

117TH CONGRESS
2D SESSION

H. R. 5657

AN ACT

To amend the Controlled Substances Act to make marijuana accessible for use by qualified marijuana researchers for medical purposes, and for other purposes.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

1 **SECTION 1. SHORT TITLE.**

2 This Act may be cited as the “Medical Marijuana Re-
3 search Act”.

4 **SEC. 2. FACILITATING MARIJUANA RESEARCH.**

5 (a) **PRODUCTION AND SUPPLY.**—The Secretary of
6 Health and Human Services—

7 (1) until the date on which the Secretary deter-
8 mines that manufacturers and distributors (other
9 than the Federal Government) can ensure a suffi-
10 cient supply of marijuana (as defined in section 102
11 of the Controlled Substances Act (21 U.S.C. 802),
12 as amended by section 8) intended for research by
13 qualified marijuana researchers registered pursuant
14 to paragraph (3) of section 303(f) of the Controlled
15 Substances Act (21 U.S.C. 823(f)), as added by sec-
16 tion 3, shall—

17 (A) continue, through grants, contracts, or
18 cooperative agreements, to produce marijuana
19 through the National Institute on Drug Abuse
20 Drug Supply Program;

21 (B) not later than one year after the date
22 of enactment of this Act, act jointly with the
23 Attorney General of the United States to estab-
24 lish and implement a specialized process for
25 manufacturers and distributors, notwith-
26 standing the registration requirements of sec-

1 tion 303 of such Act (21 U.S.C. 823), to supply
2 qualified marijuana researchers with marijuana
3 products—

4 (i) available through State-authorized
5 marijuana programs; and

6 (ii) consistent with the guidance
7 issued under subsection (c); and

8 (C) not later than 60 days after the date
9 of enactment of this Act, jointly convene with
10 the Attorney General a meeting to initiate the
11 development of the specialized process described
12 in subparagraph (B); and

13 (2) beyond the date specified in paragraph (1),
14 may, at the Secretary's discretion, continue—

15 (A) through grants, contracts, or coopera-
16 tive agreements, to so produce marijuana; and

17 (B) to implement such specialized process.

18 (b) REQUIREMENT TO VERIFY REGISTRATION.—Be-
19 fore supplying marijuana to any person through the Na-
20 tional Institute on Drug Abuse Drug Supply Program or
21 through implementation of the specialized process estab-
22 lished under subsection (a)(1)(B), the Secretary of Health
23 and Human Services shall—

24 (1) require the person to submit documentation
25 demonstrating that the person is a qualified mari-

1 juana researcher seeking to conduct research pursu-
2 ant to section 303(f)(3) of the Controlled Substances
3 Act, as added by subsection (d) of this section, or
4 a manufacturer duly registered under section 303(l)
5 of the Controlled Substances Act, as added by sec-
6 tion 3 of this Act; and

7 (2) not later than 60 days after receipt of such
8 documentation, review such documentation and
9 verify that the marijuana will be used for such re-
10 search (and for no other purpose authorized pursu-
11 ant to this Act or the amendments made by this
12 Act).

13 (c) GUIDANCE ON USE OF STATE-AUTHORIZED
14 MARIJUANA PROGRAMS.—Not later than 180 days after
15 the date of the enactment of this Act, the Secretary of
16 Health and Human Services shall issue guidance related
17 to marijuana from State-authorized marijuana programs
18 for research.

