

stance Use Policy Laboratory established under section 290aa-0 of this title, or the Assistant Secretary for Planning and Evaluation, including pursuant to the evaluation and recommendations under section 6021 of the Helping Families in Mental Health Crisis Reform Act of 2016 or priorities identified in the strategic plan under section 290aa(7) of this title.

**(c) Requirements**

The Assistant Secretary may establish minimum requirements for the applications submitted under subsection (b), including applications related to the submission of research and evaluation.

**(d) Review and rating**

**(1) In general**

The Assistant Secretary shall review applications prior to public posting in accordance with subsection (a), and may prioritize the review of applications for evidence-based programs and practices that are related to topics included in the notice provided under subsection (b)(2).

**(2) System**

In carrying out paragraph (1), the Assistant Secretary may utilize a rating and review system, which may include information on the strength of evidence associated with the evidence-based programs and practices and a rating of the methodological rigor of the research supporting the applications.

**(3) Public access to metrics and rating**

The Assistant Secretary shall make the metrics used to evaluate applications under this section, and any resulting ratings of such applications, publicly available.

(July 1, 1944, ch. 373, title V, § 543A, as added Pub. L. 114-255, div. B, title VII, § 7002, Dec. 13, 2016, 130 Stat. 1222.)

REFERENCES IN TEXT

Section 6021 of the Helping Families in Mental Health Crisis Reform Act of 2016, referred to in subsec. (b)(2), is section 6021 of Pub. L. 114-255, which is set out as a note under section 290aa of this title.

**§ 290dd-3. Grants for reducing overdose deaths**

**(a) Establishment**

**(1) In general**

The Secretary shall award grants to eligible entities to expand access to drugs or devices approved or cleared under the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.] for emergency treatment of known or suspected opioid overdose.

**(2) Maximum grant amount**

A grant awarded under this section may not be for more than \$200,000 per grant year.

**(3) Eligible entity**

For purposes of this section, the term “eligible entity” means a Federally qualified health center (as defined in section 1395x(aa) of this title), an opioid treatment program under part 8 of title 42, Code of Federal Regulations, any practitioner dispensing narcotic drugs pursu-

ant to section 823(g) of title 21, or any other entity that the Secretary deems appropriate.

**(4) Prescribing**

For purposes of this section, the term “prescribing” means, with respect to a drug or device approved or cleared under the Federal Food, Drug, and Cosmetic Act for emergency treatment of known or suspected opioid overdose, the practice of prescribing such drug or device—

(A) in conjunction with an opioid prescription for patients at an elevated risk of overdose;

(B) in conjunction with an opioid agonist approved under section 505 of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 355] for the treatment of opioid use disorder;

(C) to the caregiver or a close relative of patients at an elevated risk of overdose from opioids; or

(D) in other circumstances in which a provider identifies a patient is at an elevated risk for an intentional or unintentional drug overdose from heroin or prescription opioid therapies.

**(b) Application**

To be eligible to receive a grant under this section, an eligible entity shall submit to the Secretary, in such form and manner as specified by the Secretary, an application that describes—

(1) the extent to which the area to which the entity will furnish services through use of the grant is experiencing significant morbidity and mortality caused by opioid abuse;

(2) the criteria that will be used to identify eligible patients to participate in such program; and

(3) a plan for sustaining the program after Federal support for the program has ended.

**(c) Use of funds**

An eligible entity receiving a grant under this section may use amounts under the grant for any of the following activities, but may use not more than 20 percent of the grant funds for activities described in paragraphs (3) and (4):

(1) To establish a program for prescribing a drug or device approved or cleared under the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.] for emergency treatment of known or suspected opioid overdose.

(2) To train and provide resources for health care providers and pharmacists on the prescribing of drugs or devices approved or cleared under the Federal Food, Drug, and Cosmetic Act for emergency treatment of known or suspected opioid overdose.

(3) To purchase drugs or devices approved or cleared under the Federal Food, Drug, and Cosmetic Act for emergency treatment of known or suspected opioid overdose, for distribution under the program described in paragraph (1).

(4) To offset the co-payments and other cost sharing associated with drugs or devices approved or cleared under the Federal Food, Drug, and Cosmetic Act for emergency treatment of known or suspected opioid overdose.

(5) To establish protocols to connect patients who have experienced a drug overdose

with appropriate treatment, including medication-assisted treatment and appropriate counseling and behavioral therapies.

**(d) Evaluations by recipients**

As a condition of receipt of a grant under this section, an eligible entity shall, for each year for which the grant is received, submit to the Secretary an evaluation of activities funded by the grant which contains such information as the Secretary may reasonably require.

**(e) Reports by the Secretary**

Not later than 5 years after the date on which the first grant under this section is awarded, the Secretary shall submit to the appropriate committees of the House of Representatives and of the Senate a report aggregating the information received from the grant recipients for such year under subsection (d) and evaluating the outcomes achieved by the programs funded by grants awarded under this section.

