# ASSEMBLY BILL

No. 1037

### **Introduced by Assembly Member Elhawary**

February 20, 2025

An act to amend Section 4145.5 of the Business and Professions Code, to amend Section 1714.22 of the Civil Code, and to amend Sections 1797.197, 11364, 11364.5, 11364.7, 11372.7, 11832, 11834.026, 11834.26, 11999, 11999.1, 120780.1, 120780.2, 120780.5, and 121349, of, to amend the heading of Division 10.5 (commencing with Section 11750) of, to amend the heading of Part 4 (commencing with Section 120775) of Division 105 of, and to amend the heading of Chapter 1.5 (commencing with Section 120780) of Part 4 of Division 105 of, and to repeal and add Section 11999.2 of, and to repeal and add the heading of Division 10.7 (commencing with Section 11999) of, the Health and Safety Code, relating to public health.

#### LEGISLATIVE COUNSEL'S DIGEST

AB 1037, as introduced, Elhawary. Public health: substance use disorder.

(1) Existing law, until January 1, 2026, authorizes a physician or pharmacist, without a prescription or permit, to furnish hypodermic needles and syringes for human use to a person 18 years of age or older, and authorizes a person 18 years of age or older to, without a prescription or license, obtain hypodermic needles and syringes solely for personal use from a physician or pharmacist, as a public health measure, as specified. Existing law requires a pharmacist that provides nonprescription syringes to provide information on access to testing and treatment for HIV and hepatitis C.

This bill would extend this authority indefinitely and would additionally require a pharmacist to provide information on other bloodborne diseases.

(2) Under existing law, a licensed health care provider who is authorized by law to prescribe an opioid antagonist may issue standing orders for the distribution of an opioid antagonist to a person at risk of an opioid-related overdose or to a family member, friend, or other person in a position to assist a person at risk of an opioid-related overdose. Existing law requires that a person who receives an opioid antagonist pursuant to a standing order or otherwise possesses an opioid antagonist receive training, as specified. Existing law provides that a person who is trained in the use of an opioid antagonist and acts with reasonable care and in good faith is not subject to professional review, liable in a civil action, or subject to criminal prosecution.

This bill would expand the above-described authorizations to those who are at risk of or who may be in a position to assist a person experiencing any overdose and would strike the requirement that those who receive and possess opioid antagonists receive training. The bill would authorize a person in a position to assist a person at risk of an overdose to possess an opioid antagonist and subsequently dispense or distribute an opioid antagonist to a person at risk of an overdose or another person in a position to assist a person at risk of an overdose. The bill would instead exempt a person who administers an opioid antagonist with reasonable care and in good faith, whether or not they were trained, from professional review, liability in a civil action, or criminal prosecution.

(3) Existing law defines drug paraphernalia to include testing equipment designed for use or marketed for use in identifying, or in analyzing the strength, effectiveness, or purity of, controlled substances, subject to exceptions, and prohibits, among other things, the manufacture, sale, and possession, as specified, of drug paraphernalia. Existing law exempts from these prohibitions specified professionals, such as pharmacists, manufacturers, and dentists, under certain circumstances. Existing law authorizes a court, in determining whether an object is drug paraphernalia, to consider specified facts and circumstances, such as the expert testimony.

This bill would remove testing equipment from the definition of drug paraphernalia and would expand the group of individuals exempt from drug paraphernalia prohibition to include a person who is at risk of overdose, a family member, friend, or other person in a position to assist a person at risk of overdose, in order to reduce the spread of HIV, viral hepatitis, and other bloodborne infections among the intravenous drug user population within California. The bill would additionally authorize a court, in determining whether an object is paraphernalia, to consider whether the possession of an object is related to or a result of a substance use disorder treatment or recovery program, harm reduction program, syringe exchange program, or consistent with best clinical practices, as specified.

(4) Existing law imposes a drug program fee for each separate controlled substance offense, as specified, to be deposited by the county treasurer in a drug program fund. Existing law requires that a portion of the fund be allocated to primary prevention programs in the community.

This bill would state that primary prevention programs may include those activities as determined by the Substance Abuse and Mental Health Services Administration.

(5) Existing law requires the State Department of Health Care Services to license and regulate facilities that provide residential nonmedical services to adults who are recovering from problems related to alcohol, drug, or alcohol and drug misuse or abuse, and who need alcohol, drug, or alcohol and drug recovery treatment or detoxification services. Existing law requires these programs to be certified, except as specified. Existing law authorizes a licensed alcohol or other drug recovery or treatment facility to permit incidental medical services, as defined, to be provided to a resident at the facility premises by a licensed physician and surgeon or other health care practitioner under specified limited circumstances, including that the resident has signed an admission agreement. Existing law requires a licensee to develop a plan to address when a resident relapses, including when a resident is on the licensed premises after consuming alcohol or using illicit drugs.

This bill would require the department, on or before July 1, 2026, to offer a combined application for entities to be certified as an alcohol or other drug program and to provide incidental medical services, as defined. The bill would prohibit an admission agreement from requiring a person to be abstinent and not intoxicated in order to be admitted to care or continue treatment. The bill would require a licensee to prioritize the individual maintaining some level of connection to treatment, rather than disconnection from treatment, following a relapse.

Existing law defines "drug- or alcohol-related program" as any program designed to reduce the unlawful use of, or assist those who

engage in the unlawful use of, drugs or alcohol, through various means, such as intervention, treatment, and enforcement, among others. Existing law prohibits the encumbrance of state funds for a drug- or alcohol-related program unless it contains a component that explains that there is no unlawful use of drugs or alcohol and requires all aspects of a drug- or alcohol-related program receiving state funds to be consistent with the "no lawful use" message.

This bill would redefine that term to mean any program designed to assist persons with substance use disorders and would strike enforcement from the specified means. The bill would repeal the above-described provisions related to the "no lawful use" message and would instead require that a drug- or alcohol-related program be consistent with the best clinical practices as determined by the Substance Abuse and Mental Health Services Administration and the American Society of Addiction Medicine in order to receive state funds.

(6) Existing law authorizes a public entity, as defined, that receives General Fund money from the Office of AIDS in the State Department of Public Health for HIV prevention and education to use that money to support clean needle and syringe exchange projects authorized by the public entity. Existing law authorizes the money to be used for the purchase of sterile hypodermic needles and syringes, subject to specified conditions, such as the portion of funds used for purchasing sterile hypodermic needles and syringes does not exceed 7.5% of the total amount of the funds received by the entity for HIV prevention. Existing law requires that an entity apply for authorization to provide hypodermic needle and syringe services and requires that an entity demonstrate in its application that it complies with certain minimum standards, including that it has adequate funding to provide certain services at reasonably projected program participation levels.

This bill would expand the diseases that a public entity receiving this General Fund money may focus on to include viral hepatitis and other bloodborne diseases and would strike the above-described specified conditions to instead authorize that the money may be used for the purchase of sterile hypodermic needles and syringes as part of a clean needle and syringe exchange program in alignment with primary prevention activities as determined by the Substance Abuse and Mental Health Services Administration in the course of administering the Substance Use Prevention, Treatment, and Recovery Services Block Grant. The bill would require an entity applying for authorization to provide hypodermic needle and syringe exchange services to

demonstrate in its application that it complies with certain minimum standards, including that it has the ability to provide certain services at reasonably projected program participation levels within 3 months of authorization.

Vote: majority. Appropriation: no. Fiscal committee: yes. State-mandated local program: no.

## The people of the State of California do enact as follows:

SECTION 1. Section 4145.5 of the Business and Professions
 Code is amended to read:

3 4145.5. (a) Notwithstanding any other provision of law, a pharmacist or physician may, without a prescription or a permit, 4 furnish hypodermic needles and syringes for human use, and a 5 person may, without a prescription or license, obtain hypodermic 6 7 needles and syringes from a pharmacist or physician for human 8 use, if the furnisher has previously been provided a prescription 9 or other proof of a legitimate medical need requiring a hypodermic 10 needle or syringe to administer a medicine or treatment.

