SENATE SUBSTITUTE

FOR

SENATE COMMITTEE SUBSTITUTE

FOR

HOUSE COMMITTEE SUBSTITUTE

FOR

HOUSE BILL NO. 1682

AN ACT


BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF MISSOURI, AS FOLLOWS:

The month of May is hereby designated as "Mental Health Awareness Month". The citizens of this state are encouraged to participate in appropriate awareness and educational activities that emphasize the importance of good mental health and the effects of mental illness on Missourians.

The month of July shall be known as "Minority Mental Health Awareness Month". The citizens of this state are encouraged to observe the month with appropriate events and activities to raise awareness of the effects of mental illness on minorities.

The month of September shall be designated as "Deaf Awareness Month" and the last week of September shall be designated as "Deaf Awareness Week" in Missouri. The citizens of this state are encouraged to participate in appropriate activities and events to commemorate the first World Congress of the World Federation of the Deaf in 1951 and to increase awareness of deaf issues, people, and culture.

1. This section shall be known and may be cited as the "Public Access to Automated External Defibrillator Act".

2. [A person or entity who acquires an automated external defibrillator shall ensure that:

(1) Expected defibrillator users receive training by the American Red Cross or American Heart Association in cardiopulmonary resuscitation and the use of automated external defibrillators, or an equivalent nationally recognized course in defibrillator use and cardiopulmonary resuscitation;

(2) The defibrillator is maintained and tested according to
the manufacturer's operational guidelines;

(3) Any person who renders emergency care or treatment on a
person in cardiac arrest by using an automated external
defibrillator activates the emergency medical services system as
soon as possible; and

(4) Any person or entity that owns an automated external
defibrillator that is for use outside of a health care facility
shall have a physician review and approve the clinical protocol
for the use of the defibrillator, review and advise regarding the
training and skill maintenance of the intended users of the
defibrillator and assure proper review of all situations when the
defibrillator is used to render emergency care.

3. Any person or entity who acquires an automated external
defibrillator shall notify the emergency communications district
or the ambulance dispatch center of the primary provider of
emergency medical services where the automated external
defibrillator is to be located.

4. A person or entity that acquires an automated external
defibrillator shall:

(1) Comply with all regulations governing the placement of
an automated external defibrillator;

(2) Ensure that the automated external defibrillator is
maintained and tested according to the operation and maintenance
guidelines set forth by the manufacturer;

(3) Ensure that the automated external defibrillator is
tested at least every two years and after each use; and

(4) Ensure that an inspection is made of all automated
external defibrillators on the premises at least every ninety
days for potential issues related to the operation of the device, including a blinking light or other obvious defect that may suggest tampering or that another problem has arisen with the functionality of the automated external defibrillator.

3. Any person who gratuitously and in good faith renders emergency care by use of or provision of an automated external defibrillator shall not be held liable for any civil damages or subject to any criminal penalty as a result of such care or treatment, unless the person acts in a willful and wanton or reckless manner in providing the care, advice, or assistance. The person who or entity that provides training to the person using an automated external defibrillator, the person or entity responsible for the site where the automated external defibrillator is located, and the person or entity that owns the automated external defibrillator[; the person or entity that provided clinical protocol for automated external defibrillator sites or programs, and the licensed physician who reviews and approves the clinical protocol] shall likewise not be held liable for civil damages or subject to any criminal penalty resulting from the use of an automated external defibrillator. [Nothing in this section shall affect any claims brought pursuant to chapter 537 or 538.]

[5.] 4. All basic life support ambulances and stretcher vans operated in the state of Missouri shall be equipped with an automated external defibrillator and be staffed by at least one individual trained in the use of an automated external defibrillator.

[6.] 5. The provisions of this section shall apply in all
1 counties within the state and any city not within a county.

190.094. 1. Any ambulance licensed in this state, when used as an ambulance and staffed with volunteer staff, shall be staffed with a minimum of one emergency medical technician and one other crew member who may be a licensed emergency medical technician, registered nurse, physician, physician assistant, or someone who has an emergency medical responder certification.

2. When transporting a patient, at least one licensed emergency medical technician, registered nurse, physician assistant, or physician shall be in attendance with the patient in the patient compartment at all times.

3. For purposes of this section, "volunteer" shall mean an individual who performs hours of service without promise, expectation or receipt of compensation for services rendered. Compensation such as a nominal stipend per call to compensate for fuel, uniforms, and training shall not nullify the volunteer status.

190.105. 1. No person, either as owner, agent or otherwise, shall furnish, operate, conduct, maintain, advertise, or otherwise be engaged in or profess to be engaged in the business or service of the transportation of patients by ambulance in the air, upon the streets, alleys, or any public way or place of the state of Missouri unless such person holds a currently valid license from the department for an ambulance service issued pursuant to the provisions of sections 190.001 to 190.245.

2. No ground ambulance shall be operated for ambulance purposes, and no individual shall drive, attend or permit it to
be operated for such purposes in the state of Missouri unless the
ground ambulance is under the immediate supervision and direction
of a person who is holding a currently valid Missouri license as
an emergency medical technician. Nothing in this section shall
be construed to mean that a duly registered nurse or a duly
licensed physician, or a duly licensed physician assistant be
required to hold an emergency medical technician's license. When
a physician assistant is in attendance with a patient on an
ambulance, the physician assistant shall be exempt from any
mileage limitations in any collaborative practice arrangement
prescribed under law. Each ambulance service is responsible for
assuring that any person driving its ambulance is competent in
emergency vehicle operations and has a safe driving record. Each
ground ambulance shall be staffed with at least two licensed
individuals when transporting a patient, except as provided in
section 190.094. In emergency situations which require
additional medical personnel to assist the patient during
transportation, an emergency medical responder, firefighter, or
law enforcement personnel with a valid driver's license and prior
experience with driving emergency vehicles may drive the ground
ambulance provided the ground ambulance service stipulates to
this practice in operational policies.

3. No license shall be required for an ambulance service,
or for the attendant of an ambulance, which:

   (1) Is rendering assistance in the case of an emergency,
   major catastrophe or any other unforeseen event or series of
   events which jeopardizes the ability of the local ambulance
   service to promptly respond to emergencies; or
(2) Is operated from a location or headquarters outside of Missouri in order to transport patients who are picked up beyond the limits of Missouri to locations within or outside of Missouri, but no such outside ambulance shall be used to pick up patients within Missouri for transportation to locations within Missouri, except as provided in subdivision (1) of this subsection.

4. The issuance of a license pursuant to the provisions of sections 190.001 to 190.245 shall not be construed so as to authorize any person to provide ambulance services or to operate any ambulances without a franchise in any city not within a county or in a political subdivision in any county with a population of over nine hundred thousand inhabitants, or a franchise, contract or mutual-aid agreement in any other political subdivision which has enacted an ordinance making it unlawful to do so.

5. Sections 190.001 to 190.245 shall not preclude the adoption of any law, ordinance or regulation not in conflict with such sections by any city not within a county, or at least as strict as such sections by any county, municipality or political subdivision except that no such regulations or ordinances shall be adopted by a political subdivision in a county with a population of over nine hundred thousand inhabitants except by the county's governing body.