**(f) Authorization of appropriations**

There is authorized to be appropriated to carry out this section, \$5,000,000 for the period of fiscal years 2017 through 2021.

(July 1, 1944, ch. 373, title V, §544, as added Pub. L. 114-198, title I, §107(a), July 22, 2016, 130 Stat. 703.)

REFERENCES IN TEXT

The Federal Food, Drug, and Cosmetic Act, referred to in subssecs. (a)(1) and (c)(1) to (4), is act June 25, 1938, ch. 675, 52 Stat. 1040, which is classified generally to chapter 9 (§301 et seq.) of Title 21, Food and Drugs. For complete classification of this Act to the Code, see section 301 of Title 21 and Tables.

PRIOR PROVISIONS

A prior section 290dd-3, act July 1, 1944, ch. 373, title V, §544, formerly Pub. L. 91-616, title III, §333, Dec. 31, 1970, 84 Stat. 1853, as amended Pub. L. 93-282, title I, §122(a), May 14, 1974, 88 Stat. 131; Pub. L. 94-581, title I, §111(c)(4), Oct. 21, 1976, 90 Stat. 2852; renumbered §523 of act July 1, 1944, Apr. 26, 1983, Pub. L. 98-24, §2(b)(13), 97 Stat. 181; Aug. 27, 1986, Pub. L. 99-401, title I, §106(a), 100 Stat. 907; renumbered §544, July 22, 1987, Pub. L. 100-77, title VI, §611(2), 101 Stat. 516; June 13, 1991, Pub. L. 102-54, §13(q)(1)(A)(ii), 105 Stat. 278, which related to confidentiality of patient records for alcohol abuse and alcoholism programs, was omitted in the general revision of this part by Pub. L. 102-321. See section 290dd-2 of this title.

IMPROVING ACCESS TO OVERDOSE TREATMENT

Pub. L. 114-198, title I, §107(b), July 22, 2016, 130 Stat. 705, provided that:

“(1) INFORMATION ON BEST PRACTICES.—Not later than 180 days after the date of enactment of this Act [July 22, 2016]:

“(A) The Secretary of Health and Human Services may provide information to prescribers within Federally qualified health centers (as defined in paragraph (4) of section 1861(aa) of the Social Security Act (42 U.S.C. 1395x(aa))), and the health care facilities of the Indian Health Service, on best practices for prescribing or co-prescribing a drug or device approved or cleared under the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.) for emergency treatment of known or suspected opioid overdose, including for patients receiving chronic opioid therapy and patients being treated for opioid use disorders.

“(B) The Secretary of Defense may provide information to prescribers within Department of Defense medical facilities on best practices for prescribing or

co-prescribing a drug or device approved or cleared under the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.) for emergency treatment of known or suspected opioid overdose, including for patients receiving chronic opioid therapy and patients being treated for opioid use disorders.

“(C) The Secretary of Veterans Affairs may provide information to prescribers within Department of Veterans Affairs medical facilities on best practices for prescribing or co-prescribing a drug or device approved or cleared under the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.) for emergency treatment of known or suspected opioid overdose, including for patients receiving chronic opioid therapy and patients being treated for opioid use disorders.

“(2) RULE OF CONSTRUCTION.—Nothing in this subsection should be construed to establish or contribute to a medical standard of care.”

**§ 290dd-4. Program to support coordination and continuation of care for drug overdose patients**

**(a) In general**

The Secretary of Health and Human Services (referred to in this section as the “Secretary”) shall identify or facilitate the development of best practices for—

(1) emergency treatment of known or suspected drug overdose;

(2) the use of recovery coaches, as appropriate, to encourage individuals who experience a non-fatal overdose to seek treatment for substance use disorder and to support coordination and continuation of care;

(3) coordination and continuation of care and treatment, including, as appropriate, through referrals, of individuals after a drug overdose; and

(4) the provision or prescribing of overdose reversal medication, as appropriate.

**(b) Grant establishment and participation**

**(1) In general**

The Secretary shall award grants on a competitive basis to eligible entities to support implementation of voluntary programs for care and treatment of individuals after a drug overdose, as appropriate, which may include implementation of the best practices described in subsection (a).

**(2) Eligible entity**

In this section, the term “eligible entity” means—

(A) a State substance abuse agency;

(B) an Indian Tribe or tribal organization; or

(C) an entity that offers treatment or other services for individuals in response to, or following, drug overdoses or a drug overdose, such as an emergency department, in consultation with a State substance abuse agency.

**(3) Application**

An eligible entity desiring a grant under this section shall submit an application to the Secretary, at such time and in such manner as the Secretary may require, that includes—

(A) evidence that such eligible entity carries out, or is capable of contracting and coordinating with other community entities to carry out, the activities described in paragraph (4);