beths Hospital, prior to repeal by Pub. L. 98–621, § 10(s), Nov. 8, 1984, 98 Stat. 3381, effective Oct. 1, 1987. Subsequent to repeal, section 2104 of title XXI of act July 1, 1944, was renumbered section 2304 of title XXIII of that act by section 311(a) of Pub. L. 99–660.

A prior section 300cc–4, acts July 1, 1944, ch. 373, title XXI, § 2105, formerly title V, § 505, 58 Stat. 710; 1953 Reorg. Plan No. 1, §§ 5, 8, eff. Apr. 11, 1953, 18 F.R. 2053, 67 Stat. 631; renumbered title XXI, § 2105, Apr. 26, 1983, Pub. L. 98–24, § 2(a)(1), 97 Stat. 176, provided procedures under which the Secretary could settle claims for damages from collisions or incident to the operation of vessels within a year of the accrual of such claims and not to exceed $3,000, prior to repeal by Pub. L. 99–117, § 12(f), Oct. 7, 1985, 99 Stat. 495. Subsequent to repeal, section 2105 of title XXI of act July 1, 1944, was renumbered section 2305 of title XXIII of that act by Pub. L. 99–660, title III, § 311(a), Nov. 14, 1986, 100 Stat. 3755.

Prior sections 300cc–5 to 300cc–10, act July 1, 1944, §§ 2306 to 2311, respectively, were successively renumbered by subsequent acts and transferred, see sections 238c to 238h of this title.

AMENDMENTS

1993—Pub. L. 103–43, § 2008(d)(1)(A), substituted ''Research Advisory Committee'' for ''Clinical Research Review Committee'' in section catchline.

Subsec. (a). Pub. L. 103–43, § 2008(d)(1)(B), substituted ''AIDS Research Advisory Committee'' for ''AIDS Clinical Research Review Committee''.

Subsec. (c)(1). Pub. L. 103–43, § 1811(1), in introductory provisions inserted ''(and may provide advice to the Directors of other agencies of the National Institutes of Health, as appropriate)'' after ''Director of such Institute'' and in subpar. (A) inserted before semicolon at end '', including recommendations on the projects of research with respect to diagnosing immune deficiency and with respect to predicting, diagnosing, preventing, and treating opportunistic cancers and infectious diseases''.

1988—Subsec. (c)(2)(B). Pub. L. 100–690 substituted semicolon for period.

EFFECTIVE DATE OF 1988 AMENDMENT

Amendment by Pub. L. 100–690 effective immediately after enactment of Pub. L. 100–607, which was approved Nov. 4, 1988, see section 2600 of Pub. L. 100–690, set out as a note under section 242m of this title.

TERMINATION OF ADVISORY COMMITTEES

Advisory committees established after Jan. 5, 1973, to terminate not later than the expiration of the 2-year period beginning on the date of their establishment, unless, in the case of a committee established by the President or an officer of the Federal Government, such committee is renewed by appropriate action prior to the expiration of such 2-year period, or in the case of a committee established by the Congress, its duration is otherwise provided by law. See section 14 of Pub. L. 92–463, Oct. 6, 1972, 86 Stat. 776, set out in the Appendix to Title 5, Government Organization and Employees.

Pub. L. 93–641, § 6, Jan. 4, 1975, 88 Stat. 2275, set out as a note under section 217a of this title, provided that an advisory committee established pursuant to the Public Health Service Act shall terminate at such time as may be specifically prescribed by an Act of Congress enacted after Jan. 4, 1975.

PART B—RESEARCH AUTHORITY

§ 300cc–11. Clinical evaluation units at National Institutes of Health

(a) In general

The Secretary, acting through the Director of the National Cancer Institute and the Director of the National Institute of Allergy and Infectious Diseases, shall for each such Institute establish a clinical evaluation unit at the Clinical Center at the National Institutes of Health. Each of the clinical evaluation units—

(1) shall conduct clinical evaluations of experimental treatments for acquired immune deficiency syndrome developed within the preclinical drug development program, including evaluations of methods of diagnosing immune deficiency and evaluations of methods of predicting, diagnosing, preventing, and treating opportunistic cancers and infectious diseases; and

(2) may conduct clinical evaluations of experimental treatments for such syndrome that are developed by any other national research institute of the National Institutes of Health or by any other entity.

(b) Personnel and administrative support

(1) For the purposes described in subsection (a) of this section, the Secretary, acting through the Director of the National Institutes of Health, shall provide each of the clinical evaluation units required in such subsection—

(A)(i) with not less than 50 beds; or

(ii) with an outpatient clinical capacity equal to not less than twice the outpatient clinical capacity, with respect to acquired immune deficiency syndrome, possessed by the Clinical Center of the National Institutes of Health on June 1, 1988; and

(B) with such personnel, such administrative support, and such other support services as may be necessary.

(2) Facilities, personnel, administrative support, and other support services provided pursuant to paragraph (1) shall be in addition to the number or level of facilities, personnel, administrative support, and other support services that otherwise would be available at the Clinical Center at the National Institutes of Health for the provision of clinical care for individuals with diseases or disorders.

(c) Authorization of appropriations

For the purpose of carrying out this section, there are authorized to be appropriated such sums as may be necessary.

(July 1, 1944, ch. 373, title XXIII, § 2311, as added Pub. L. 100–607, title II, § 201(4), Nov. 4, 1988, 102 Stat. 3066; amended Pub. L. 103–43, title XVIII, § 1811(2), June 10, 1993,107 Stat. 199.)

PRIOR PROVISIONS

A prior section 300cc–11, act July 1, 1944, § 2312, was successively renumbered by subsequent acts and transferred, see section 238i of this title.

AMENDMENTS

1993—Subsec. (a)(1). Pub. L. 103–43 inserted before semicolon at end '', including evaluations of methods of diagnosing immune deficiency and evaluations of methods of predicting, diagnosing, preventing, and treating opportunistic cancers and infectious diseases''.

§ 300cc–12. Use of investigational new drugs with respect to acquired immune deficiency syndrome

(a) Encouragement of applications with respect to clinical trials

(1) If, in the determination of the Secretary, there is preliminary evidence that a new drug