6. In a county with a population of over nine hundred thousand inhabitants, the governing body of the county shall set the standards for all ambulance services which shall comply with subsection 5 of this section. All such ambulance services must
be licensed by the department. The governing body of such county
shall not prohibit a licensed ambulance service from operating in
the county, as long as the ambulance service meets county
standards.

7. An ambulance service or vehicle when operated for the
purpose of transporting persons who are sick, injured, or
otherwise incapacitated shall not be treated as a common or
contract carrier under the jurisdiction of the Missouri division
of motor carrier and railroad safety.

8. Sections 190.001 to 190.245 shall not apply to, nor be
construed to include, any motor vehicle used by an employer for
the transportation of such employer's employees whose illness or
injury occurs on private property, and not on a public highway or
property, nor to any person operating such a motor vehicle.

9. A political subdivision that is authorized to operate a
licensed ambulance service may establish, operate, maintain and
manage its ambulance service, and select and contract with a
licensed ambulance service. Any political subdivision may
contract with a licensed ambulance service.

10. Except as provided in subsections 5 and 6, nothing in
section 67.300, or subsection 2 of section 190.109, shall be
construed to authorize any municipality or county which is
located within an ambulance district or a fire protection
district that is authorized to provide ambulance service to
promulgate laws, ordinances or regulations related to the
provision of ambulance services. This provision shall not apply
to any municipality or county which operates an ambulance service
established prior to August 28, 1998.
11. Nothing in section 67.300 or subsection 2 of section 190.109 shall be construed to authorize any municipality or county which is located within an ambulance district or a fire protection district that is authorized to provide ambulance service to operate an ambulance service without a franchise in an ambulance district or a fire protection district that is authorized to provide ambulance service which has enacted an ordinance making it unlawful to do so. This provision shall not apply to any municipality or county which operates an ambulance service established prior to August 28, 1998.

12. No provider of ambulance service within the state of Missouri which is licensed by the department to provide such service shall discriminate regarding treatment or transportation of emergency patients on the basis of race, sex, age, color, religion, sexual preference, national origin, ancestry, handicap, medical condition or ability to pay.

13. No provision of this section, other than subsections 5, 6, 10 and 11 of this section, is intended to limit or supersede the powers given to ambulance districts pursuant to this chapter or to fire protection districts pursuant to chapter 321, or to counties, cities, towns and villages pursuant to chapter 67.

14. Upon the sale or transfer of any ground ambulance service ownership, the owner of such service shall notify the department of the change in ownership within thirty days of such sale or transfer. After receipt of such notice, the department shall conduct an inspection of the ambulance service to verify compliance with the licensure standards of sections 190.001 to 190.245.
190.143. 1. Notwithstanding any other provisions of law, the department may grant a ninety-day temporary emergency medical technician license to all levels of emergency medical technicians who meet the following:

(1) Can demonstrate that they have, or will have, employment requiring an emergency medical technician license;

(2) Are not currently licensed as an emergency medical technician in Missouri or have been licensed as an emergency medical technician in Missouri and fingerprints need to be submitted to the Federal Bureau of Investigation to verify the existence or absence of a criminal history, or they are currently licensed and the license will expire before a verification can be completed of the existence or absence of a criminal history;

(3) Have submitted a complete application upon such forms as prescribed by the department in rules adopted pursuant to sections 190.001 to 190.245;

(4) Have not been disciplined pursuant to sections 190.001 to 190.245 and rules promulgated pursuant to sections 190.001 to 190.245;

(5) Meet all the requirements of rules promulgated pursuant to sections 190.001 to 190.245.

2. A temporary emergency medical technician license shall only authorize the license to practice while under the immediate supervision of a licensed emergency medical technician, registered nurse, physician assistant, or physician who is currently licensed, without restrictions, to practice in Missouri.

3. A temporary emergency medical technician license shall
automatically expire either ninety days from the date of issuance or upon the issuance of a five-year emergency medical technician license.

190.196. 1. No employer shall knowingly employ or permit any employee to perform any services for which a license, certificate or other authorization is required by sections 190.001 to 190.245, or by rules adopted pursuant to sections 190.001 to 190.245, unless and until the person so employed possesses all licenses, certificates or authorizations that are required.

2. Any person or entity that employs or supervises a person's activities as an emergency medical responder, emergency medical dispatcher, emergency medical technician, registered nurse, physician assistant, or physician shall cooperate with the department's efforts to monitor and enforce compliance by those individuals subject to the requirements of sections 190.001 to 190.245.

3. Any person or entity who employs individuals licensed by the department pursuant to sections 190.001 to 190.245 shall report to the department within seventy-two hours of their having knowledge of any charges filed against a licensee in their employ for possible criminal action involving the following felony offenses:

(1) Child abuse or sexual abuse of a child;

(2) Crimes of violence; or

(3) Rape or sexual abuse.

4. Any licensee who has charges filed against him or her for the felony offenses in subsection 3 of this section shall
report such an occurrence to the department within seventy-two hours of the charges being filed.

5. The department will monitor these reports for possible licensure action authorized pursuant to section 190.165.

190.606. The following persons and entities shall not be subject to civil, criminal, or administrative liability and are not guilty of unprofessional conduct for the following acts or omissions that follow discovery of an outside the hospital do-not-resuscitate identification upon a patient, or upon being presented with an outside the hospital do-not-resuscitate order from Missouri, another state, the District of Columbia, or a territory of the United States; provided that the acts or omissions are done in good faith and in accordance with the provisions of sections 190.600 to 190.621 and the provisions of an outside the hospital do-not-resuscitate order executed under sections 190.600 to 190.621:

(1) Physicians, persons under the direction or authorization of a physician, emergency medical services personnel, or health care facilities that cause or participate in the withholding or withdrawal of cardiopulmonary resuscitation from such patient; and

(2) Physicians, persons under the direction or authorization of a physician, emergency medical services personnel, or health care facilities that provide cardiopulmonary resuscitation to such patient under an oral or written request communicated to them by the patient or the patient's representative.

190.612. 1. Emergency medical services personnel are
authorized to comply with the outside the hospital
do-not-resuscitate protocol when presented with an outside the
cardiac or respiratory arrest, the desire
to be resuscitated.

2. Emergency medical services personnel are authorized to
comply with the outside the hospital do-not-resuscitate protocol
when presented with an outside the hospital do-not-resuscitate
order from another state, the District of Columbia, or a
territory of the United States if such order is on a standardized
written form:

   (1) Signed by the patient or the patient's representative
   and a physician who is licensed to practice in the other state,
   the District of Columbia, or the territory of the United States;
   and

   (2) Such form has been previously reviewed and approved by
   the department of health and senior services to authorize
   emergency medical services personnel to withhold or withdraw
   cardiopulmonary resuscitation from the patient in the event of a
cardiac or respiratory arrest.

Emergency medical services personnel shall not comply with an
outside the hospital do-not-resuscitate order from another state,
the District of Columbia, or a territory of the United States or
the outside the hospital do-not-resuscitate protocol when the
patient or patient's representative expresses to such personnel
in any manner, before or after the onset of a cardiac or
respiratory arrest, the desire to be resuscitated.