19 (d) RESEARCH.—Section 303(f) of the Controlled
20 Substances Act (21 U.S.C. 823(f)) is amended—

21 (1) by redesignating paragraphs (1) through
22 (5) as subparagraphs (A) through (E), respectively;

23 (2) by striking “(f) The Attorney General” and
24 inserting “(f)(1) The Attorney General”;

1 (3) by striking “Registration applications” and
2 inserting the following:

3 “(2) Registration applications”;

4 (4) in paragraph (2), as so designated, by strik-
5 ing “schedule I” each place that term appears and
6 inserting “schedule I, except marijuana,”;

7 (5) by striking “Article 7” and inserting the
8 following:

9 “(4) Article 7”; and

10 (6) by inserting before paragraph (4), as so
11 designated, the following:

12 “(3)(A) The Attorney General shall register the ap-
13 plicant to conduct research with marijuana (including any
14 derivative, extract, preparation, and compound thereof) if,
15 irrespective of whether the applicant is registered pursu-
16 ant to paragraphs (1) and (2)—

17 “(i) the applicant meets the requirements for
18 being registered under such paragraphs to dispense,
19 or conduct research with respect to, controlled sub-
20 stances in schedule I, II, III, IV, or V;

21 “(ii) the applicant is compliant with, and au-
22 thorized to conduct the activities described in clause
23 (i) under, the laws of the State in which the appli-
24 cant practices; and

1 “(iii) in the case of an applicant pursuing clinical research, the applicant’s clinical research protocol has been reviewed and authorized to proceed by the Secretary under section 505(i) of the Federal Food, Drug, and Cosmetic Act.

6 “(B) An applicant registered under subparagraph (A) shall be referred to in this section as a ‘qualified marijuana researcher’.

9 “(C)(i) Not later than 60 days after the date on which the Attorney General receives a complete application for registration under this paragraph, the Attorney General shall approve or deny the application.

13 “(ii) For purposes of clause (i), an application shall be deemed complete when the applicant has submitted documentation showing that the requirements under subparagraph (A) are satisfied.

17 “(iii) In the case of a denial under clause (i), the Attorney General shall provide a written explanation of the basis for the denial.

20 “(D) The Attorney General shall grant an application for registration under this paragraph unless the Attorney General determines that the issuance of the registration would be inconsistent with the public interest. In determining the public interest, the following factors shall be considered:

1 “(i) The applicant’s experience in dispensing, or
2 conducting research with respect to, controlled sub-
3 stances.

4 “(ii) The applicant’s conviction record under
5 Federal or State laws relating to the manufacture,
6 distribution, or dispensing of controlled substances.

7 “(iii) Compliance with applicable State or local
8 laws relating to controlled substance misuse or diver-
9 sion.

10 “(iv) Such other conduct which may threaten
11 the public health and safety.

12 “(E)(i) A qualified marijuana researcher shall store
13 marijuana to be used in research in a securely locked, sub-
14 stantially constructed cabinet.

15 “(ii) Except as provided in clause (i), any security
16 measures required by the Attorney General for applicants
17 conducting research with marijuana pursuant to a reg-
18 istration under this paragraph shall be consistent with the
19 security measures for applicants conducting research on
20 other controlled substances in schedule II that have a
21 similar risk of diversion and abuse.

22 “(F)(i) If the Attorney General grants an application
23 for registration under this paragraph, the applicant may
24 amend or supplement the research protocol and proceed
25 with the research under such amended or supplemented

1 protocol, without additional review or approval by the At-
2 torney General or the Secretary of Health and Human
3 Services if the applicant does not change the type of mari-
4 juana (including any derivative, extract, preparation, and
5 compound thereof), the source of the marijuana, or the
6 conditions under which the marijuana is stored, tracked,
7 or administered.

8 “(ii) If an applicant amends or supplements the re-
9 search protocol or initiates research on a new research
10 protocol under clause (i), the applicant shall, in order to
11 renew the registration under this paragraph, provide no-
12 tice to the Attorney General of the amended or supple-
13 mented research protocol or any new research protocol in
14 the applicant’s renewal materials.

15 “(iii)(I) If an applicant amends or supplements a re-
16 search protocol and the amendment or supplement in-
17 volves a change to the type of marijuana, the source of
18 the marijuana, or conditions under which the marijuana
19 is stored, tracked, or administered, the applicant shall pro-
20 vide notice to the Attorney General not later than 30 days
21 before proceeding on such amended or supplemental re-
22 search or new research protocol, as the case may be.

