



Councilmember Kenyan R. McDuffie

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A BILL

IN THE COUNCIL OF THE DISTRICT OF COLUMBIA

To amend, on an emergency basis, due to congressional review, the Legalization of Marijuana for Medical Treatment Initiative of 1999 to allow the Alcoholic Beverage and Cannabis Board (“ABC Board”) to issue temporary non-resident registration identification cards that are valid for periods between 3 days and no longer than one year in length, allow licensed testing laboratories to receive and test samples of medical cannabis products from qualifying patients, allow licensed testing laboratories to conduct quality assurance or research and development testing for cultivation centers and manufacturers, amend the definition of a social equity applicant to include arrests and convictions of qualifying family members for a cannabis or drug offense, expand the list of eligible family members under the social equity applicant definition to include siblings and grandparents, clarify that existing licensed cultivation centers and retailers and applicants that scored 150 points or more during the open application period that occurred between November 29, 2021 and March 28, 2022, that are authorized by statute to receive a cultivation center, manufacturer, or retailer license apart from a designated open application period are not counted in calculating the 50% set aside requirement, clarify that the 5 cultivation center registration applicants that scored 150 points or more during the same open application period shall automatically receive a manufacturer license provided that they pay the annual fee and register with the ABC Board, allow the Alcoholic Beverage and Cannabis Administration to issue conditional licenses to testing laboratory applicants, and to waive the application fee for testing laboratory licenses.

BE IT ENACTED BY THE COUNCIL OF THE DISTRICT OF COLUMBIA, That this act may be cited as the “Medical Cannabis Clarification and Non-Resident Patient Access Congressional Review Emergency Amendment Act of 2023”.

40 Sec. 2. The Legalization of Marijuana for Medical Treatment Initiative of 1999, effective
41 February 25, 2010 (D.C. Law 13-315; D.C. Official Code § 7-1671.01 *et seq.*), is amended as
42 follows:

43 (a) Section 2 (D.C. Official Code § 7-1671.01) is amended as follows:

44 (1) Paragraph (13B)(B) is amended by striking the phrase “30-day registration
45 identification card” and inserting the phrase "registration identification card valid for periods
46 established by the ABC Board by rulemaking, which are between 3 days and no longer than one
47 year in length” in its place.

48 (2) Paragraph (20C)(B) is amended by striking the phrase “or has a non-parent
49 legal guardian who is or has been incarcerated” and inserting the phrase “or has a non-parent
50 legal guardian, or a grandparent or a sibling who is or has been arrested, convicted, or
51 incarcerated”.

52 (b) Section 6(b) (D.C. Official Code § 7-1671.05(b)) is amended as follows:

53 (1) Paragraph (4) is amended as follows:

54 (A) Subparagraph (A) is amended by striking the phrase “30 days” and
55 inserting the phrase "periods established by the ABC Board by rulemaking, which are between 3
56 days and no longer than one year in length”.

57 (B) Subparagraph (B) is amended by striking the phrase "30-day”.

58 (2) Paragraph (5)(C) is amended by striking the phrase "3 years.” and inserting
59 the phrase “3 years, except for temporary non-resident registration identification cards that are
60 valid for periods established by the ABC Board by rulemaking, which shall be between 3 days
61 and no longer than one year in length.” in its place.

62 (3) A new paragraph (11A) is added to read as follows:

63 “(11A) Allow testing laboratories to:

64 “(A) Receive and test samples of medical cannabis products from
65 qualifying patients; provided, that the qualifying patient must present proof that he or she is
66 currently registered, and that the medical cannabis product was purchased from a retailer or
67 internet retailer licensed with ABCA.

68 “(B) Receive and test samples of medical cannabis products from licensed
69 cultivation centers or manufacturers for purposes of quality assurance or research and
70 development. Samples collected for quality assurance or research and development testing may
71 be selected by the cultivation center or manufacturer non-randomly. Any tests conducted for
72 purposes of quality assurance or research and development shall not satisfy the requirements of
73 paragraphs (8) through (11) of this subsection.”.

74 (c) Section 7 (D.C. Official Code § 7-1671.06) is amended as follows:

75 (1) Subsection (h) is amended by striking the phrase "cultivation centers who
76 receive a manufacturer’s license pursuant to subsection (d) of this section.” and inserting the
77 phrase "cultivation centers and retailers, and applicants who scored 150 points or more during
78 the ABC Board open application period that occurred between November 29, 2021 and March
79 28, 2022, who receive a cultivation center, manufacturer, or retailer’s license pursuant to
80 subsections (d), (w), (x) and (y) of this section.”.

81 (2) Subsection (k)(1) is amended to read as follows:

82 “(k)(1) The ABC Board shall be authorized to issue a one-year conditional license for a
83 cultivation center, retailer, internet retailer, manufacturer, courier, or testing laboratory that does
84 not currently have a proposed location.”.

85 (3) Subsection (n)(2) is amended to read as follows:

86 “(2)(A) The ABC Board shall, by rules issued pursuant to section 14, establish the
87 initial application and renewal fees for cultivation center, manufacturer, retailer, internet retailer,
88 and courier licenses. The ABC Board may revise these fees as considered necessary.

89 “(B) There shall be no initial application fee for a testing laboratory
90 license. Renewal fees for a testing laboratory license shall be established by rules issued pursuant
91 to sub-paragraph (A) of this paragraph.”.

92 (4) A new subsection (y) is added to read as follows:

93 “(y) The 5 cultivation center registration applicants that submitted medical
94 cannabis facility registration applications to the ABC Board between November 29, 2021 and
95 March 28, 2022, that scored 150 points or more shall automatically receive a manufacturer
96 license provided that the annual fee is paid after the effective date of the Medical Cannabis
97 Manufacturer Clarification Emergency Amendment Act of 2023, effective June 29, 2023 (D.C.
98 Act 25-152; 70 DCR ___); provided, that the applicant registers on a form provided by ABCA
99 with the ABC Board by May 1, 2024.”.

100 Sec. 3. Repealer.

101 The Medical Cannabis Manufacturer Clarification Temporary Amendment Act of 2023,
102 effective August 23, 2023 (D.C. Law 25-43; 70 DCR 9685), is repealed.

103

104 Sec. 4. Fiscal impact statement.

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

108 Sec. 5. 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), and shall remain in effect for no longer than
111 90 days, as provided for emergency acts of the Council of the District of Columbia in section
112 412(a) of the District of Columbia Home Rule Act, approved December 24, 1973 (87 Stat. 788;
113 D.C. Official Code § 1-204.12(a)).