3. If a physician or a health care facility other than a
hospital admits or receives a patient with an outside the
hospital do-not-resuscitate identification or an outside the
hospital do-not-resuscitate order, and the patient or patient's
representative has not expressed or does not express to the
physician or health care facility the desire to be resuscitated,
and the physician or health care facility is unwilling or unable
to comply with the outside the hospital do-not-resuscitate order,
the physician or health care facility shall take all reasonable
steps to transfer the patient to another physician or health care
facility where the outside the hospital do-not-resuscitate order
will be complied with.

190.1005. Notwithstanding any other provision of law to the
 contrary, any training or course in cardiopulmonary resuscitation
 shall also include instruction on the proper use of automated
 external defibrillators. Such training or course shall follow
 the standards created by the American Red Cross or the American
 Heart Association, or equivalent evidence-based standards from a
 nationally recognized organization.

191.775. No person shall smoke or otherwise use tobacco
products, or vapor products, as such term is
defined in section 407.925, in any indoor area of a public
elementary or secondary school building or educational facility,
excluding institutions of higher education, or on buses used
solely to transport students to or from school or to transport
students to or from any place for educational purposes. Any
school board of any school district may set policy on the
permissible uses of tobacco products or vapor products in any
other nonclassroom or nonstudent occupant facility, and on the
school grounds or outdoor facility areas as the school board
deems proper. [Any person who violates the provisions of this
section shall be guilty of an infraction.]
establish an informed diagnosis as though the medical interview and physical examination has been performed in person; and

(2) Prior to providing treatment, including issuing prescriptions or physician certifications under article XIV of the Missouri Constitution, a physician who uses telemedicine shall interview the patient, collect or review relevant medical history, and perform an examination sufficient for the diagnosis and treatment of the patient. A questionnaire completed by the patient, whether via the internet or telephone, does not constitute an acceptable medical interview and examination for the provision of treatment by telehealth.

192.2305. 1. There is hereby established within the department of health and senior services the "Office of State Ombudsman for Long-Term Care Facility Residents", for the purpose of helping to assure the adequacy of care received by residents of long-term care facilities and Missouri veterans' homes, as defined in section 42.002, and to improve the quality of life experienced by them, in accordance with the federal Older Americans Act, 42 U.S.C. Section 3001, et seq.

2. The office shall be administered by the state ombudsman, who shall devote his or her entire time to the duties of his or her position.

3. The office shall establish and implement procedures for receiving, processing, responding to, and resolving complaints made by or on behalf of residents of long-term care facilities and Missouri veterans' homes relating to action, inaction, or decisions of providers, or their representatives, of long-term care services, of public agencies or of social service agencies,
which may adversely affect the health, safety, welfare or rights of such residents.

4. The department shall establish and implement procedures for resolution of complaints. The ombudsman or representatives of the office shall have the authority to:

   (1) Enter any long-term care facility or Missouri veterans' homes and have access to residents of the facility at a reasonable time and in a reasonable manner. The ombudsman shall have access to review resident records, if given permission by the resident or the resident's legal guardian. Residents of the facility shall have the right to request, deny, or terminate visits with an ombudsman;

   (2) Make the necessary inquiries and review such information and records as the ombudsman or representative of the office deems necessary to accomplish the objective of verifying these complaints.

5. The office shall acknowledge complaints, report its findings, make recommendations, gather and disseminate information and other material, and publicize its existence.

6. The ombudsman may recommend to the relevant governmental agency changes in the rules and regulations adopted or proposed by such governmental agency which do or may adversely affect the health, safety, welfare, or civil or human rights of any resident in a facility. The office shall analyze and monitor the development and implementation of federal, state and local laws, regulations and policies with respect to long-term care facilities and services and Missouri veterans' homes in the state and shall recommend to the department changes in such laws,
regulations and policies deemed by the office to be appropriate.

7. The office shall promote community contact and involvement with residents of facilities through the use of volunteers and volunteer programs directed by the regional ombudsman coordinators.

8. The office shall develop and establish by regulation of the department statewide policies and standards for implementing the activities of the ombudsman program, including the qualifications and the training of regional ombudsman coordinators and ombudsman volunteers.

9. The office shall develop and propose programs for use, training and coordination of volunteers in conjunction with the regional ombudsman coordinators and may:
   (1) Establish and conduct recruitment programs for volunteers;
   (2) Establish and conduct training seminars, meetings and other programs for volunteers; and
   (3) Supply personnel, written materials and such other reasonable assistance, including publicizing their activities, as may be deemed necessary.

10. The regional ombudsman coordinators and ombudsman volunteers shall have the authority to report instances of abuse and neglect to the ombudsman hotline operated by the department.

11. If the regional ombudsman coordinator or volunteer finds that a nursing home administrator is not willing to work with the ombudsman program to resolve complaints, the state ombudsman shall be notified. The department shall establish procedures by rule in accordance with chapter 536 for
implementation of this subsection.

12. The office shall prepare and distribute to each facility written notices which set forth the address and telephone number of the office, a brief explanation of the function of the office, the procedure to follow in filing a complaint and other pertinent information.

13. The administrator of each facility shall ensure that such written notice is given to every resident or the resident's guardian upon admission to the facility and to every person already in residence, or to his or her guardian. The administrator shall also post such written notice in a conspicuous, public place in the facility in the number and manner set forth in the regulations adopted by the department.

14. The office shall inform residents, their guardians or their families of their rights and entitlements under state and federal laws and rules and regulations by means of the distribution of educational materials and group meetings.

195.070. 1. A physician, podiatrist, dentist, a registered optometrist certified to administer pharmaceutical agents as provided in section 336.220, or an assistant physician in accordance with section 334.037 or a physician assistant in accordance with section 334.747 in good faith and in the course of his or her professional practice only, may prescribe, administer, and dispense controlled substances or he or she may cause the same to be administered or dispensed by an individual as authorized by statute.

2. An advanced practice registered nurse, as defined in section 335.016, but not a certified registered nurse anesthetist
as defined in subdivision (8) of section 335.016, who holds a certificate of controlled substance prescriptive authority from the board of nursing under section 335.019 and who is delegated the authority to prescribe controlled substances under a collaborative practice arrangement under section 334.104 may prescribe any controlled substances listed in Schedules III, IV, and V of section 195.017, and may have restricted authority in Schedule II. Prescriptions for Schedule II medications prescribed by an advanced practice registered nurse who has a certificate of controlled substance prescriptive authority are restricted to only those medications containing hydrocodone. However, no such certified advanced practice registered nurse shall prescribe controlled substance for his or her own self or family. Schedule III narcotic controlled substance and Schedule II - hydrocodone prescriptions shall be limited to a one hundred twenty-hour supply without refill.

3. A veterinarian, in good faith and in the course of the veterinarian's professional practice only, and not for use by a human being, may prescribe, administer, and dispense controlled substances and the veterinarian may cause them to be administered by an assistant or orderly under his or her direction and supervision.

