

(July 1, 1944, ch. 373, title III, §377A, as added Pub. L. 108-216, §4, Apr. 5, 2004, 118 Stat. 585.)

§ 274f-2. Grants regarding hospital organ donation coordinators

(a) Authority

(1) In general

The Secretary may award grants to qualified organ procurement organizations and hospitals under section 273 of this title to establish programs coordinating organ donation activities of eligible hospitals and qualified organ procurement organizations under section 273 of this title. Such activities shall be coordinated to increase the rate of organ donations for such hospitals.

(2) Eligible hospital

For purposes of this section, the term “eligible hospital” means a hospital that performs significant trauma care, or a hospital or consortium of hospitals that serves a population base of not fewer than 200,000 individuals.

(b) Administration of coordination program

A condition for the receipt of a grant under subsection (a) of this section is that the applicant involved agree that the program under such subsection will be carried out jointly—

(1) by representatives from the eligible hospital and the qualified organ procurement organization with respect to which the grant is made; and

(2) by such other entities as the representatives referred to in paragraph (1) may designate.

(c) Requirements

Each entity receiving a grant under subsection (a) of this section shall—

(1) establish joint organ procurement organization and hospital designated leadership responsibility and accountability for the project;

(2) develop mutually agreed upon overall project performance goals and outcome measures, including interim outcome targets; and

(3) collaboratively design and implement an appropriate data collection process to provide ongoing feedback to hospital and organ procurement organization leadership on project progress and results.

(d) Rule of construction

Nothing in this section shall be construed to interfere with regulations in force on April 5, 2004.

(e) Evaluations

Within 3 years after the award of grants under this section, the Secretary shall ensure an evaluation of programs carried out pursuant to subsection (a) of this section in order to determine the extent to which the programs have increased the rate of organ donation for the eligible hospitals involved.

(f) Matching requirement

The Secretary may not award a grant to a qualifying organ donation entity under this section unless such entity agrees that, with respect to costs to be incurred by the entity in carrying

out activities for which the grant was awarded, the entity shall contribute (directly or through donations from public or private entities) non-Federal contributions in cash or in kind, in an amount equal to not less than 30 percent of the amount of the grant awarded to such entity.

(g) Funding

For the purpose of carrying out this section, there are authorized to be appropriated \$3,000,000 for fiscal year 2005, and such sums as may be necessary for each of fiscal years 2006 through 2009.

(July 1, 1944, ch. 373, title III, §377B, as added Pub. L. 108-216, §4, Apr. 5, 2004, 118 Stat. 586.)

§ 274f-3. Studies relating to organ donation and the recovery, preservation, and transportation of organs

(a) Development of supportive information

The Secretary, acting through the Director of the Agency for Healthcare Research and Quality, shall develop scientific evidence in support of efforts to increase organ donation and improve the recovery, preservation, and transportation of organs.

(b) Activities

In carrying out subsection (a) of this section, the Secretary shall—

(1) conduct or support evaluation research to determine whether interventions, technologies, or other activities improve the effectiveness, efficiency, or quality of existing organ donation practice;

(2) undertake or support periodic reviews of the scientific literature to assist efforts of professional societies to ensure that the clinical practice guidelines that they develop reflect the latest scientific findings;

(3) ensure that scientific evidence of the research and other activities undertaken under this section is readily accessible by the organ procurement workforce; and

(4) work in coordination with the appropriate professional societies as well as the Organ Procurement and Transplantation Network and other organ procurement and transplantation organizations to develop evidence and promote the adoption of such proven practices.

(c) Research and dissemination

The Secretary, acting through the Director of the Agency for Healthcare Research and Quality, as appropriate, shall provide support for research and dissemination of findings, to—

(1) develop a uniform clinical vocabulary for organ recovery;

(2) apply information technology and telecommunications to support the clinical operations of organ procurement organizations;

(3) enhance the skill levels of the organ procurement workforce in undertaking quality improvement activities; and

(4) assess specific organ recovery, preservation, and transportation technologies.

(d) Authorization of appropriations

For the purpose of carrying out this section, there are authorized to be appropriated \$2,000,000

for fiscal year 2005, and such sums as may be necessary for each of fiscal years 2006 through 2009.

(July 1, 1944, ch. 373, title III, §377C, as added Pub. L. 108-216, §5, Apr. 5, 2004, 118 Stat. 587.)