(b) Notwithstanding any other provision of law, and until 11 12 January 1, 2026, law, as a public health measure intended to 13 prevent the transmission of HIV, viral hepatitis, and other 14 bloodborne diseases among persons who use syringes and hypodermic needles, and to prevent subsequent infection of sexual 15 partners, newborn children, or other persons, a physician or 16 pharmacist may, without a prescription or a permit, furnish 17 18 hypodermic needles and syringes for human use to a person 18 19 years of age or older, and a person 18 years of age or older may, 20 without a prescription or license, obtain hypodermic needles and 21 syringes solely for personal use from a physician or pharmacist. (c) Notwithstanding any other provision of law, a pharmacist, 22

veterinarian, or person licensed pursuant to Section 4141 may,
without a prescription or license, furnish hypodermic needles and
syringes for use on animals, and a person may, without a
prescription or license, obtain hypodermic needles and syringes
from a pharmacist, veterinarian, or person licensed pursuant to
Section 4141 for use on animals.

29 (d) A pharmacy that furnishes nonprescription hypodermic 30 needles and syringes shall store hypodermic needles and syringes

in a manner that ensures that they are available only to authorized 1 2

personnel, and are not accessible to other persons.

3 (e) In order to provide for the safe disposal of hypodermic 4 needles and syringes, a pharmacy or hypodermic needle and syringe 5 exchange program that furnishes nonprescription hypodermic 6 needles and syringes shall counsel consumers on safe disposal and 7 provide consumers with one or more of the following disposal 8 options:

9 (1) It shall establish an onsite, safe, hypodermic needle and 10 syringe collection and disposal program that meets applicable state 11 and federal standards for collection and disposal of medical sharps 12 waste.

13 (2) It shall furnish, or make available, mail-back sharps 14 containers authorized by the United States Postal Service that meet 15 applicable state and federal requirements for the transport of medical sharps waste, and shall provide tracking forms to verify 16 17 destruction at a certified disposal facility.

(3) It shall furnish, or make available, a sharps container that 18 meets applicable state and federal standards for collection and 19 20 disposal of medical sharps waste.

21 (f) Until January 1, 2026, a A pharmacy that furnishes 22 nonprescription syringes shall provide written information or verbal 23 counseling to consumers at the time of furnishing or sale of nonprescription hypodermic needles or syringes on how to do the 24 25 following:

26 (1) Access drug treatment.

27 (2) Access testing and treatment for HIV and hepatitis C. HIV,

28 viral hepatitis, and other bloodborne diseases.

29 (3) Safely dispose of sharps waste.

30 SEC. 2. Section 1714.22 of the Civil Code is amended to read:

31 1714.22. (a) For purposes of this section, the following 32 definitions apply:

33 (1) "Opioid antagonist" means naloxone hydrochloride or any 34 other opioid antagonist that is approved by the United States Food 35 and Drug Administration for the treatment of an opioid overdose. 36 (2) "Opioid overdose prevention and treatment training

37 program" means any program operated by a local health 38 jurisdiction or that is registered by a local health jurisdiction to

39 train individuals to prevent, recognize, and respond to an opiate

- 1 overdose, and that provides, at a minimum, training in all of the 2 following:
- 3 (A) The causes of an opiate overdose.
- 4 (B) Mouth to mouth resuscitation.
- 5 (B) Basic life support.
- 6 (C) How to contact appropriate emergency medical services.
- 7 (D) How to administer an opioid antagonist.
- 8 (b) A licensed health care provider who is authorized by law to
- 9 prescribe an opioid antagonist may, if acting with reasonable care, 10 prescribe and subsequently dispense or distribute an opioid 11 antagonist to a person at risk of an opioid-related overdose or to 12 a family member, friend, or other person in a position to assist a
- 13 person at risk of an opioid-related overdose.
- 14 (c) (1) A licensed health care provider who is authorized by
- 15 law to prescribe an opioid antagonist may issue standing orders
- 16 for the distribution of an opioid antagonist to a person at risk of
- 17 an opioid-related overdose or to a family member, friend, or other
- 18 person in a position to assist a person at risk of an opioid-related19 overdose.
- 20 (2) A licensed health care provider who is authorized by law to
- 21 prescribe an opioid antagonist may issue standing orders for the
- 22 administration of an opioid antagonist to a person at risk of an
- 23 opioid-related overdose by a family member, friend, or other person
- 24 in a position to assist a person experiencing or reasonably suspected
- 25 of experiencing an <del>opioid</del> overdose.
- 26 (d) (1) A person who is prescribed or possesses an opioid
- 27 antagonist pursuant to a standing order shall receive the training
- provided by an opioid overdose prevention and treatment training
   program.
- 30 (2) A person who is prescribed an opioid antagonist directly
- 31 from a licensed prescriber shall not be required to receive training
- 32 from an opioid prevention and treatment training program.
- 33 (3) A person who is at risk of an overdose, a family member,
- 34 friend, or other person in a position to assist a person at risk of
- 35 an overdose may possess an opioid antagonist and subsequently
- 36 dispense or distribute an opioid antagonist to a person at risk of
- 37 an overdose or to a family member, friend, or other person in a
- 38 position to assist a person at risk of an overdose.
- 39 <del>(e)</del>

1 (d) A licensed health care provider or a person who is at risk 2 of an overdose, or a family member, friend, or other person in a 3 position to assist a person at risk of an overdose who acts with 4 reasonable care shall not be subject to professional review, be 5 liable in a civil action, or be subject to criminal prosecution for 6 issuing a prescription or order or for possession or distributing an 7 opioid antagonist pursuant to subdivision (b) or (c). 8 (f)

9 (e) Notwithstanding any other law, a person who possesses or distributes an opioid antagonist-pursuant to a prescription or 10 standing order shall not be subject to professional review, be liable 11 12 in a civil action, or be subject to criminal prosecution for this 13 possession or distribution. Notwithstanding any other law, a person 14 not otherwise licensed to administer an opioid-antagonist, but 15 trained as required under paragraph (1) of subdivision (d), antagonist who acts with reasonable care in administering an opioid 16 17 antagonist, in good faith and not for compensation, to a person 18 who is experiencing or is suspected of experiencing an overdose 19 shall not be subject to professional review, be liable in a civil 20 action, or be subject to criminal prosecution for this administration. 21 SEC. 3. Section 1797.197 of the Health and Safety Code is 22 amended to read:

23 1797.197. (a) The authority shall establish training and 24 standards for all prehospital emergency medical care personnel, 25 as defined in paragraph (2) of subdivision (a) of Section 1797.189, 26 regarding the characteristics and method of assessment and 27 treatment of anaphylactic reactions and the use of epinephrine. 28 The authority shall promulgate regulations regarding these matters 29 for use by all prehospital emergency medical care personnel. 30 (b) (1) The authority shall develop and, after approval by the

31 commission pursuant to Section 1799.50, adopt training and 32 standards for all prehospital emergency medical care personnel, as defined in paragraph (2) of subdivision (a) of Section 1797.189, 33 34 regarding the use and administration of naloxone hydrochloride 35 and other opioid antagonists. The authority shall promulgate 36 regulations regarding these matters for use by all prehospital 37 emergency medical care personnel. The authority may adopt 38 existing training and standards for prehospital emergency medical 39 care personnel regarding the statewide use and administration of

naloxone hydrochloride or another opioid antagonist to satisfy the
 requirements of this section.

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3 (2) The medical director of a local EMS agency may, pursuant 4 to Section 1797.221, approve or conduct a trial study of the use 5 and administration of naloxone hydrochloride or other opioid 6 antagonists by any level of prehospital emergency medical care personnel. Training received by prehospital emergency medical 7 8 care personnel specific to the use and administration of naloxone 9 hydrochloride or another opioid antagonist during this trial study 10 may be used towards satisfying the training requirements 11 established pursuant to paragraph (1) regarding the use and 12 administration of naloxone hydrochloride and other opioid 13 antagonists by prehospital emergency medical care personnel.