4. A practitioner shall not accept any portion of a controlled substance unused by a patient, for any reason, if such practitioner did not originally dispense the drug, except:

(1) When the controlled substance is delivered to the practitioner to administer to the patient for whom the medication is prescribed as authorized by federal law. Practitioners shall
maintain records and secure the medication as required by this chapter and regulations promulgated pursuant to this chapter; or

(2) As provided in section 195.265.

5. An individual practitioner shall not prescribe or dispense a controlled substance for such practitioner's personal use except in a medical emergency.

195.417. 1. The limits specified in this section shall not apply to any quantity of such product, mixture, or preparation which must be dispensed, sold, or distributed in a pharmacy pursuant to a valid prescription.

2. Within any thirty-day period, no person shall sell, dispense, or otherwise provide to the same individual, and no person shall purchase, receive, or otherwise acquire more than the following amount: any number of packages of any drug product containing any detectable amount of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts or optical isomers, or salts of optical isomers, either as:

(1) The sole active ingredient; or

(2) One of the active ingredients of a combination drug; or

(3) A combination of any of the products specified in subdivisions (1) and (2) of this subsection;

in any total amount greater than [nine] seven and two-tenths grams, without regard to the number of transactions.

3. Within any twenty-four-hour period, no pharmacist, intern pharmacist, or registered pharmacy technician shall sell, dispense, or otherwise provide to the same individual, and no person shall purchase, receive, or otherwise acquire more than
the following amount: any number of packages of any drug product
containing any detectable amount of ephedrine,
phenylpropanolamine, or pseudoephedrine, or any of their salts or
optical isomers, or salts of optical isomers, either as:

(1) The sole active ingredient; or
(2) One of the active ingredients of a combination drug; or
(3) A combination of any of the products specified in
subdivisions (1) and (2) of this subsection;

in any total amount greater than three and six-tenths grams
without regard to the number of transactions.

4. Within any twelve-month period, no person shall sell,
dispense, or otherwise provide to the same individual, and no
person shall purchase, receive, or otherwise acquire more than
the following amount: any number of packages of any drug product
containing any detectable amount of ephedrine,
phenylpropanolamine, or pseudoephedrine, or any of their salts or
optical isomers, or salts of optical isomers, either as:

(1) The sole active ingredient; or
(2) One of the active ingredients of a combination drug; or
(3) A combination of any of the products specified in
subdivisions (1) and (2) of this subsection;

in any total amount greater than forty-three and two-tenths
grams, without regard to the number of transactions.

5. All packages of any compound, mixture, or preparation
containing any detectable quantity of ephedrine,
phenylpropanolamine, or pseudoephedrine, or any of their salts or
optical isomers, or salts of optical isomers, except those that are excluded from Schedule V in subsection 17 or 18 of section 195.017, shall be offered for sale only from behind a pharmacy counter where the public is not permitted, and only by a registered pharmacist or registered pharmacy technician under section 195.017.

[5.] 6. Each pharmacy shall submit information regarding sales of any compound, mixture, or preparation as specified in this section in accordance with transmission methods and frequency established by the department by regulation.

7. No prescription shall be required for the dispensation, sale, or distribution of any drug product containing any detectable amount of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts or optical isomers, or salts of optical isomers, in an amount within the limits described in subsections 2, 3, and 4 of this section. The superintendent of the Missouri state highway patrol shall report to the revisor of statutes and the general assembly by February first when the statewide number of methamphetamine laboratory seizure incidents exceeds three hundred incidents in the previous calendar year. The provisions of this subsection shall expire on April first of the calendar year in which the revisor of statutes receives such notification.

[6.] 8. This section shall supersede and preempt any local ordinances or regulations, including any ordinances or regulations enacted by any political subdivision of the state. This section shall not apply to the sale of any animal feed products containing ephedrine or any naturally occurring or
herbal ephedra or extract of ephedra.

9. Any local ordinances or regulations enacted by any political subdivision of the state prior to August 28, 2020, requiring a prescription for the dispensation, sale, or distribution of any drug product containing any detectable amount of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts or optical isomers, or salts of optical isomers, in an amount within the limits described in subsections 2, 3, and 4 of this section shall be void and of no effect and no such political subdivision shall maintain or enforce such ordinance or regulation.

7. All logs, records, documents, and electronic information maintained for the dispensing of these products shall be open for inspection and copying by municipal, county, and state or federal law enforcement officers whose duty it is to enforce the controlled substances laws of this state or the United States.

8. All persons who dispense or offer for sale pseudoephedrine and ephedrine products, except those that are excluded from Schedule V in subsection 17 or 18 of section 195.017, shall ensure that all such products are located only behind a pharmacy counter where the public is not permitted.

9. The penalty for a knowing or reckless violation of this section is found in section 579.060.

195.805. 1. No edible marijuana-infused product, packaging, or logo sold in Missouri pursuant to article XIV of the Missouri Constitution shall be designed in the shape of a human, animal, or fruit, including realistic, artistic,
1 caricature, or cartoon renderings. However, geometric shapes,  
2 including, but not limited to, circles, squares, rectangles, and  
3 triangles, shall be permitted.

2. Each package, or packages with a package, containing an  
edible marijuana-infused product with ten or more milligrams of  
tetrahydrocannabinols (THC) shall be stamped with a universal  
symbol for such products, which shall consist of the following:

   (1) A diamond containing the letters "THC";
   (2) The letter "M" located under the "THC" within the  
diamond, to signify that the product is for medical purposes; and
   (3) The number of milligrams of THC in the package.

   The universal symbol shall be placed on the front of the package  
in red and white print and shall measure one-half inch by one-  
half inch from point to point.

3. Any licensed or certified entity regulated by the  
department of health and senior services pursuant to article XIV  
of the Missouri Constitution found to have violated the  
provisions of this section shall be subject to department  
sanctions, including an administrative penalty, in accordance  
with the regulations promulgated by the department pursuant to  
article XIV of the Missouri Constitution.

4. The department shall promulgate rules and regulations  
prohibiting edible marijuana-infused products designed to appeal  
to persons under eighteen years of age, as well as promulgate  
rules and regulations to establish a process by which a licensed  
or certified entity may seek approval of an edible product  
design, package, or label prior to such product's manufacture or
sale in order to determine compliance with the provisions of this section and any rules promulgated pursuant to this section. Any rule or portion of a rule, as that term is defined in section 536.010 that is created under the authority delegated in this section shall become effective only if it complies with and is subject to all of the provisions of chapter 536 and, if applicable, section 536.028. This section and chapter 536 are nonseverable and if any of the powers vested with the general assembly pursuant to chapter 536 to review, to delay the effective date, or to disapprove and annul a rule are subsequently held unconstitutional, then the grant of rulemaking authority and any rule proposed or adopted after August 28, 2020, shall be invalid and void.

195.815. 1. The department of health and senior services shall require all officers, managers, contractors, employees, and other support staff of licensed or certified medical marijuana facilities, and all owners of such medical marijuana facilities who will have access to the facilities or to the facilities' medical marijuana, to submit fingerprints to the Missouri state highway patrol for the purpose of conducting a state and federal fingerprint-based criminal background check.

