and coordination of federally funded clin-
ical practice support tools for physicians and other
health care professionals in caring for and managing
the care of individuals with any of the diseases and
complications described in subsection (a), specifically
with regard to implementation of new treatments
and technologies;
“(5) evaluate programs described in subsection (a) that are in existence as of the date of the enactment of this section and determine if such programs are meeting the needs identified in paragraph (2) and, if such programs are determined as not meeting such needs, recommend programs that would be more appropriate;

“(6) recommend, with respect to the diseases and complications described in subsection (a), clinical pathways for new technologies and treatments, including future data collection activities, that may be developed and then used to evaluate—

“(A) various care models and methods;

and

“(B) the impact of such models and methods on quality of care as measured by appropriate care parameters (such as A1C, blood pressure, and cholesterol levels);

“(7) evaluate and expand education and awareness activities provided to physicians and other health care professionals regarding clinical practices for the prevention and treatment of the diseases and complications described in subsection (a);
“(8) review and recommend appropriate methods for outreach and dissemination of educational resources that—

“(A) address the diseases and complications described in subsection (a);

“(B) are funded by the Federal Government; and

“(C) are intended for health care professionals and the public; and

“(9) carry out other activities, such as activities relating to the areas of public health and nutrition, that the Commission deems appropriate with respect to the diseases and complications described in subsection (a).

“(d) OPERATING PLAN.—

“(1) INITIAL PLAN.—Not later than 90 days after its first meeting, the Commission shall submit to the Secretary and the Congress an operating plan for carrying out the activities of the Commission as described in subsection (c). Such operating plan may include—

“(A) a list of specific activities that the Commission plans to conduct for purposes of carrying out the duties described in each of the paragraphs in subsection (c);
“(B) a plan for completing the activities;

“(C) a list of members of the Commission and other individuals who are not members of the Commission who will need to be involved to conduct such activities;

“(D) an explanation of Federal agency involvement and coordination needed to conduct such activities;

“(E) a budget for conducting such activities;

“(F) a plan for evaluating the value and potential impact of the Commission’s work and recommendations, including the possible continuation of the Commission for the purposes of overseeing their implementation; and

“(G) other information that the Commission deems appropriate.

“(2) UPDATES.—The Commission shall periodically update the operating plan under paragraph (1) and submit such updates to the Secretary and the Congress.

“(e) FINAL REPORT.—By not later than 3 years after the date of the Commission’s first meeting, the Commission shall submit to the Secretary and the Congress a final report containing all of the findings and recommendations
required by this section. Not later than 120 days after
the submission of the final report, the Secretary shall re-
view the plan required by subsection (d)(1)(F) and submit
to the Congress a recommendation on whether the Com-
mission should be reauthorized to operate after fiscal year
2021.

“(f) SUNSET.—The Commission shall terminate 120
days after submitting its final report, but not later than
the end of fiscal year 2021.”.