§ 274f-4. Report relating to organ donation and the recovery, preservation, and transportation of organs

(a) In general

Not later than December 31, 2005, and every 2 years thereafter, the Secretary shall report to the appropriate committees of Congress on the activities of the Department carried out pursuant to this part, including an evaluation describing the extent to which the activities have affected the rate of organ donation and recovery.

(b) Requirements

To the extent practicable, each report submitted under subsection (a) of this section shall—

- (1) evaluate the effectiveness of activities, identify effective activities, and disseminate such findings with respect to organ donation and recovery;
- (2) assess organ donation and recovery activities that are recently completed, ongoing, or planned; and
- (3) evaluate progress on the implementation of the plan required under subsection (c)(5) of this section.

(c) Initial report requirements

The initial report under subsection (a) of this section shall include the following:

- (1) An evaluation of the organ donation practices of organ procurement organizations, States, other countries, and other appropriate organizations including an examination across all populations, including those with low organ donation rates, of—

- (A) existing barriers to organ donation; and
- (B) the most effective donation and recovery practices.

- (2) An evaluation of living donation practices and procedures. Such evaluation shall include an assessment of issues relating to informed consent and the health risks associated with living donation (including possible reduction of long-term effects).

- (3) An evaluation of—

- (A) federally supported or conducted organ donation efforts and policies, as well as federally supported or conducted basic, clinical, and health services research (including research on preservation techniques and organ rejection and compatibility); and

- (B) the coordination of such efforts across relevant agencies within the Department and throughout the Federal Government.

- (4) An evaluation of the costs and benefits of State donor registries, including the status of existing State donor registries, the effect of State donor registries on organ donation rates, issues relating to consent, and recommendations regarding improving the effectiveness of State donor registries in increasing overall organ donation rates.

(5) A plan to improve federally supported or conducted organ donation and recovery activities, including, when appropriate, the establishment of baselines and benchmarks to measure overall outcomes of these programs. Such plan shall provide for the ongoing coordination of federally supported or conducted organ donation and research activities.

(July 1, 1944, ch. 373, title III, §377D, as added Pub. L. 108-216, §6, Apr. 5, 2004, 118 Stat. 588.)

§ 274g. Authorization of appropriations

For the purpose of carrying out this part, there are authorized to be appropriated \$8,000,000 for fiscal year 1991, and such sums as may be necessary for each of the fiscal years 1992 and 1993.

(July 1, 1944, ch. 373, title III, §378, as added Pub. L. 101-616, title II, §206(a), Nov. 16, 1990, 104 Stat. 3285; amended Pub. L. 105-196, §4(1), July 16, 1998, 112 Stat. 636.)

AMENDMENTS

1998—Pub. L. 105-196 made technical amendment relating to placement of section within part H of this subchapter.

PART I—C.W. BILL YOUNG CELL TRANSPLANTATION PROGRAM

AMENDMENTS

2005—Pub. L. 109-129, §3(e), Dec. 20, 2005, 119 Stat. 2562, substituted “C.W. Bill Young Cell Transplantation Program” for “National Bone Marrow Donor Registry” in part heading.

1990—Pub. L. 101-616, title I, §101(a)(2), Nov. 16, 1990, 104 Stat. 3279, added part I “National Bone Marrow Donor Registry” and redesignated former part I “Biomedical Ethics” as J.

1985—Pub. L. 99-158, §§3(b), 11, Nov. 20, 1985, 99 Stat. 879, 883, added part I “Biomedical Ethics”, and repealed former part I “National Library of Medicine”.

1970—Pub. L. 91-212, §10(a)(2), Mar. 13, 1970, 84 Stat. 66, redesignated part H “National Library of Medicine”, as part I “National Library of Medicine”.

§ 274k. National Program

(a) Establishment

The Secretary, acting through the Administrator of the Health Resources and Services Administration, shall by one or more contracts establish and maintain a C.W. Bill Young Cell Transplantation Program (referred to in this section as the “Program”), successor to the National Bone Marrow Donor Registry, that has the purpose of increasing the number of transplants for recipients suitably matched to biologically unrelated donors of bone marrow and cord blood, and that meets the requirements of this section. The Secretary may award a separate contract to perform each of the major functions of the Program described in paragraphs (1) and (2) of subsection (d) of this section if deemed necessary by the Secretary to operate an effective and efficient system that is in the best interest of patients. The Secretary shall conduct a separate competition for the initial establishment of the cord blood functions of the Program. The Program shall be under the general supervision of the Secretary. The Secretary shall establish an Advisory Council to advise,