2. The department may require that such fingerprint submissions be made as part of a medical marijuana facility application for licensure or certification, a medical marijuana facility application for renewal of licensure or certification, and an individual's application for an identification card authorizing that individual to be an owner, officer, manager, contractor, employee, or other support staff of a medical
marijuana facility.

3. Fingerprint cards and any required fees shall be sent to the Missouri state highway patrol's central repository. The fingerprints shall be used for searching the state criminal records repository and shall also be forwarded to the Federal Bureau of Investigation for a federal criminal records search under section 43.540. The Missouri state highway patrol shall notify the department of any criminal history record information or lack of criminal history record information discovered on the individual. Notwithstanding the provisions of section 610.120 to the contrary, all records related to any criminal history information discovered shall be accessible and available to the department.

4. As used in this section, the following words shall mean:

(1) "Employee", any person performing work or service of any kind or character for hire in a medical marijuana facility;

(2) "Medical marijuana facility", an entity licensed or certified by the department of health and senior services, or its successor agency, to acquire, cultivate, process, manufacture, test, store, sell, transport, or deliver medical marijuana;

(3) "Other support staff", any person performing work or service of any kind or character, other than employees, on behalf of a medical marijuana facility if such a person would have access to the medical marijuana facility or its medical marijuana or related equipment or supplies.

196.990. 1. As used in this section, the following terms shall mean:

(1) "Administer", the direct application of an epinephrine
(2) "Authorized entity", any entity or organization at or in connection with which allergens capable of causing anaphylaxis may be present including, but not limited to, qualified first responders, as such term is defined in section 321.621, restaurants, recreation camps, youth sports leagues, amusement parks, and sports arenas. "Authorized entity" shall not include any public school or public charter school;

(3) "Epinephrine auto-injector", a single-use device used for the automatic injection of a premeasured dose of epinephrine into the human body;

(4) "Physician", a physician licensed in this state under chapter 334;

(5) "Provide", the supply of one or more epinephrine auto-injectors to an individual;

(6) "Self-administration", a person's discretionary use of an epinephrine auto-injector.

2. A physician may prescribe epinephrine auto-injectors in the name of an authorized entity for use in accordance with this section, and pharmacists, physicians, and other persons authorized to dispense prescription medications may dispense epinephrine auto-injectors under a prescription issued in the name of an authorized entity.

3. An authorized entity may acquire and stock a supply of epinephrine auto-injectors under a prescription issued in accordance with this section. Such epinephrine auto-injectors shall be stored in a location readily accessible in an emergency and in accordance with the epinephrine auto-injector's
instructions for use and any additional requirements established by the department of health and senior services by rule. An authorized entity shall designate employees or agents who have completed the training required under this section to be responsible for the storage, maintenance, and general oversight of epinephrine auto-injectors acquired by the authorized entity.

4. An authorized entity that acquires a supply of epinephrine auto-injectors under a prescription issued in accordance with this section shall ensure that:

   (1) Expected epinephrine auto-injector users receive training in recognizing symptoms of severe allergic reactions including anaphylaxis and the use of epinephrine auto-injectors from a nationally recognized organization experienced in training laypersons in emergency health treatment or another entity or person approved by the department of health and senior services;

   (2) All epinephrine auto-injectors are maintained and stored according to the epinephrine auto-injector's instructions for use;

   (3) Any person who provides or administers an epinephrine auto-injector to an individual who the person believes in good faith is experiencing anaphylaxis activates the emergency medical services system as soon as possible; and

   (4) A proper review of all situations in which an epinephrine auto-injector is used to render emergency care is conducted.

5. Any authorized entity that acquires a supply of epinephrine auto-injectors under a prescription issued in accordance with this section shall notify the emergency
communications district or the ambulance dispatch center of the primary provider of emergency medical services where the epinephrine auto-injectors are to be located within the entity's facility.

6. No person shall provide or administer an epinephrine auto-injector to any individual who is under eighteen years of age without the verbal consent of a parent or guardian who is present at the time when provision or administration of the epinephrine auto-injector is needed. Provided, however, that a person may provide or administer an epinephrine auto-injector to such an individual without the consent of a parent or guardian if the parent or guardian is not physically present and the person reasonably believes the individual shall be in imminent danger without the provision or administration of the epinephrine auto-injector.

7. The following persons and entities shall not be liable for any injuries or related damages that result from the administration or self-administration of an epinephrine auto-injector in accordance with this section that may constitute ordinary negligence:

   (1) An authorized entity that possesses and makes available epinephrine auto-injectors and its employees, agents, and other trained persons;

   (2) Any person who uses an epinephrine auto-injector made available under this section;

   (3) A physician that prescribes epinephrine auto-injectors to an authorized entity; or

   (4) Any person or entity that conducts the training
described in this section.

Such immunity does not apply to acts or omissions constituting a reckless disregard for the safety of others or willful or wanton conduct. The administration of an epinephrine auto-injector in accordance with this section shall not be considered the practice of medicine. The immunity from liability provided under this subsection is in addition to and not in lieu of that provided under section 537.037. An authorized entity located in this state shall not be liable for any injuries or related damages that result from the provision or administration of an epinephrine auto-injector by its employees or agents outside of this state if the entity or its employee or agent is not liable for such injuries or related damages under the laws of the state in which such provision or administration occurred. No trained person who is in compliance with this section and who in good faith and exercising reasonable care fails to administer an epinephrine auto-injector shall be liable for such failure.

8. All basic life support ambulances and stretcher vans operated in the state shall be equipped with epinephrine auto-injectors and be staffed by at least one individual trained in the use of epinephrine auto-injectors.

9. The provisions of this section shall apply in all counties within the state and any city not within a county.

10. Nothing in this section shall be construed as superseding the provisions of section 167.630.

196.1050. 1. The proceeds of any monetary settlement or portion of a global settlement between the attorney general of
the state and any drug manufacturers, distributors, or combination thereof to resolve an opioid-related cause of action against such drug manufacturers, distributors, or combination thereof in a state or federal court shall only be utilized to pay for opioid addiction treatment and prevention services and health care and law enforcement costs related to opioid addiction treatment and prevention. Under no circumstances shall such settlement moneys be utilized to fund other services, programs, or expenses not reasonably related to opioid addiction treatment and prevention.

2. (1) There is hereby established in the state treasury the "Opioid Addiction Treatment and Recovery Fund", which shall consist of the proceeds of any settlement described in subsection 1 of this section, as well as any funds appropriated by the general assembly, or gifts, grants, donations, or bequests. The state treasurer shall be custodian of the fund. In accordance with sections 30.170 and 30.180, the state treasurer may approve disbursements. The fund shall be a dedicated fund and money in the fund shall be used by the department of mental health, the department of health and senior services, the department of social services, and the department of public safety for the purposes set forth in subsection 1 of this section.

(2) Notwithstanding the provisions of section 33.080 to the contrary, any moneys remaining in the fund at the end of the biennium shall not revert to the credit of the general revenue fund.

(3) The state treasurer shall invest moneys in the fund in the same manner as other funds are invested. Any interest and
moneys earned on such investments shall be credited to the fund.

