

Editorial Notes

AMENDMENTS

1993—Subsec. (c). Pub. L. 103-43 struck out subsec. (c) which read as follows: “For the purpose of performing the duties specified in subsection (b) of this section, there are authorized to be appropriated \$8,000,000 for fiscal year 1989 and such sums as may be necessary for fiscal year 1990. Funds appropriated under this subsection shall remain available until expended.”

SUBPART 4—NATIONAL INFORMATION CENTER ON HEALTH SERVICES RESEARCH AND HEALTH CARE TECHNOLOGY

§ 286d. National Information Center**(a) Establishment**

There is established within the Library an entity to be known as the National Information Center on Health Services Research and Health Care Technology (in this section referred to as the “Center”).

(b) Purpose

The purpose of the Center is the collection, storage, analysis, retrieval, and dissemination of information on health services research, clinical practice guidelines, and on health care technology, including the assessment of such technology. Such purpose includes developing and maintaining data bases and developing and implementing methods of carrying out such purpose.

(c) Electronic, convenient format; criteria for inclusion

The Director of the Center shall ensure that information under subsection (b) concerning clinical practice guidelines is collected and maintained electronically and in a convenient format. Such Director shall develop and publish criteria for the inclusion of practice guidelines and technology assessments in the information center database.

(d) Coordination with Director of the Agency for Healthcare Research and Quality

The Secretary, acting through the Center, shall coordinate the activities carried out under this section through the Center with related activities of the Director of the Agency for Healthcare Research and Quality.

(July 1, 1944, ch. 373, title IV, § 478A, as added Pub. L. 103-43, title XIV, § 1421, June 10, 1993, 107 Stat. 171; amended Pub. L. 106-129, § 2(b)(2), Dec. 6, 1999, 113 Stat. 1670.)

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1999—Subsec. (d). Pub. L. 106-129 substituted “Director of the Agency for Healthcare Research and Quality” for “Administrator for Health Care Policy and Research”.

Statutory Notes and Related Subsidiaries

CONSTRUCTION

Pub. L. 103-43, § 1422(b), June 10, 1993, 107 Stat. 172, provided that: “The amendments made by section 3 of Public Law 102-410 (106 Stat. 2094) [amending section 299a-1 of this title], by section 1421 of this Act [enacting this section], and by subsection (a) of this section

[amending section 299a-1 of this title] may not be construed as terminating the information center on health care technologies and health care technology assessment established under section 904 of the Public Health Service Act [42 U.S.C. 299a-2], as in effect on the day before the date of the enactment of Public Law 102-410 [Oct. 13, 1992]. Such center shall be considered to be the center established in section 478A of the Public Health Service Act [42 U.S.C. 286d], as added by section 1421 of this Act, and shall be subject to the provisions of such section 478A.”

PART E—OTHER AGENCIES OF NIH

SUBPART 1—NATIONAL CENTER FOR ADVANCING TRANSLATIONAL SCIENCES

Editorial Notes

CODIFICATION

Pub. L. 112-74, div. F, title II, § 221(a)(1)(A), Dec. 23, 2011, 125 Stat. 1086, substituted “advancing translational sciences” for “research resources” in subpart heading.

§ 287. National Center for Advancing Translational Sciences**(a) Purpose**

The purpose of the National Center for Advancing Translational Sciences (in this subpart referred to as the “Center”) is to advance translational sciences, including—

(1) coordinating and developing resources that leverage basic research in support of translational science; and

(2) developing partnerships and working cooperatively to foster synergy in ways that do not create duplication, redundancy, and competition with industry activities.

(b) Clinical trial activities**(1) In general**

The Center may develop and provide infrastructure and resources for all phases of clinical trials research. Except as provided in paragraph (2), the Center may support clinical trials only through the end of phase IIB.

(2) Exception

The Center may support clinical trial activities through the end of phase III for a treatment for a rare disease or condition (as defined in section 360bb of title 21) so long as—

(A) the Center gives public notice for a period of at least 120 days of the Center’s intention to support the clinical trial activities in phase III;

(B) no public or private organization provides credible written intent to the Center that the organization has timely plans to further the clinical trial activities or conduct clinical trials of a similar nature beyond phase IIB; and

(C) the Center ensures that support of the clinical trial activities in phase III will not increase the Federal Government’s liability beyond the award value of the Center’s support.

(c) Biennial report

The Center shall publish a report on a biennial basis that, with respect to all research supported by the Center, includes a complete list of—