(3) The training described in paragraphs (1) and (2) shall satisfy
 the requirements of paragraph (1) of subdivision (d) of Section
 1714.22 of the Civil Code.

17 SEC. 4. Section 11364 of the Health and Safety Code is 18 amended to read:

19 11364. (a) It is unlawful to possess an opium pipe or any 20 device, contrivance, instrument, or paraphernalia used for 21 unlawfully injecting or smoking (1) a controlled substance specified 22 in subdivision (b), (c), or (e) or paragraph (1) of subdivision (f) of 23 Section 11054, specified in paragraph (14), (15), or (20) of 24 subdivision (d) of Section 11054, specified in subdivision (b) or 25 (c) of Section 11055, or specified in paragraph (2) of subdivision 26 (d) of Section 11055, or (2) a controlled substance that is a narcotic 27 drug classified in Schedule III, IV, or V. 28 (b) This section-shall does not apply to hypodermic needles or 29 syringes that have been containerized for safe disposal in a

30 container that meets state and federal standards for disposal of
 31 sharps waste.
 32 (a) This section does not early to an individual abtaining

(c) This section does not apply to an individual obtaining
controlled substance checking services, as described in Article 5
(commencing with Section 11300) of Chapter 5.

(d) Until January 1, 2026, as As a public health measure intended
to prevent the transmission of HIV, viral hepatitis, and other
bloodborne diseases among persons who use syringes and
hypodermic needles, and to prevent subsequent infection of sexual
partners, newborn children, or other persons, this section-shall
does not apply to the possession solely for personal use of

1 hypodermic-needles or syringes. needles, syringes, and controlled

2 substance checking equipment, as described in Article 5
3 (commencing with Section 11300) of Chapter 5. This section does

4 not apply to overdose prevention or treatment programs, or

5 individuals in possession of hypodermic needles or syringes or

6 controlled substance checking equipment related to an overdose

7 prevention or treatment training program, as described in Article

8 5 (commencing with Section 11300) of Chapter 5.

9 SEC. 5. Section 11364.5 of the Health and Safety Code is 10 amended to read:

11364.5. (a) Except as authorized by law, a person shall not 11 12 maintain or operate a place of business in which drug paraphernalia 13 is kept, displayed, or offered in any manner, sold, furnished, 14 transferred, or given away unless that drug paraphernalia is 15 completely and wholly kept, displayed, or offered within a separate room or enclosure to which persons under 18 years of age who are 16 17 not accompanied by a parent or legal guardian are excluded. Each 18 entrance to such a room or enclosure shall be signposted in 19 reasonably visible and legible words to the effect that drug 20 paraphernalia is kept, displayed, or offered in the room or enclosure 21 and that minors, unless accompanied by a parent or legal guardian, 22 are excluded.

(b) Except as authorized by law, an owner, manager, proprietor,
or other person in charge of a room or enclosure, within a place
of business, in which drug paraphernalia is kept, displayed, or
offered in any manner, sold, furnished, transferred, or given away
shall not permit or allow a person under 18 years of age to enter,
be in, remain in, or visit the room or enclosure unless that minor
person is accompanied by their parent or legal guardian.

30 (c) Unless authorized by law, a person under 18 years of age
31 shall not enter, be in, remain in, or visit a room or enclosure in a
32 place of business in which drug paraphernalia is kept, displayed,
33 or offered in any manner, sold, furnished, transferred, or given

34 away unless accompanied by their parent or legal guardian.

(d) As used in this section, "drug paraphernalia" means all
equipment, products, and materials of any kind which are intended
for use or designed for use, in planting, propagating, cultivating,
growing, harvesting, manufacturing, compounding, converting,

39 producing, processing, preparing, packaging, repackaging, storing,

40 containing, concealing, injecting, ingesting, inhaling, or otherwise

introducing into the human body a controlled substance. "Drug
 paraphernalia" includes, but is not limited to, all of the following:
 (1) Kits intended for use or designed for use in planting,

4 propagating, cultivating, growing, or harvesting of any species of 5 plant that is a controlled substance or from which a controlled 6 substance can be derived.

7 (2) Kits intended for use or designed for use in manufacturing,
8 compounding, converting, producing, processing, or preparing
9 controlled substances.

(3) Isomerization devices intended for use or designed for usein increasing the potency of any species of plant that is a controlledsubstance.

(4) Scales and balances intended for use or designed for use inweighing or measuring controlled substances.

(5) Diluents and adulterants, such as quinine hydrochloride,
mannitol, mannite, dextrose, and lactose, intended for use or
designed for use in cutting controlled substances.

(6) Separation gins and sifters intended for use or designed foruse in removing twigs and seeds from, or in otherwise cleaning orrefining, cannabis.

(7) Blenders, bowls, containers, spoons, and mixing devices
intended for use or designed for use in compounding controlled
substances.

(8) Capsules, balloons, envelopes, and other containers intended
for use or designed for use in packaging small quantities of
controlled substances.

(9) Containers and other objects intended for use or designedfor use in storing or concealing controlled substances.

(10) Hypodermic syringes, needles, and other objects intended
 for use or designed for use *used* in parenterally injecting controlled
 substances into the human body.

(11) Objects intended for use or designed for use in ingesting,inhaling, or otherwise introducing cannabis, cocaine, hashish, or

34 hashish oil into the human body, such as the following:

(A) Metal, wooden, acrylic, glass, stone, plastic, or ceramic
 pipes with or without screens, permanent screens, hashish heads,

37 or punctured metal bowls.

38 (B) Water pipes.

39 (C) Carburetion tubes and devices.

40 (D) Smoking and carburetion masks.

- 1 (E) Roach clips, meaning objects used to hold burning material,
- 2  $\,$  such as a cannabis cigarette that has become too small or too short  $\,$
- 3 to be held in the hand.
- 4 (F) Miniature cocaine spoons, and cocaine vials.
- 5 (G) Chamber pipes.
- 6 (H) Carburetor pipes.
- 7 (I) Electric pipes.
- 8 (J) Air-driven pipes.
- 9 (K) Chillums.
- 10 (L) Bongs.
- 11 (M) Ice pipes or chillers.
- 12 (12) Testing equipment designed for use or marketed for use in
- 13 identifying, or in analyzing the strength, effectiveness, or purity
- 14 of, controlled substances, except as otherwise provided in
- 15 subdivision (g).
- (e) In determining whether an object is drug paraphernalia, acourt or other authority may consider, in addition to all other
- 18 logically relevant factors, the following:
- (1) Statements by an owner or by anyone in control of the objectconcerning its use.
- (2) Prior convictions, if any, of an owner, or of anyone in control
  of the object, under any state or federal law relating to any
  controlled substance.
- (3) Direct or circumstantial evidence of the intent of an owner,
  or of anyone in control of the object, to deliver it to persons whom
  they know, or should reasonably know, intend to use the object to
  facilitate a violation of this section. The innocence of an owner,
  or of anyone in control of the object, as to a direct violation of this
  section shall not prevent a finding that the object is intended for
  use, or designed for use, as drug paraphernalia.
- 31 (4) Instructions, oral or written, provided with the object
  32 concerning its use.
- 32 concerning its use.
  33 (5) Descriptive materials, accompanying the object which that
  34 explain or depict its use.
- 35 (6) National and local advertising concerning its use.
- 36 (7) The manner in which the object is displayed for sale.
- 37 (8) Whether the owner or anyone in control of the object is a
- 38 legitimate supplier of like or related items to the community, such
- 39 as a licensed distributor or dealer of tobacco products.

1 (9) The existence and scope of legitimate uses for the object in 2 the community.

3 (10) Expert testimony concerning its use.

4 (11) If the possession of the object is related to, or is a result 5 of, a substance use disorder treatment or recovery program, harm 6 reduction program, syringe exchange program or syringe services 7 program, or consistent with best clinical practices in alignment 8 with the Substance Abuse and Mental Health Services 9 Administration and the American Society of Addiction Medicine.