205.202. 1. The governing body of any hospital district established under sections 205.160 to 205.379 in any county of the third classification without a township form of government and with more than thirteen thousand five hundred but fewer than thirteen thousand six hundred inhabitants may, by resolution, abolish the property tax levied in such district under this chapter and impose a sales tax on all retail sales made within the district which are subject to sales tax under chapter 144. The tax authorized in this section shall be not more than one percent, and shall be imposed solely for the purpose of funding the hospital district. The tax authorized in this section shall be in addition to all other sales taxes imposed by law, and shall be stated separately from all other charges and taxes.

2. No such resolution adopted under this section shall become effective unless the governing body of the hospital district submits to the voters residing within the district at a state general, primary, or special election a proposal to authorize the governing body of the district to impose a tax under this section. If a majority of the votes cast on the question by the qualified voters voting thereon are in favor of the question, then the tax shall become effective on the first day of the second calendar quarter after the director of revenue receives notification of adoption of the local sales tax. If a majority of the votes cast on the question by the qualified voters voting thereon are opposed to the question, then the tax shall not become effective unless and until the question is resubmitted under this section to the qualified voters and such
question is approved by a majority of the qualified voters voting
on the question.

3. All revenue collected under this section by the director
of the department of revenue on behalf of the hospital district,
except for one percent for the cost of collection which shall be
deposited in the state's general revenue fund, shall be deposited
in a special trust fund, which is hereby created and shall be
known as the "Hospital District Sales Tax Fund", and shall be
used solely for the designated purposes. Moneys in the fund
shall not be deemed to be state funds, and shall not be
commingled with any funds of the state. The director may make
refunds from the amounts in the fund and credited to the district
for erroneous payments and overpayments made, and may redeem
dishonored checks and drafts deposited to the credit of such
district. Any funds in the special fund which are not needed for
current expenditures shall be invested in the same manner as
other funds are invested. Any interest and moneys earned on such
investments shall be credited to the fund.

4. The governing body of any hospital district that has
adopted the sales tax authorized in this section may submit the
question of repeal of the tax to the voters on any date available
for elections for the district. If a majority of the votes cast
on the question by the qualified voters voting thereon are in
favor of the repeal, that repeal shall become effective on
December thirty-first of the calendar year in which such repeal
was approved. If a majority of the votes cast on the question by
the qualified voters voting thereon are opposed to the repeal,
then the sales tax authorized in this section shall remain
effective until the question is resubmitted under this section to
the qualified voters and the repeal is approved by a majority of
the qualified voters voting on the question.

5. Whenever the governing body of any hospital district
that has adopted the sales tax authorized in this section
receives a petition, signed by a number of registered voters of
the district equal to at least ten percent of the number of
registered voters of the district voting in the last
gubernatorial election, calling for an election to repeal the
sales tax imposed under this section, the governing body shall
submit to the voters of the district a proposal to repeal the
tax. If a majority of the votes cast on the question by the
qualified voters voting thereon are in favor of the repeal, the
repeal shall become effective on December thirty-first of the
calendar year in which such repeal was approved. If a majority
of the votes cast on the question by the qualified voters voting
thereon are opposed to the repeal, then the sales tax authorized
in this section shall remain effective until the question is
resubmitted under this section to the qualified voters and the
repeal is approved by a majority of the qualified voters voting
on the question.

6. If the tax is repealed or terminated by any means other
than by a dissolution of a hospital district as described in
subsection 7 of this section, all funds remaining in the special
trust fund shall continue to be used solely for the designated
purposes, and the hospital district shall notify the director of
the department of revenue of the action at least ninety days
before the effective date of the repeal and the director may
order retention in the trust fund, for a period of one year, of
two percent of the amount collected after receipt of such notice
to cover possible refunds or overpayment of the tax and to redeem
dishonored checks and drafts deposited to the credit of such
accounts. After one year has elapsed after the effective date of
abolition of the tax in such district, the director shall remit
the balance in the account to the district and close the account
of that district. The director shall notify each district of
each instance of any amount refunded or any check redeemed from
receipts due the district.

7. Upon the dissolution of a hospital district levying a
sales tax pursuant to this section, the sales tax shall be
automatically repealed and all funds remaining in the special
trust fund shall be distributed as follows:
   (1) Twenty-five percent shall be distributed to the county
       public health center established pursuant to sections 205.010 to
       205.150; and
   (2) Seventy-five percent shall be distributed to a
       federally qualified health center, as defined in 42 U.S.C.
       Section 1396d(l)(1) and (2), located in the county.

208.909. 1. Consumers receiving personal care assistance
services shall be responsible for:
   (1) Supervising their personal care attendant;
   (2) Verifying wages to be paid to the personal care
       attendant;
   (3) Preparing and submitting time sheets, signed by both
       the consumer and personal care attendant, to the vendor on a
       biweekly basis;
(4) Promptly notifying the department within ten days of any changes in circumstances affecting the personal care assistance services plan or in the consumer's place of residence;
(5) Reporting any problems resulting from the quality of services rendered by the personal care attendant to the vendor. If the consumer is unable to resolve any problems resulting from the quality of service rendered by the personal care attendant with the vendor, the consumer shall report the situation to the department; [and]
(6) Providing the vendor with all necessary information to complete required paperwork for establishing the employer identification number;
(7) Allowing the vendor to comply with its quality assurance and supervision process, which shall include, but not be limited to, annual face-to-face home visits and monthly case management activities; and
(8) Report to the department significant changes in their health and ability to self-direct care.

2. Participating vendors shall be responsible for:
(1) Collecting time sheets or reviewing reports of delivered services and certifying the accuracy thereof;
(2) The Medicaid reimbursement process, including the filing of claims and reporting data to the department as required by rule;
(3) Transmitting the individual payment directly to the personal care attendant on behalf of the consumer;
(4) Monitoring the performance of the personal care assistance services plan. Such monitoring shall occur during the
annual face-to-face home visit under section 208.918. The vendor shall document whether services are being provided to the consumer as set forth in the plan of care. If the attendant was not providing services as set forth in the plan of care, the vendor shall notify the department and the department may suspend services to the consumer; and

(5) Report to the department significant changes in the consumer's health or ability to self-direct care.

3. No state or federal financial assistance shall be authorized or expended to pay for services provided to a consumer under sections 208.900 to 208.927, if the primary benefit of the services is to the household unit, or is a household task that the members of the consumer's household may reasonably be expected to share or do for one another when they live in the same household, unless such service is above and beyond typical activities household members may reasonably provide for another household member without a disability.

4. No state or federal financial assistance shall be authorized or expended to pay for personal care assistance services provided by a personal care attendant who has not undergone the background screening process under section 192.2495. If the personal care attendant has a disqualifying finding under section 192.2495, no state or federal assistance shall be made, unless a good cause waiver is first obtained from the department in accordance with section 192.2495.