23 “(II) If the Attorney General does not object during
24 the 30-day period following a notification under subclause

1 (I), the applicant may proceed with the amended or sup-
2 plemental research or new research protocol.

3 “(iv) The Attorney General may object to an amend-
4 ed or supplemental protocol or a new research protocol
5 under clause (i) or (iii) only if additional security meas-
6 ures are needed to safeguard against diversion or abuse.

7 “(G) If marijuana is listed on a schedule other than
8 schedule I, the provisions of paragraphs (1), (2), and (4)
9 that apply to research with a controlled substance in the
10 applicable schedule shall apply to research with marijuana
11 or that compound, as applicable, in lieu of the provisions
12 of subparagraphs (A) through (F) of this paragraph.

13 “(H) Nothing in this paragraph shall be construed
14 as limiting the authority of the Secretary under section
15 505(i) of the Federal Food, Drug, and Cosmetic Act or
16 over requirements related to research protocols, including
17 changes in—

18 “(i) the method of administration of marijuana;

19 “(ii) the dosing of marijuana; and

20 “(iii) the number of individuals or patients in-
21 volved in research.”.

1 **SEC. 3. MANUFACTURE AND DISTRIBUTION OF MARIJUANA**
2 **FOR USE IN LEGITIMATE RESEARCH.**

3 Section 303 of the Controlled Substances Act (21
4 U.S.C. 823), as amended by section 2, is further amended
5 by adding at the end the following:

6 “(1) REGISTRATION OF PERSONS TO MANUFACTURE
7 AND DISTRIBUTE MARIJUANA FOR USE IN LEGITIMATE
8 RESEARCH.—

9 “(1) REGISTRATION OF MANUFACTURERS.—

10 “(A) IN GENERAL.—Beginning not later
11 than the day that is 1 year after the date of en-
12 actment of the Medical Marijuana Research
13 Act, the Attorney General, pursuant to sub-
14 section (f)(3) and subject to subparagraph (B)
15 of this paragraph, shall register an applicant to
16 manufacture marijuana (including any deriva-
17 tive, extract, preparation, and compound there-
18 of) that is intended for—

19 “(i) use by qualified marijuana re-
20 searchers for research pursuant to sub-
21 section (f)(3); or

22 “(ii) subsequent downstream manu-
23 facture by a duly registered manufacturer
24 for use by qualified marijuana researchers
25 for research pursuant to subsection (f)(3).

1 “(B) PUBLIC INTEREST.—The Attorney
2 General shall register an applicant under sub-
3 paragraph (A) unless the Attorney General de-
4 termines that the issuance of such registration
5 is inconsistent with the public interest. In deter-
6 mining the public interest, the Attorney General
7 shall take into consideration—

8 “(i) maintenance of effective controls
9 against diversion of marijuana and any
10 controlled substance compounded there-
11 from into other than legitimate medical,
12 scientific, or research channels;

13 “(ii) compliance with applicable State
14 and local laws relating to controlled sub-
15 stance misuse and diversion;

16 “(iii) prior conviction record of the
17 applicant under Federal or State laws re-
18 lating to the manufacture, distribution, or
19 dispensing of such substances; and

20 “(iv) such other conduct which may
21 threaten the public health and safety.

22 “(2) REGISTRATION OF DISTRIBUTORS.—

23 “(A) IN GENERAL.—Beginning not later
24 than the day that is 1 year after the date of en-
25 actment of the Medical Marijuana Research

1 Act, the Attorney General shall register an ap-
2 plicant to distribute marijuana (including any
3 derivative, extract, preparation, and compound
4 thereof) that is intended for use by qualified
5 marijuana researchers for research pursuant to
6 subsection (f)(3) or intended for subsequent
7 downstream manufacture by a duly registered
8 manufacturer for use by qualified marijuana re-
9 searchers for research pursuant to such sub-
10 section, unless the Attorney General determines
11 that the issuance of such registration is incon-
12 sistent with the public interest.