10 (f) This section shall not apply to any of the following:

(1) Any pharmacist or other authorized person who sells or
furnishes drug paraphernalia described in paragraph (10) of
subdivision (d) upon the prescription of a physician, dentist,
podiatrist, or veterinarian.

(2) Any physician, dentist, podiatrist, or veterinarian who
furnishes or prescribes drug paraphernalia described in paragraph
(10) of subdivision (d) to a patient.

(3) Any manufacturer, wholesaler, or retailer licensed by the
California State Board of Pharmacy to sell or transfer drug
paraphernalia described in paragraph (10) of subdivision (d).

21 (4) A person who is at risk of overdose, a family member, friend,

22 or other person in a position to assist a person at risk of overdose,

23 in order to reduce the spread of HIV, viral hepatitis, and other

bloodborne infections among the intravenous drug user populationwithin California.

(g) Notwithstanding paragraph (12) of subdivision (a), "drug
paraphernalia" "Drug paraphernalia" does not include any testing
equipment designed, marketed, intended to be used, or used, to
test a substance for the presence of contaminants, toxic substances,
hazardous compounds, or other adulterants, or controlled
substances that include, without limitation, fentanyl, ketamine,
gamma hydroxybutyric acid, or any analog of fentanyl.

33 (h) Notwithstanding any other law, including Section 11374, 34 violation of this section shall not constitute a criminal offense, but 35 operation of a business in violation of the provisions of this section 36 shall be grounds for revocation or nonrenewal of any license, 37 permit, or other entitlement previously issued by a city, county, 38 or city and county for the privilege of engaging in such business 39 and shall be grounds for denial of any future license, permit, or 40 other entitlement authorizing the conduct of such business or any

1 other business, if the business includes the sale of drug 2 paraphernalia.

3 SEC. 6. Section 11364.7 of the Health and Safety Code is 4 amended to read:

5 11364.7. (a) (1) Except as authorized by law, a person who delivers, furnishes, or transfers, possesses with intent to deliver, 6 7 furnish, or transfer, or manufactures with the intent to deliver, 8 furnish, or transfer, drug paraphernalia, knowing, or under 9 circumstances where one reasonably should know, that it will be used to plant, propagate, cultivate, grow, harvest, compound, 10 convert, produce, process, prepare, test, analyze, pack, repack, 11 12 store, contain, conceal, inject, ingest, inhale, or otherwise introduce 13 into the human body a controlled substance, except as provided 14 in subdivision (b), in violation of this division, is guilty of a 15 misdemeanor.

16 (2) A public entity, its agents, or employees shall not be subject 17 to criminal prosecution for distribution of hypodermic needles or syringes or any materials deemed by a local or state health 18 department to be necessary to prevent the spread of communicable 19 diseases, or to prevent drug overdose, injury, or disability to 20 21 participants in clean needle and syringe exchange projects 22 authorized by the public entity pursuant to Chapter 18 23 (commencing with Section 121349) of Part 4 of Division 105.

(b) Except as authorized by law, a person who manufactures 24 25 with intent to deliver, furnish, or transfer drug paraphernalia 26 knowing, or under circumstances where one reasonably should 27 know, that it will be used to plant, propagate, cultivate, grow, 28 harvest, manufacture, compound, convert, produce, process, 29 prepare, test, analyze, pack, repack, store, contain, conceal, inject, 30 ingest, inhale, or otherwise introduce into the human body cocaine, 31 cocaine base, heroin, phencyclidine, or methamphetamine in 32 violation of this division shall be punished by imprisonment in a 33 county jail for not more than one year, or in the state prison.

(c) Except as authorized by law, a person, 18 years of age or over, who violates subdivision (a) by delivering, furnishing, or transferring drug paraphernalia to a person under 18 years of age who is at least three years their junior, or who, upon the grounds of a public or private elementary, vocational, junior high, or high school, possesses a hypodermic needle, as defined in Section 11014.5, with the intent to deliver, furnish, or transfer the

1 hypodermic needle, knowing, or under circumstances where one

2 reasonably should know, that it will be used by a person under 18

3 years of age to inject into the human body a controlled substance,

4 is guilty of a misdemeanor and shall be punished by imprisonment

5 in a county jail for not more than one year, by a fine of not more

6 than one thousand dollars (\$1,000), or by both that imprisonment7 and fine.

8 (d) The violation, or the causing or the permitting of a violation,
9 of subdivision (a), (b), or (c) by a holder of a business or liquor
10 license issued by a city, county, or city and county, or by the State

of California, and in the course of the licensee's business shall be grounds for the revocation of that license.

(e) All drug paraphernalia defined in Section 11014.5 is subject
to forfeiture and may be seized by any peace officer pursuant to
Section 11471 unless its distribution has been authorized pursuant
to subdivision (a).

17 (f) If any provision of this section or the application thereof to 18 any person or circumstance is held invalid, it is the intent of the 19 Legislature that the invalidity shall not affect other provisions or 20 applications of this section that can be given effect without the 21 invalid provision or application and to this end the provisions of 22 this section are severable.

23 SEC. 7. Section 11372.7 of the Health and Safety Code is 24 amended to read:

25 11372.7. (a) Except as otherwise provided in subdivision (b) 26 or (e), each person who is convicted of a violation of this chapter 27 shall pay a drug program fee in an amount not to exceed one 28 hundred fifty dollars (\$150) for each separate offense. The court 29 shall increase the total fine, if necessary, to include this increment, 30 which shall be in addition to any other penalty prescribed by law. 31 (b) The court shall determine whether or not the person who is 32 convicted of a violation of this chapter has the ability to pay a drug 33 program fee. If the court determines that the person has the ability 34 to pay, the court may set the amount to be paid and order the person to pay that sum to the county in a manner that the court believes 35 36 is reasonable and compatible with the person's financial ability. 37 In its determination of whether a person has the ability to pay, the 38 court shall take into account the amount of any fine imposed upon

39 that person and any amount that person has been ordered to pay

40 in restitution. If the court determines that the person does not have

1 the ability to pay a drug program fee, the person shall not be 2 required to pay a drug program fee.

3 (c) The county treasurer shall maintain a drug program fund. 4 For every drug program fee assessed and collected pursuant to 5 subdivisions (a) and (b), an amount equal to this assessment shall be deposited into the fund for every conviction pursuant to this 6 7 chapter, in addition to fines, forfeitures, and other moneys-which 8 that are transmitted by the courts to the county treasurer pursuant 9 to Sections 11372.5 and 11502. These deposits shall be made prior 10 to any transfer pursuant to Section 11502. Amounts deposited in 11 the drug program fund shall be allocated by the administrator of 12 the county's drug program to drug abuse programs in the schools 13 and the community, subject to the approval of the board of 14 supervisors, as follows:

(1) The moneys in the fund shall be allocated through the
planning process established pursuant to Sections 11983, 11983.1,
11983.2, and 11983.3.

18 (2) A minimum of 33 percent of the fund shall be allocated to 19 primary prevention programs in the schools and the community. Primary prevention programs developed and implemented under 20 21 this article shall emphasize cooperation in planning and program 22 implementation among schools and community drug abuse 23 agencies, and shall demonstrate coordination through an 24 interagency agreement among county offices of education, school 25 districts, and the county drug program administrator. These primary 26 prevention programs may include:

27 (A) School- and classroom-oriented programs, including, but 28 not limited to, programs designed to encourage sound 29 decisionmaking, an awareness of values, an awareness of drugs 30 and their effects, enhanced self-esteem, social and practical skills 31 that will assist students toward maturity, enhanced or improved 32 school climate and relationships among all school personnel and 33 students, and furtherance of cooperative efforts of school- and 34 community-based personnel.

(B) School- or community-based nonclassroom alternative
programs, or both, including, but not limited to, positive peer group
programs, programs involving youth and adults in constructive
activities designed as alternatives to drug use, and programs for
special target groups, such as women, ethnic minorities, and other
high-risk, high-need populations.