5. (1) All vendors shall, by July 1, 2015, have, maintain, and use a telephone tracking system for the purpose of reporting and verifying the delivery of consumer-directed services as
authorized by the department of health and senior services or its
designee. [Use of such a system prior to July 1, 2015, shall be
voluntary.] The telephone tracking system shall be used to
process payroll for employees and for submitting claims for
reimbursement to the MO HealthNet division. At a minimum, the
telephone tracking system shall:

(a) Record the exact date services are delivered;
(b) Record the exact time the services begin and exact time
the services end;
(c) Verify the telephone number from which the services are
registered;
(d) Verify that the number from which the call is placed is
a telephone number unique to the client;
(e) Require a personal identification number unique to each
personal care attendant;
(f) Be capable of producing reports of services delivered,
tasks performed, client identity, beginning and ending times of
service and date of service in summary fashion that constitute
adequate documentation of service; and
(g) Be capable of producing reimbursement requests for
consumer approval that assures accuracy and compliance with
program expectations for both the consumer and vendor.

(2) [The department of health and senior services, in
collaboration with other appropriate agencies, including centers
for independent living, shall establish telephone tracking system
pilot projects, implemented in two regions of the state, with one
in an urban area and one in a rural area. Each pilot project
shall meet the requirements of this section and section 208.918.

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The department of health and senior services shall, by December 31, 2013, submit a report to the governor and general assembly detailing the outcomes of these pilot projects. The report shall take into consideration the impact of a telephone tracking system on the quality of the services delivered to the consumer and the principles of self-directed care.

(3) As new technology becomes available, the department may allow use of a more advanced tracking system, provided that such system is at least as capable of meeting the requirements of this subsection.

[(4)] (3) The department of health and senior services shall promulgate by rule the minimum necessary criteria of the telephone tracking system. Any rule or portion of a rule, as that term is defined in section 536.010, that is created under the authority delegated in this section shall become effective only if it complies with and is subject to all of the provisions of chapter 536 and, if applicable, section 536.028. This section and chapter 536 are nonseverable and if any of the powers vested with the general assembly pursuant to chapter 536 to review, to delay the effective date, or to disapprove and annul a rule are subsequently held unconstitutional, then the grant of rulemaking authority and any rule proposed or adopted after August 28, 2010, shall be invalid and void.

[6. In the event that a consensus between centers for independent living and representatives from the executive branch cannot be reached, the telephony report issued to the general assembly and governor shall include a minority report which shall detail those elements of substantial dissent from the main
report.

7. No interested party, including a center for independent living, shall be required to contract with any particular vendor or provider of telephony services nor bear the full cost of the pilot program.

208.918. 1. In order to qualify for an agreement with the department, the vendor shall have a philosophy that promotes the consumer's ability to live independently in the most integrated setting or the maximum community inclusion of persons with physical disabilities, and shall demonstrate the ability to provide, directly or through contract, the following services:

(1) Orientation of consumers concerning the responsibilities of being an employer[,] and supervision of personal care attendants including the preparation and verification of time sheets. Such orientation shall include notifying customers that falsification of attendant visit verification records shall be considered fraud and shall be reported to the department. Such orientation shall take place in the presence of the personal care attendant, to the fullest extent possible;

(2) Training for consumers about the recruitment and training of personal care attendants;

(3) Maintenance of a list of persons eligible to be a personal care attendant;

(4) Processing of inquiries and problems received from consumers and personal care attendants;

(5) Ensuring the personal care attendants are registered with the family care safety registry as provided in sections
210.900 to [210.937] 210.936; and

(6) The capacity to provide fiscal conduit services through a telephone tracking system by the date required under section 208.909.

2. In order to maintain its agreement with the department, a vendor shall comply with the provisions of subsection 1 of this section and shall:

(1) Demonstrate sound fiscal management as evidenced on accurate quarterly financial reports and an annual financial statement audit [submitted to the department] performed by a certified public accountant if the vendor's annual gross revenue is two hundred thousand dollars or more or, if the vendor's annual gross revenue is less than two hundred thousand dollars, an annual financial statement audit or annual financial statement review performed by a certified public accountant. Such reports, audits, and reviews shall be completed and made available upon request to the department; [and]

(2) Demonstrate a positive impact on consumer outcomes regarding the provision of personal care assistance services as evidenced on accurate quarterly and annual service reports submitted to the department;

(3) Implement a quality assurance and supervision process that ensures program compliance and accuracy of records including, but not limited to:

(a) The department of health and senior services shall promulgate by rule a consumer-directed services division provider certification manager course; and

(b) The vendor shall perform ongoing monitoring of the
provision of services in the plan of care and shall assess the
good quality of care being delivered. Such monitoring shall include
at least one annual face-to-face visit and may include electronic
monitoring, telephone checks, written case notes, or other
department-approved methods. The ongoing monitoring shall not
preclude the vendor's responsibility of ongoing diligence of case
management activity oversight;

(4) Comply with all provisions of sections 208.900 to
208.927, and the regulations promulgated thereunder; and

(5) Beginning July 1, 2022, maintain a business location
which shall comply with any and all applicable city, county,
state, and federal requirements.

3. No state or federal funds shall be authorized or
expended to pay for personal care assistance services under
sections 208.900 to 208.927 if any direct employee of the
consumer-directed services vendor conducts the face-to-face home
visit of a consumer for whom such employee is also the personal
care attendant, unless such person provides services solely on a
temporary basis on no more than three days in a thirty-day
period.

208.924. 1. A consumer's personal care assistance services
may be discontinued under circumstances such as the following:

(1) The department learns of circumstances that require
closure of a consumer's case, including one or more of the
following: death, admission into a long-term care facility, no
longer needing service, or inability of the consumer to
consumer-direct personal care assistance service;

(2) The consumer has falsified records; provided false
information of his or her condition, functional capacity, or
level of care needs; or committed fraud;

(3) The consumer is noncompliant with the plan of care.
Noncompliance requires persistent actions by the consumer which
negate the services provided in the plan of care;

(4) The consumer or member of the consumer's household
threatens or abuses the personal care attendant or vendor to the
point where their welfare is in jeopardy and corrective action
has failed;

(5) The maintenance needs of a consumer are unable to
continue to be met because the plan of care hours exceed
availability; and

(6) The personal care attendant is not providing services
as set forth in the personal care assistance services plan and
attempts to remedy the situation have been unsuccessful.

2. The personal care attendant shall report to the
department if he or she witnesses significant deterioration of
the health of the consumer or if he or she has a belief that the
consumer is no longer capable of self-directed care.

208.935. Subject to appropriations, the department of
health and senior services shall develop, or contract with a
state agency or third party to develop an interactive assessment
tool, which may include mobile as well as centralized
functionality, for utilization when implementing the assessment
and authorization process for MO HealthNet home and community-
based services authorized by the division of senior and
disability services.

321.621. 1. For the purposes of this section, "qualified
first responder" shall mean any state and local law enforcement agency staff, fire department personnel, fire district personnel, or licensed emergency medical technician who is acting under the directives and established protocols of a medical director who comes in contact with a person suffering from an anaphylactic reaction and who has received training in recognizing and responding to anaphylactic reactions and the administration of epinephrine auto-injector devices to a person suffering from an apparent anaphylactic reaction. "Qualified first responder agencies" shall mean any state or local law enforcement agency, fire department, or ambulance service that provides documented training to its staff related to the administration of epinephrine auto-injector devices in an apparent anaphylactic reaction.