13 “(B) PUBLIC INTEREST.—In determining
14 the public interest under subparagraph (A), the
15 Attorney General shall take into consider-
16 ation—

17 “(i) the factors specified in clauses (i),
18 (ii), (iii), and (iv) of paragraph (1)(B); and

19 “(ii) past experience in the distribu-
20 tion of controlled substances, and the exist-
21 ence of effective controls against diversion.

22 “(3) NO LIMIT ON NUMBER OF MANUFACTUR-
23 ERS AND DISTRIBUTORS.—Notwithstanding any
24 other provision of law, the Attorney General shall
25 not impose or implement any limit on the number of

1 persons eligible to be registered to manufacture or
2 distribute marijuana pursuant to paragraph (1) or
3 (2).

4 “(4) REQUIREMENT TO VERIFY USE FOR LE-
5 GITIMATE RESEARCH.—As a condition of registra-
6 tion under this section to manufacture or distribute
7 marijuana, the Attorney General shall require the
8 registrant—

9 “(A) to require any person to whom the
10 marijuana will be supplied to submit docu-
11 mentation demonstrating that the marijuana
12 (including any derivative, extract, preparation,
13 and compound thereof) will be used by qualified
14 marijuana researchers for research pursuant to
15 subsection (f)(3) or for subsequent downstream
16 manufacture by a duly registered manufacturer
17 for use by qualified marijuana researchers for
18 research pursuant to such subsection;

19 “(B) in the case of distribution, to com-
20 plete, with respect to that distribution, the ap-
21 propriate order form in accordance with section
22 308 and to upload such forms to the system
23 used by the Drug Enforcement Administration
24 for such distribution;

1 “(C) to include in the labeling of any mari-
2 juana so manufactured or distributed—

3 “(i) the following statement: ‘This
4 material is for biomedical and scientific re-
5 search purposes only.’; and

6 “(ii) the name of the requestor of the
7 marijuana;

8 “(D) to limit the transfer and sale of any
9 marijuana under this subsection—

10 “(i) to researchers who are registered
11 under this Act to conduct research with
12 marijuana or to manufacturers duly reg-
13 istered under this subsection; and

14 “(ii) for purposes of use in preclinical
15 research or in a clinical investigation pur-
16 suant to an investigational new drug ex-
17 emption under 505(i) of the Federal Food,
18 Drug, and Cosmetic Act or for the pur-
19 poses of further manufacturing of mari-
20 juana; and

21 “(E) to transfer or sell any marijuana
22 manufactured under this subsection only with
23 prior, written consent for the transfer or sale by
24 the Attorney General.

1 “(5) TIMING.—Not later than 60 days after re-
 2 ceipt of a request for registration under this sub-
 3 section to manufacture or distribute marijuana, the
 4 Attorney General shall—

5 “(A) grant or deny the request; and

6 “(B) in the case of a denial, provide a
 7 written explanation of the basis for the denial.

8 “(6) DEEMED APPROVAL.—If the Attorney
 9 General fails to grant or deny a request for registra-
 10 tion under this subsection to manufacture or dis-
 11 tribute marijuana within the 60-day period referred
 12 to in paragraph (5), such request is deemed ap-
 13 proved.”.

14 **SEC. 4. TERMINATION OF INTERDISCIPLINARY REVIEW**
 15 **PROCESS FOR NON-NIH-FUNDED QUALIFIED**
 16 **MARIJUANA RESEARCHERS.**

17 The Secretary of Health and Human Services may
 18 not—

19 (1) reinstate the Public Health Service inter-
 20 disciplinary review process described in the guidance
 21 entitled “Guidance on Procedures for the Provision
 22 of Marijuana for Medical Research” (issued on May
 23 21, 1999); or

24 (2) create an additional review of scientific pro-
 25 tocols that is only conducted for research on mari-

1 juana other than the review of research protocols
2 performed at the request of a qualified marijuana
3 researcher conducting nonhuman research that is
4 not federally funded, in accordance with section
5 303(f)(3)(A) of the Controlled Substances Act, as
6 added by section 2 of this Act.