1 (C) Family-oriented programs, including, but not limited to, 2 programs aimed at improving family relationships and involving parents constructively in the education and nurturing of their 3 4 children, as well as in specific activities aimed at preventing drug 5 abuse. substance use disorders. 6 (D) Primary prevention activities identified by the Substance 7 Abuse and Mental Health Services Administration in the course 8 of administering the Substance Use Prevention, Treatment, and 9 Recovery Services Block Grant, authorized by Section 1921 of 10 Subparts II and III of Part B of Title XIX of the Public Health Service Act. 11 12 (d) Moneys deposited into a county drug program fund pursuant 13 to this section shall supplement, and shall not supplant, any local funds made available to support the county's drug abuse prevention 14 15 and treatment efforts. 16 (e) This section shall not apply to any person convicted of a 17 violation of subdivision (b) of Section 11357 of the Health and 18 Safety Code. SEC. 8. The heading of Division 10.5 (commencing with 19 20 Section 11750) of the Health and Safety Code is amended to read: 21 22 DIVISION 10.5. ALCOHOL AND OTHER DRUG 23 PROGRAMS 24 25 SEC. 9. Section 11832 of the Health and Safety Code is 26 amended to read: 27 11832. (a) The department has the sole authority in state 28 government to certify alcohol or other drug programs. 29 (b) In administering this chapter, the department shall issue 30 certifications for a period of two years to those alcohol or other 31 drug programs that meet the requirements set forth in this chapter. 32 (c) The department shall, on or before July 1, 2026, offer a 33 combined application for entities seeking certification as an alcohol 34 or other drug program to simultaneously apply for certification 35 to provide incidental medical services as defined in Section 36 11834.026. 37 (d) An additional fee shall not be charged for the combined 38 application described in subdivision (c) in excess of the charge 39 authorized in Sections 11832.1 and 11832.4.

1 (e) The department shall post on its internet website a timeline 2 with the relative dates of key milestones in the permit application 3 review process and the average processing times for the 4 department of each stage of key milestones in the permit 5 application review process. The department shall note on its 6 internet website that these times are estimates, and shall update 7 the times as necessary.

8 (f) The department shall provide written notices of estimated 9 dates of key milestones in the permit application review process 10 to the applicant and the local continuum of care.

11 (g) Key milestones in the permit application review process 12 shall include, but are not limited to, all of the following:

(1) Initial indication of whether the application is complete orincomplete within 45 working days of receipt of the application.

15 (A) If the application is incomplete, the department shall specify 16 the information or documentation that is missing in a notice to the

17 applicant within 45 working days of receipt of the application.

18 (B) The applicant shall have 60 working days from the date of 19 the notification to provide the missing information or 20 documentation.

(2) Indication of whether the application for certification to
 provide incidental medical services is complete or incomplete
 within 45 working days of receipt of the application.

(A) If the application for certification to provide incidental
medical services is incomplete, the department shall specify the
information or documentation that is missing in a notice to the
applicant within 45 working days of receipt of the application.

28 (B) The applicant shall have 60 working days from the date of 29 the notification to provide the missing information or 30 documentation.

31 (3) Issuance of a certification to provide incidental medical
32 services or a written notification of denial of certification within
33 120 working days of determining that the application is complete.

34 (4) Issuance of a license by certified mail or a written
 35 notification of denial of licensure within 120 working days of

36 *determining that the application is complete.* 

37 (*h*) On or before December 31, 2026, the department shall post

38 to its internet website the average processing times, as described

39 in subdivision (e), for each application under review by the

40 *department*.

1 (i) Any necessary rules and regulations for the purpose of 2 implementing this section may be adopted as emergency 3 regulations in accordance with the Administrative Procedure Act 4 (Chapter 3.5 (commencing with Section 11340) of Part 1 of 5 Division 3 of Title 2 of the Government Code). The adoption of 6 emergency regulations pursuant to this section shall be deemed to be an emergency and necessary for the immediate preservation 7 8 of the public peace, health and safety, or general welfare.

9 SEC. 10. Section 11834.026 of the Health and Safety Code is 10 amended to read:

11834.026. (a) As used in this section, "incidental medical 11 12 services" means services that are in compliance with the 13 community standard of practice and are not required to be 14 performed in a licensed clinic or licensed health facility, as defined 15 by Section 1200 or 1250, respectively, to address medical issues associated with either detoxification from alcohol or other drugs 16 17 or the provision of alcohol or other drug recovery or treatment 18 services, including all of the following categories of services that 19 the department shall further define by regulation:

20 (1) Obtaining medical histories.

(2) Monitoring health status to determine whether the health
 status warrants transfer of the patient in order to receive urgent or
 emergent care.

24 (3) Testing associated with detoxification from alcohol or *other*25 drugs.

26 (4) Providing alcohol or other drug recovery or treatment27 services.

28 (5) Overseeing patient self-administered medications.

29 (6) Treating substance use disorders, including detoxification.

30 (b) Incidental medical services do not include the provision of

31 general primary medical care.

32 (c) Notwithstanding any other law, a licensed alcohol or other 33 drug recovery or treatment facility may permit incidental medical 34 services to be provided to a resident at the facility premises by, or under the supervision of, one or more physicians and surgeons 35 36 licensed by the Medical Board of California or the Osteopathic 37 Medical Board who are knowledgeable about addiction medicine, 38 or one or more other health care practitioners acting within the 39 scope of practice of their license and under the direction of a

1 physician and surgeon, and who are also knowledgeable about 2 addiction medicine, if all of the following conditions are met:

3 (1) The facility, in the judgment of the department, has the 4 ability to comply with the requirements of this chapter and all other 5 applicable laws and regulations to meet the needs of a resident receiving incidental medical services pursuant to this chapter. The 6 7 department shall specify in regulations the minimum requirements 8 that a facility shall meet in order to be approved to permit the 9 provision of incidental medical services on its premises. The license 10 of a facility approved to permit the provision of incidental medical 11 services shall reflect that those services are permitted at the facility 12 premises.

13 (2) The physician and surgeon and any other health care 14 practitioner has signed an acknowledgment on a form provided 15 by the department that they have been advised of and understand the statutory and regulatory limitations on the services that may 16 17 legally be provided at a licensed alcohol or other drug recovery or 18 treatment facility and the statutory and regulatory requirements 19 and limitations for the physician and surgeon or other health care practitioner and for the facility, related to providing incidental 20 21 medical services. The licensee shall maintain a copy of the signed 22 form at the facility for a physician and surgeon or other health care 23 practitioner providing incidental medical services at the facility 24 premises.

(3) A physician and surgeon or other health care practitioner
shall assess a resident, prior to that resident receiving incidental
medical services, to determine whether it is medically appropriate
for that resident to receive these services at the premises of the
licensed facility. A copy of the form provided by the department
shall be signed by the physician and surgeon and maintained in
the resident's file at the facility.

32 (4) The resident has signed an admission agreement. The

(A) *The* admission agreement, at a minimum, shall describe the
incidental medical services that the facility may permit to be
provided and shall state that the permitted incidental medical
services will be provided by, or under the supervision of, a
physician and surgeon. The

38 (*B*) *The admission agreement shall not require a person to have* 39 *been abstinent, to not be intoxicated, or to otherwise not be under* 

the influence in order to be admitted into care, be considered for
 treatment, or continue treatment.

3 (*C*) *The* department shall specify in regulations, at a minimum, 4 the content and manner of providing the admission agreement, and 5 any other information that the department deems appropriate. The 6 facility shall maintain a copy of the signed admission agreement 7 in the resident's file.

8 (5) Once incidental medical services are initiated for a resident, 9 the physician and surgeon and facility shall monitor the resident 10 to ensure that the resident remains appropriate to receive those 11 services. If the physician and surgeon determines that a change in 12 the resident's medical condition requires other medical services 13 or that a higher level of care is required, the facility shall 14 immediately arrange for the other medical services or higher level 15 of care, as appropriate.

