

A BILL

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IN THE COUNCIL OF THE DISTRICT OF COLUMBIA

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To amend the Uniform Controlled Substances Act of 1981 to limit initial prescription of opiates for acute conditions to seven days or less, create additional requirements for subsequent prescription of opiates for acute conditions, limit the strength of opiates for other than acute conditions, and require the Mayor to issue rules to limit dispensing of opiates prescribed out-of-State and track the number of prescriptions prescribed by practitioners.

BE IT ENACTED BY THE COUNCIL OF THE DISTRICT OF COLUMBIA, That this act may be cited as the "Opioid Abuse Prevention Amendment Act of 2017".

Sec. 2. The Uniform Controlled Substances Act of 1981, effective August 5, 1981 (D.C. Law 4-29; D.C. Official Code § 48-901.01 *et seq.*), is amended as follows:

(a) Section 102 (D.C. Official Code § 48-901.02) is amended as follows:

(1) A new paragraph 1A is added to read as follows:

"(1A) "Acute pain" means pain, whether resulting from disease, accidental or intentional trauma, or other cause, that the practitioner reasonably expects to last only a short period of time, but does not include chronic pain, pain being treated as part of cancer care, hospice or other end of life care, or pain being treated as part of palliative care.

(2) A new paragraph 12A is added to read as follows:

37                   “(12A) “Initial prescription” means a prescription issued to a patient who, as  
38 determined by the practitioner after consultation with the patient:

39                   “(A) Has never previously been issued a prescription for the drug or its  
40 pharmaceutical equivalent; or

41                   “(B) Was previously issued a prescription for the drug or its  
42 pharmaceutical equivalent, but the date on which the current prescription is being issued is more  
43 than one year after the date the patient last used or was administered the drug or its equivalent.

44           (b) Section 308 (D.C. Official Code § 48-903.08) is amended by adding a new paragraph  
45 (f) to read as follows:

46           “(f)(1) A practitioner shall not issue an initial prescription for an opiate in a quantity  
47 exceeding a seven-day supply, with no refill, for treatment of acute pain.

48           “(2) No less than six days after issuing the initial prescription pursuant to  
49 paragraph (1) of this subsection, the practitioner, after consultation with the patient, may issue a  
50 subsequent prescription for the drug to the patient in any quantity that complies with applicable  
51 laws, provided that:

52                   “(A) The subsequent prescription would not be deemed an initial  
53 prescription under this section;

54                   “(B) The practitioner determines the prescription is necessary and  
55 appropriate to the patient’s treatment needs and documents the rationale for the issuance of the  
56 subsequent prescription; and

57                   “(C) The practitioner determines that issuance of the subsequent  
58 prescription does not present an undue risk of abuse, addiction, or diversion and documents that  
59 determination.

60 “(3) Prior to issuing a second prescription of a Schedule II opiate in a course of  
61 treatment for acute pain, a practitioner shall:

62 “(A) Take and document the results of a thorough medical history,  
63 including the patient’s experience with non-opioid medication and non-pharmacological pain  
64 management approaches and substance abuse history;

65 “(B) Conduct, as appropriate, and document the results of a physical  
66 examination; and

67 “(C) Develop a treatment plan, with particular attention focused on  
68 determining the cause of the patient’s pain.

69 “(4) At the time of the issuance of a prescription after the second prescription for  
70 an opiate for acute pain, or when a Schedule II opiate is continuously prescribed for three months  
71 or more for chronic pain, the practitioner shall:

72 “(A) Review, at a minimum of every three months, the course of  
73 treatment, any new information about the etiology of the pain, and the patient’s progress toward  
74 treatment objectives and document the results of that review;

75 “(B) Assess the patient prior to every renewal to determine whether the  
76 patient is experiencing problems associated with physical and psychological dependence and  
77 document the results of that assessment;

78 “(C) Periodically make reasonable efforts, unless clinically  
79 contraindicated, to either stop the use of the controlled substance, decrease the dosage, try other  
80 drugs or treatment modalities in an effort to reduce the potential for abuse or the development of  
81 physical or psychological dependence and document with specificity the efforts undertaken;

82 “(5) Except when dispensed directly by a practitioner, other than a pharmacy, to  
83 an ultimate user, no opiate in Schedule II may be dispensed with a strength in excess of 90  
84 morphine milligram equivalents per day for treatment of acute pain.

85 “(7) This subsection shall not apply to a prescription for a patient who is  
86 currently in active treatment for cancer, receiving hospice care from a licensed hospice or  
87 palliative care, or is a resident of a long term care facility, or to any medications that are being  
88 prescribed for use in the treatment of substance abuse or opioid dependence.

89 Sec. 4. Rulemaking.

90 (a) The Mayor shall promulgate regulations to limit the amount and strength of opiate  
91 prescription drugs dispensed by a pharmacy in the District of Columbia as a result of a  
92 prescription prescribed by an individual licensed to prescribe opiates in a jurisdiction not in the  
93 District of Columbia who is not subject to the requirements of this subsection. Such regulations,  
94 at a minimum, shall:

95 (1) Limit to seven days an initial prescription opiate dispensed by a pharmacy for  
96 an acute condition;

97 (2) Limit the number of prescription opiates that may be dispensed by a  
98 pharmacy after an initial prescription for an acute condition;

99 (3) Limit the strength of any prescription opiate dispensed by a pharmacy for an  
100 acute condition.

101 (b) The Mayor shall promulgate regulations to require practitioners to disclose annually  
102 to the Department of Health the number of opiate prescription doses prescribed in a manner that  
103 does not disclose any identifying information patients.

104 Sec. 5. Fiscal impact statement.

105           The Council adopts the fiscal impact statement in the committee report as the fiscal  
106 impact statement required by section 4a of the General Legislative Procedures Act of 1975,  
107 approved October 16, 2006 (120 Stat. 2038; D.C. Official Code § 1-301.47a).

108           Sec. 6. Effective date.

109           This act shall take effect following approval by the Mayor (or in the event of veto by the  
110 mayor, action by the Council to override the veto), a 30-day period of Congressional review as  
111 provided in section 602(c)(1) of the District of Columbia Home Rule Act, approved December  
112 24, 1973, (87 Stat. 813; D.C. Official Code § 1-206.02(c)(1)), and publication in the District of  
113 Columbia Register.