7 **SEC. 5. CONSIDERATION OF RESULTS OF RESEARCH.**

8 Immediately upon the approval by the Food and
9 Drug Administration of an application for a drug that
10 contains marijuana (as defined in section 102 of the Con-
11 trolled Substances Act (21 U.S.C. 802), as amended by
12 section 8 of this Act) under section 505 of the Federal
13 Food, Drug, and Cosmetic Act (21 U.S.C. 355), and (irre-
14 spective of whether any such approval is granted) not later
15 than the date that is 5 years after the date of enactment
16 of this Act, the Secretary of Health and Human Services
17 shall—

18 (1) conduct a review of existing medical and
19 other research with respect to marijuana;

20 (2) submit a report to the Congress on the re-
21 sults of such review; and

22 (3) include in such report whether, taking into
23 consideration the factors listed in section 201(c) of
24 the Controlled Substances Act (21 U.S.C. 811(c)),
25 as well as any potential for medical benefits, any

1 gaps in research, and any impacts of Federal restric-
2 tions and policy on research, marijuana should be
3 transferred to a schedule other than schedule I (if
4 marijuana has not been so transferred already).

5 **SEC. 6. PRODUCTION QUOTAS FOR MARIJUANA GROWN**
6 **FOR LEGITIMATE, SCIENTIFIC RESEARCH.**

7 Section 306 of the Controlled Substances Act (21
8 U.S.C. 826) is amended by adding at the end the fol-
9 lowing:

10 “(j) The Attorney General may only establish a quota
11 for production of marijuana that is manufactured and dis-
12 tributed in accordance with the Medical Marijuana Re-
13 search Act that meets the changing medical, scientific, and
14 industrial needs for marijuana.”.

15 **SEC. 7. ARTICLE 28 OF THE SINGLE CONVENTION ON NAR-**
16 **COTIC DRUGS.**

17 Article 28 of the Single Convention on Narcotic
18 Drugs shall not be construed to prohibit, or impose addi-
19 tional restrictions upon, research involving marijuana, or
20 the manufacture, distribution, or dispensing of marijuana,
21 that is conducted in accordance with the Controlled Sub-
22 stances Act (21 U.S.C. 801 et seq.), this Act, and the
23 amendments made by this Act.

1 **SEC. 8. DEFINITIONS.**

2 (a) QUALIFIED MARIJUANA RESEARCHER.—In this
3 Act, the term “qualified marijuana researcher” has the
4 meaning given the term in section 303(f)(3) of the Con-
5 trolled Substances Act, as added by section 2(d) of this
6 Act.

7 (b) UPDATING TERM.—Section 102(16) of the Con-
8 trolled Substances Act (21 U.S.C. 802(16)) is amended—

9 (1) in subparagraph (A), by striking “the term
10 ‘marihuana’ means” and inserting “the terms ‘mari-
11 huana’ and ‘marijuana’ mean”; and

12 (2) in subparagraph (B), by striking “The term
13 ‘marihuana’ does not” and inserting “The terms
14 ‘marihuana’ and ‘marijuana’ do not”.

15 **SEC. 9. DETERMINATION OF BUDGETARY EFFECTS.**

16 The budgetary effects of this Act, for the purpose of
17 complying with the Statutory Pay-As-You-Go Act of 2010,
18 shall be determined by reference to the latest statement
19 titled “Budgetary Effects of PAYGO Legislation” for this
20 Act, submitted for printing in the Congressional Record
21 by the Chairman of the House Budget Committee, pro-

- 1 vided that such statement has been submitted prior to the
- 2 vote on passage.

Passed the House of Representatives April 4, 2022.

Attest:

Clerk.

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