16 (6) The facility maintains in its files a copy of the relevant 17 professional license or other written evidence of licensure to 18 practice medicine or perform medical services in the state for the 19 physician and surgeon and any other health care practitioner 20 providing incidental medical services at the facility.

(d) The department is not required to evaluate or have any
responsibility or liability with respect to evaluating the incidental
medical services provided by a physician and surgeon or other
health care practitioner at a licensed facility. This section does not
limit the department's ability to report suspected misconduct by
a physician and surgeon or other health care practitioner to the
appropriate licensing entity or to law enforcement.

28 (e) A facility licensed and approved by the department to allow 29 provision of incidental medical services shall not by offering 30 approved incidental medical services be deemed a clinic or health 31 facility within the meaning of Section 1200 or 1250, respectively. 32 (f) Other than incidental medical services permitted to be 33 provided or any urgent or emergent care required in the case of a 34 life-threatening emergency, including the administration of naloxone hydrochloride, or any other opioid antagonist that is 35 36 approved by the United States Food and Drug Administration for 37 treatment of an opioid overdose, this section does not authorize 38 the provision at the premises of the facility of any medical or health 39 care services or any other services that require a higher level of

- 1 care than the care that may be provided within a licensed alcohol
- 2 or other drug recovery or treatment facility.
- 3 (g) This section does not require a residential treatment facility
- 4 licensed by the department to provide incidental medical services5 or any services not otherwise permitted by law.
- 6 (h) (1) On or before July 1, 2024, the department shall adopt
- 7 regulations to implement this section in accordance with the 8 Administrative Procedure Act (Chapter 3.5 (commencing with
- 9 Section 11340) of Part 1 of Division 3 of Title 2 of the Government
- 10 Code).
- (2) Notwithstanding the rulemaking provisions of the
  Administrative Procedure Act, the department may, if it deems
  appropriate, implement, interpret, or make specific this section by
  means of provider bulletins, written guidelines, or similar
- 15 instructions from the department until regulations are adopted.
- 16 SEC. 11. Section 11834.26 of the Health and Safety Code is 17 amended to read:
- 18 11834.26. (a) The licensee shall provide at least one of the 19 following *alcohol- or other drug-related* nonmedical services:
- 20 (1) Recovery services.
- 21 (2) Treatment services.
- 22 (3) Detoxification services.
- (b) The department shall adopt regulations requiring records
  and procedures that are appropriate for each of the services
  specified in subdivision (a). The records and procedures may
  include all of the following:
- 27 (1) Admission criteria.
- 28 (2) Intake process.
- 29 (3) Assessments.
- 30 (4) Recovery, treatment, or detoxification planning.
- 31 (5) Referral.
- 32 (6) Documentation of provision of recovery, treatment, or 33 detoxification services.
- 34 (7) Discharge and continuing care planning.
- 35 (8) Indicators of recovery, treatment, or detoxification outcomes.
- 36 (c) A licensee shall not deny admission to any individual based
- 37 solely on-the either of the following:
- 38 (1) The individual having a valid prescription from a licensed
- 39 health care professional for a medication approved by the federal
- 40 Food and Drug Administration for the purpose of narcotic
  - 99

1 replacement treatment or medication-assisted treatment of 2 substance use disorders.

3 (2) The individual having consumed, used, or otherwise been 4 under the influence of alcohol or other drugs, as these 5 circumstances represent symptoms of the condition of substance 6 use disorders.

7 (d) A licensee shall develop a plan to address when a resident
8 relapses, including when a resident is on the licensed premises
9 after consuming alcohol or using illegal using alcohol or other
10 drugs. The

(1) The plan shall include details of how the treatment stay and treatment plan of the resident will be adjusted to address the relapse episode and how the resident will be treated and supervised while under the influence of alcohol or-illegal other drugs, as well as discharge and continuing care planning, including when a licensee determines that a resident requires services beyond the scope of the licensee. This

(2) This subdivision does not require a licensee to discharge a
resident. resident, as relapse, lapses, and momentary reengagement
with alcohol or other drugs are symptoms of the condition of
substance use disorders.

(3) In developing a plan pursuant to this subdivision, the
licensee shall consider options to avoid disconnection of the
resident from treatment and shall prioritize the individual
maintaining some level of connection to treatment, rather than a
complete disconnection from treatment.

(e) The department shall have the authority to implement
subdivisions (d) and (f) by bulletin or all-county or all-provider
letter, after stakeholder input, until regulations are promulgated.
The department shall promulgate regulations to implement
subdivisions (d) and (f) no later than July 1, 2024.

32 (f) (1) A licensee shall, at all times, maintain at least two 33 unexpired doses of naloxone hydrochloride, or any other opioid 34 antagonist that is approved by the United States Food and Drug 35 Administration for treatment of an opioid overdose, on the premises and shall, at all times, have at least one staff member on the 36 37 premises who knows the specific location of the naloxone 38 hydrochloride, or other opioid antagonist that is approved by the 39 United States Food and Drug Administration for treatment of an 40 opioid overdose, and who has been trained on the administration

of naloxone hydrochloride, or the other opioid antagonist that is 1 2 approved by the United States Food and Drug Administration for 3 treatment of an opioid overdose, in accordance with the training 4 requirements set forth by the department. Proof of completion of 5 training on the administration of naloxone hydrochloride, or other opioid antagonist that is approved by the United States Food and 6 7 Drug Administration for treatment of an opioid overdose, shall be 8 documented in the staff member's individual personnel file. 9 (2) A trained staff member shall not be liable for damages in a 10 civil action or subject to criminal prosecution for the administration, in good faith, of naloxone hydrochloride, or any 11 12 other opioid antagonist that is approved by the United States Food 13 and Drug Administration for treatment of an opioid overdose, to 14 a person appearing to experience an opioid-related overdose. This 15 paragraph shall not apply in a case where the person who renders emergency care treatment by the use of naloxone hydrochloride, 16 17 or any other opioid antagonist that is approved by the United States 18 Food and Drug Administration for treatment of an opioid overdose, 19 acts with gross negligence or engages in willful and wanton 20 misconduct. 21 (g) In the development of regulations implementing this section, 22 the written record requirements shall be modified or adapted for 23 social model programs. SEC. 12. The heading of Division 10.7 (commencing with 24 25 Section 11999) of the Health and Safety Code is repealed. 26 27 **DIVISION 10.7. ILLEGAL USE OF DRUGS AND ALCOHOL** 28 29 SEC. 13. The heading of Division 10.7 (commencing with 30 Section 11999) is added to the Health and Safety Code, to read: 31 32 **DIVISION 10.7. SUBSTANCE USE DISORDER** 33 PREVENTION, TREATMENT, AND RECOVERY PROGRAMS 34 35 SEC. 14. Section 11999 of the Health and Safety Code is 36 amended to read: 37 11999. The Legislature finds and declares all of the following: 38 (a) The Legislature has established various drug- and 39 alcohol-related programs which provide for education, prevention, 40 intervention, treatment, or enforcement.

(b) The Legislature has classified certain substances as
 controlled substances and has defined the lawful and unlawful use
 of controlled substances which are commonly referred to as, but
 not limited to, anabolic steroids, marijuana, and cocaine.

5 (c) The Legislature has classified certain substances as imitation 6 controlled substances which are commonly referred to as, but not 7 limited to, designer drugs.

8 (d) The Legislature has determined that the possession with the 9 intent to be under the influence, or being under the influence of 10 toluene, or any substance or material containing toluene, or any 11 substance with similar toxic qualities, is unlawful. Some substances 12 or materials containing toluene, or substances with similar toxic 13 qualities are commonly referred to, but not limited to, inhalants 14 such as cement, glue, and paint thinner.

(e) The Legislature has determined that the purchase, possession,or use of alcohol by persons under 21 years of age is unlawful.

17 (f) Public and private agencies that provide information 18 pertaining to the drug- and alcohol-related programs provide mixed 19 messages and misinformation relating to the unlawful use of drugs 20 and alcohol. It is the intent of the Legislature that the messages 21 and information provided by the drug and alcohol programs 22 promote no unlawful use of any drugs or alcohol. Mixed messages 23 mean communications discussing how to use or when to use 24 unlawful drugs or alcohol.

(g) Any material, curricula, teachings, or promotion of
 responsible use, if the use is unlawful, of drugs or alcohol is
 inconsistent with the law.

(h) The "no unlawful use" message applies to all drug and
alcohol programs for the people of the State of California. These
materials are to teach and promote that any unlawful use of drugs
and alcohol is illegal and dangerous.

(f) Substance use disorder should be viewed and treated as a
health problem, as well as a public safety problem as described
in Section 11760.5.

(g) Comprehensive prevention and treatment services for
individuals experiencing or recovering from substance use
disorders must be medically accurate, culturally congruent, and
evidence based.

39 (h) Naloxone, a life-saving opioid antagonist medication used 40 to reverse an opioid overdose, including heroin, fentanyl, and

1 prescription opioid medications, is safe and easy to use, works

2 almost immediately, and is not addictive. Naloxone has very few

3 negative effects, and has no effect if opioids are not in a person's

4 system.

5 (i) With the establishment of the Naloxone Distribution Program

6 and the United States Food and Drug Administration's approval

7 for over-the-counter, nonprescription use of naloxone for the

8 reversal of an opioid overdose, the Legislature further finds that

9 carrying naloxone provides an extra layer of protection for those

10 at a higher risk for overdose. Although most professional first

11 responders and emergency departments carry naloxone, they may

12 not arrive in time to reverse an opioid overdose. Anyone can carry

naloxone, give it to someone having an overdose, and potentially
save a life. Bystanders such as friends, family, non-health care

14 save a tife. Bystanders such as friends, family, non-neutin care 15 providers, and persons who use drugs can reverse an opioid

16 *overdose with naloxone.* 

17 SEC. 15. Section 11999.1 of the Health and Safety Code is 18 amended to read:

19 11999.1. For the purpose of this division, the following20 definitions apply:

21 (a) "Drug" means all of the following:

(1) Any controlled substance as defined in Division 10(commencing with Section 11000).

(2) Any imitation controlled substance as defined in Chapter 1(commencing with Section 11670) of Division 10.1.

(3) Toluene or any substance or material containing toluene or
any substance with similar toxic qualities as set forth in Sections
380 and 381 of the Penal Code.

29 (b) "Drug- or alcohol-related program" means any program

30 designed to reduce the unlawful use of, or assist those who engage 31 in the unlawful use of, drugs or alcohol, assist persons with

32 substance use disorders whether through education, prevention,

33 intervention, treatment, enforcement, or other means.

34 (c) "Local agency" shall include, but is not limited to, a county,35 a city, a city and county, and school district.

36 (d) "State agency" shall include the State Department of Health
 37 Care Services, the State Department of Education, the Department

38 of Justice, the Office of Criminal Justice Planning, and the Office

39 of Traffic Safety. Any other state agency or department may

40 comply with this division.

1 SEC. 16. Section 11999.2 of the Health and Safety Code is 2 repealed. 3 11999.2. (a) Notwithstanding any other provision of law, 4 commencing July 1, 1990, no state funds shall be encumbered by 5 a state agency for allocation to any entity, whether public or 6 private, for a drug- or alcohol-related program, unless the drug-7 or alcohol-related program contains a component that clearly 8 explains in written materials that there shall be no unlawful use of 9 drugs or alcohol. No aspect of a drug- or alcohol-related program 10 shall include any message on the responsible use, if the use is 11 unlawful, of drugs or alcohol. 12 (b) All aspects of a drug- or alcohol-related program shall be 13 consistent with the "no unlawful use" message, including, but not 14 limited to, program standards, curricula, materials, and teachings. 15 These materials and programs may include information regarding 16 the health hazards of use of illegal drugs and alcohol, concepts 17 promoting the well-being of the whole person, risk reduction, the 18 addictive personality, development of positive self-esteem, 19 productive decisionmaking skills, and other preventive concepts consistent with the "no unlawful use" of drugs and alcohol 20 21 message. 22 (c) The "no unlawful use" of drugs and alcohol message 23 contained in drug- or alcohol-related programs shall apply to the 24 use of drugs and alcohol prohibited by law. 25 (d) This section does not apply to any program funded by the 26 state that provides education and prevention outreach to intravenous 27 drug users with AIDS or AIDS-related conditions, or persons at 28 risk of HIV-infection through intravenous drug use. 29 SEC. 17. Section 11999.2 is added to the Health and Safety 30 Code, to read: 31 11999.2. (a) Notwithstanding any other law, an alcohol or 32 other drug-related program shall be consistent with best clinical 33 practices in alignment with the Substance Abuse and Mental Health 34 Services Administration and the American Society of Addiction Medicine in order for state funds to be encumbered by a state 35 36 agency for allocation to any entity, whether public or private. 37 (b) This section includes any program funded by the state that 38 provides education and prevention outreach to persons at risk of

39 HIV-infection, viral hepatis, or other bloodborne infections through

40 intravenous drug use, or an opioid overdose prevention and

1	treatment training program as defined in paragraph (2) of subdivision (a) of Section 1714.22 of the Civil Code.
2 3	SEC. 18. The heading of Part 4 (commencing with Section
4	120775) of Division 105 of the Health and Safety Code is amended
5	to read:
6	
7	PART 4. HUMAN IMMUNODEFICIENCY VIRUS (HIV) AND
8	OTHER BLOODBORNE DISEASES
9	
10	SEC. 19. The heading of Chapter 1.5 (commencing with
11	Section 120780) of Part 4 of Division 105 of the Health and Safety
12	Code is amended to read:
13	
14	CHAPTER 1.5. STATE HIV, VIRAL HEPATITIS, AND OTHER
15	<b>Bloodborne Disease</b> Prevention and Education Funds
16 17	SEC 20 Section 120780.1 of the Health and Sefety Code is
17	SEC. 20. Section 120780.1 of the Health and Safety Code is amended to read:
19	120780.1. A public entity that receives General Fund money
20	from the State Department of Public Health for <del>-HIV</del> <i>HIV</i> , <i>viral</i>
20	hepatitis, and other bloodborne disease prevention and education
22	may use that money to support clean needle and syringe exchange
23	programs authorized pursuant to existing law. The money may be
24	used for, but is not limited to, the purchase of sterile hypodermic
25	needles and syringes as part of a clean needle and syringe exchange
26	program only if all of the following conditions are met: in
27	alignment with primary prevention activities as determined by the
28	Substance Abuse and Mental Health Services Administration for
29	the purposes of the administration of the Substance Use Prevention,
30	Treatment, and Recovery Services Block Grant, authorized by
31	Section 1921 of Subparts II and III of Part B of Title XIX of the
32	Public Health Service Act.
33	(a) The General Fund money used for purchasing the sterile
34	hypodermic needles and syringes does not supplant any other
35	public or private funds or other resources for this purpose.
36	(b) The amount of the General Fund money used for purchasing the starile hypothermia needles and suringes does not exceed 7.5
37 38	the sterile hypodermic needles and syringes does not exceed 7.5
30	percent of the total amount of the General Fund money received

39 by the public entity for HIV prevention and education.

1 (c) Each dollar of General Fund money used for purchasing the

2 sterile hypodermic needles and syringes is matched by forty-three
 3 cents (\$0.43) of moneys from nonstate public funds or private
 4 funds.

5 (d) The allocation of General Fund money for the purchase of

6 sterile hypodermic needles and syringes is based upon

7 epidemiological data as reported by the health jurisdiction in its

8 local HIV prevention plan submitted to the Office of AIDS within

9 the department.

10 SEC. 21. Section 120780.2 of the Health and Safety Code is 11 amended to read:

12 120780.2. In order to reduce the spread of HIV, hepatitis C, 13 viral hepatitis, and other potentially deadly bloodborne pathogens, the State Department of Public Health may purchase sterile 14 15 hypodermic needles and syringes, and other supplies, for distribution to syringe exchange programs authorized pursuant to 16 17 law and support any costs associated with distribution of supplies. 18 Supplies provided to programs, including those administered by 19 local health departments, are not subject to the formulas and limits 20 of Section 120780.1. 21 SEC. 22. Section 120780.5 of the Health and Safety Code is

21 SEC. 22. Section 120780.5 of the Health and Safety Code is 22 amended to read:

23 120780.5. (a) Upon an appropriation in the annual Budget 24 Act, the State Department of Public Health shall award funding, 25 on a competitive basis, to community-based organizations or local 26 health jurisdictions to provide comprehensive HIV prevention and 27 control activities for the most vulnerable and underserved 28 individuals living with, or at high risk for, HIV infection. 29 Applicants may include individual community-based organizations 30 and local health jurisdictions, as well as collaborations between 31 community-based organizations and local health jurisdictions.

32 (b) Entities located in any county are eligible to receive grant33 funding.

34 (c) Comprehensive <u>HIV</u> *HIV, viral hepatitis, and other* 35 *bloodborne disease* prevention and control activities may include,

36 but are not limited to, any of the following:

37 (1) HIV testing, including the purchase of HIV test kits.

38 (2) Linkage to and retention in care for people living with HIV.

39 (3) Pre-exposure prophylaxis (PrEP)-related and post-exposure

40 prophylaxis (PEP)-related activities.

1 (4) Syringe services programs.

(d) The department shall determine the funding levels of each 2 3 award based on scope and geographic area. Priority for grants shall 4 be given to community-based organizations or local health 5 jurisdictions that, through their applications, demonstrate expertise, history, and credibility at working successfully in engaging the 6 7 most vulnerable and underserved individuals living with, or at high 8 risk for, HIV infection. HIV, viral hepatitis, or other bloodborne 9 infections. 10 (e) Funds shall be allocated in a manner that balances the need

10 (c) Funds shall be anocated in a manner that balances the need 11 to spread funding to as many local health jurisdictions and 12 community-based organizations as possible and the need to provide 13 meaningful activities to each recipient. Not less than 50 percent 14 of the funds allocated shall be provided to community-based 15 organizations, for purposes consistent with this section.

(f) The department shall determine the application process,selection criteria, and any reporting requirements for the grant,consistent with this section.

(g) The department shall develop measures for each local health
 jurisdiction and community-based organization funded pursuant
 to this section to demonstrate accountability.

(h) This section shall be operative only if funds are explicitlyappropriated in the annual Budget Act specifically for purposesof this section.

25 SEC. 23. Section 121349 of the Health and Safety Code is 26 amended to read:

27 121349. (a) The Legislature finds and declares that scientific 28 data from needle exchange programs in the United States and in Europe have shown that the exchange of used hypodermic needles 29 30 and syringes for clean hypodermic needles and syringes does not 31 increase drug use in the population, can serve as an important 32 bridge to treatment and recovery from drug abuse, substance use 33 disorder, and can curtail the spread of human immunodeficiency 34 virus (HIV) infection among the intravenous drug user population. 35 (b) In order to reduce the spread of HIV infection and bloodborne hepatitis among the intravenous drug user population 36 within California, the Legislature hereby authorizes a clean needle 37 38 and syringe exchange project pursuant to this chapter in any city, 39 county, or city and county upon the action of a county board of 40 supervisors and the local health officer or health commission of

1 that county, or upon the action of the city council, the mayor, and

2 the local health officer of a city with a health department, or upon

3 the action of the city council and the mayor of a city without a

4 health department.

5 (c) In order to reduce the spread of HIV infection, viral hepatitis, 6 and other potentially deadly bloodborne infections, the State

7 Department of Public Health may, notwithstanding any other law,

8 authorize entities that provide services set forth in paragraph (1)

9 of subdivision (d), and that have sufficient staff and capacity to 10 provide the services described in Section 121349.1, as determined

11 by the department, to apply for authorization under this chapter to

12 provide hypodermic needle and syringe exchange services

consistent with state standards in any location where the departmentdetermines that the conditions exist for the rapid spread of HIV,

15 viral hepatitis, or any other potentially deadly or disabling

16 infections that are spread through the sharing of used hypodermic

17 needles and syringes. Authorization shall be made after

18 consultation with the local health officer and local law enforcement

19 leadership, and after a period of public comment, as described in

20 subdivision (e). In making the determination, the department shall

balance the concerns of law enforcement with the public healthbenefits. The authorization shall not be for more than two years.

23 Before the end of the two-year period, the department may

reauthorize the program in consultation with the local health officer

25 and local law enforcement leadership.

(d) In order for an entity to be authorized to conduct a project
pursuant to this chapter, its application to the department shall
demonstrate that the entity complies with all of the following

## 29 minimum standards:

30 (1) The entity provides, directly or through referral, all of the 31 following services:

- 32 (A) <del>Drug abuse</del> *Substance use disorder* treatment services.
- 33 (B) HIV or hepatitis screening.
- 34 (C) Hepatitis A and hepatitis B vaccination.
- 35 (D) Screening for sexually transmitted infections.
- 36 (E) Housing services for the homeless, for victims of domestic 37 violence, or other similar housing services.
- 38 (F) Services related to provision of education and materials for

39 the reduction of sexual risk behaviors, including, but not limited

40 to, the distribution of condoms.

1 (2) The entity has the capacity to commence needle and syringe 2 exchange services within three months of authorization.

3 (3) The entity has adequate funding can demonstrate that it has

4 *the ability* to do all of the following at reasonably projected 5 program participation levels: levels within three months of 6 *authorization*:

7 (A) Provide needles and syringe exchange services for all of its8 participants.

9 (B) Provide HIV and viral hepatitis prevention education 10 services for all of its participants.

(C) Provide for the safe recovery and disposal of used syringesand sharps waste from all of its participants.

(4) The entity has the capacity, and an established plan, to collect
evaluative data in order to assess program impact, including, but
not limited to, all of the following:

16 (A) The total number of persons served.

17 (B) The total number of needles and syringes distributed, 18 recovered, and disposed of.

19 (C) The total numbers and types of referrals to drug treatment20 and other services.

(e) If the application is provisionally deemed appropriate by thedepartment, the department shall, at least 45 days prior to approval

of the application, provide for a period of public comment as follows:

(1) Post on the department's internet website the name of theapplicant, the nature of the services, and the location where theapplying entity will provide the services.

(2) Send a written and an email notice to the local health officerof the affected jurisdiction.

30 (3) Send a written and an email notice to the chief of police, the
31 sheriff, or both, as appropriate, of the jurisdictions in which the
32 program will operate.

33 (f) The department shall establish and maintain on its internet

34 website the address and contact information of programs providing

35 hypodermic needle and syringe exchange services pursuant to this36 chapter.

37 (g) The authorization provided under this section is only for a

38 clean needle and syringe exchange project as described in Section

39 121349.1.

(h) (1) Needle and syringe exchange services application
submissions, authorizations, and operations performed pursuant
to this chapter shall be exempt from review under the California
Environmental Quality Act, Division 13 (commencing with Section
21000) of the Public Resources Code.

6 (2) This subdivision is intended to be declaratory of existing 7 law.

8 (i) If the department, in its discretion, determines that a state 9 authorized syringe exchange program continues to meet all 10 standards set forth in subdivision (d) and that a public health need 11 exists, it may administratively approve amendments to a program's 12 operations including, but not limited to, modifications to the time, 13 location, and type of services provided, including the designation 14 as a fixed site or a mobile site. The amendment approval is not 15 subject to the noticing requirements of subdivision (e).

(j) The department shall have 30 business days to review andrespond to the applicant's request for amendment of theauthorization. If the department does not respond in writing within

19 30 business days, the request shall be deemed denied.

20 (k) The provisions of this section are severable. If any provision

21 of this section or its application is held invalid, that invalidity shall

not affect other provisions or applications that can be given effect

23 without the invalid provision or application.

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