2. The director of the department of health and senior services, if a licensed physician, may issue a statewide standing order for epinephrine auto-injector devices for adult patients to fire protection districts in nonmetropolitan areas in Missouri as such areas are determined according to the United States Census Bureau's American Community Survey, based on the most recent of five-year period estimate data in which the final year of the estimate ends in either zero or five. If the director of the department of health and senior services is not a licensed physician, the department of health and senior services may employ or contract with a licensed physician who may issue such a statewide order with the express consent of the director.

3. Possession and use of epinephrine auto-injector devices for adult patients shall be limited as follows:
(1) No person shall use an epinephrine auto-injector device pursuant to this section unless such person has successfully completed a training course in the use of epinephrine auto-injector devices for adult patients approved by the director of the department of health and senior services. Nothing in this section shall prohibit the use of an epinephrine auto-injector device:

(a) By a health care professional licensed or certified by this state who is acting within the scope of his or her practice; or

(b) By a person acting pursuant to a lawful prescription;

(2) Every person, firm, organization and entity authorized to possess and use epinephrine auto-injector devices for adult patients pursuant to this section shall use, maintain and dispose of such devices for adult patients in accordance with the rules of the department;

(3) Every use of an epinephrine auto-injector device pursuant to this section shall immediately be reported to the emergency health care provider as defined in section 190.246.

4. (1) Use of an epinephrine auto-injector device pursuant to this section shall be considered first aid or emergency treatment for the purpose of any law relating to liability.

(2) Purchase, acquisition, possession or use of an epinephrine auto-injector device pursuant to this section shall not constitute the unlawful practice of medicine or the unlawful practice of a profession.

(3) Any person otherwise authorized to sell or provide an epinephrine auto-injector device may sell or provide it to a
person authorized to possess it pursuant to this section.

5. (1) There is hereby created in the state treasury the
"Epinephrine Auto-injector Devices for Fire Personnel Fund",
which shall consist of money collected under this section. The
state treasurer shall be custodian of the fund. In accordance
with sections 30.170 and 30.180, the state treasurer may approve
disbursements. The moneys in the fund as set forth in this
section shall be subject to appropriation by the general assembly
for the particular purpose for which collected. The fund shall
be a dedicated fund and money in the fund shall be used solely by
the department of health and senior services for the purposes of
providing epinephrine auto-injector devices for adult patients to
qualified first responder agencies as used in this section.

(2) Notwithstanding the provisions of section 33.080 to the
contrary, any moneys remaining in the fund at the end of the
biennium shall not revert to the credit of the general revenue
fund.

(3) The state treasurer shall invest moneys in the fund in
the same manner as other funds are invested. Any interest and
moneys earned on such investments shall be credited to the fund.

338.035. 1. Every person who desires to be licensed as an
intern pharmacist shall file with the board of pharmacy an
application, on a form to be provided by the board of pharmacy.

2. If an applicant for an intern pharmacist license has
complied with the requirements of this section and with the rules
and regulations of the board of pharmacy and is not denied a
license on any of the grounds listed in section 338.055, the
board of pharmacy may issue to him a license to practice as an
intern pharmacist.

3. Any intern pharmacist who wishes to renew his license shall within thirty days before the license expiration date file an application for a renewal.

4. A licensed intern pharmacist may practice pharmacy only under the direct supervision of a pharmacist licensed by the board; provided, however, that an intern pharmacist working at a remote dispensing site pharmacy may be remotely supervised by a pharmacist working at a supervising pharmacy as provided for in section 338.215.

5. The board of pharmacy shall promulgate rules and regulations which shall further regulate the duties of intern pharmacists and shall set the amount of the fees which shall accompany the license and renewal applications for intern pharmacists.

6. No rule or portion of a rule promulgated under the authority of this chapter shall become effective unless it has been promulgated pursuant to the provisions of section 536.024.

338.210. 1. Pharmacy refers to any location where the practice of pharmacy occurs or such activities are offered or provided by a pharmacist or another acting under the supervision and authority of a pharmacist, including every premises or other place:

(1) Where the practice of pharmacy is offered or conducted;

(2) Where drugs, chemicals, medicines, any legend drugs under 21 U.S.C. Section 353, prescriptions, or poisons are compounded, prepared, dispensed or sold or offered for sale at retail;
(3) Where the words "pharmacist", "apothecary", "drugstore", "drugs", and any other symbols, words or phrases of similar meaning or understanding are used in any form to advertise retail products or services;

(4) Where patient records or other information is maintained for the purpose of engaging or offering to engage in the practice of pharmacy or to comply with any relevant laws regulating the acquisition, possession, handling, transfer, sale or destruction of drugs, chemicals, medicines, prescriptions or poisons;

(5) Where the practice of pharmacy occurs or is offered at a remote dispensing pharmacy site.

2. All activity or conduct involving the practice of pharmacy as it relates to an identifiable prescription or drug order shall occur at the pharmacy location where such identifiable prescription or drug order is first presented by the patient or the patient's authorized agent for preparation or dispensing, unless otherwise expressly authorized by the board.

3. The requirements set forth in subsection 2 of this section shall not be construed to bar the complete transfer of an identifiable prescription or drug order pursuant to a verbal request by or the written consent of the patient or the patient's authorized agent.

4. The board is hereby authorized to enact rules waiving the requirements of subsection 2 of this section and establishing such terms and conditions as it deems necessary, whereby any activities related to the preparation, dispensing or recording of an identifiable prescription or drug order may be shared between
5. If a violation of this chapter or other relevant law occurs in connection with or adjunct to the preparation or dispensing of a prescription or drug order, any permit holder or pharmacist-in-charge at any facility participating in the preparation, dispensing, or distribution of a prescription or drug order may be deemed liable for such violation.

6. Nothing in this section shall be construed to supersede the provisions of section 197.100.

338.215. 1. For purposes of this section, the following terms mean:

(1) "Remote dispensing site pharmacy", any location in this state where the practice of pharmacy occurs and that is licensed as a pharmacy to dispense prescription drugs and is staffed by one or more qualified pharmacy technicians, as defined by the board, or intern pharmacists, whose activities are supervised by a pharmacist at a supervising pharmacy through a continuous real-time audio and video link. "Remote dispensing site pharmacy" does not include the office of a dispensing prescriber or an automated device;

(2) "Supervising pharmacy", a pharmacy licensed in this state under the provisions of chapter 338 that oversees the dispensation activities of a remote dispensing site pharmacy.

2. A supervising pharmacy that operates a remote dispensing site pharmacy, and the remote dispensing site pharmacy, shall be licensed as a pharmacy by the board of pharmacy. The board shall issue a license to a remote dispensing site pharmacy that meets the requirements of this subsection. The remote dispensing site
pharmacy shall:

(1) Submit an application and pay the licensing fee established by the board;

(2) Be jointly owned by a supervising pharmacy; and

(3) Maintain a policy and procedures manual that includes the following:

(a) A description of how the supervising pharmacy and remote dispensing site pharmacy will comply with federal and state laws, rules, and regulations;

(b) The procedure for the supervising pharmacy to supervise the remote dispensing site pharmacy and counsel patients in accordance with the laws of this state prior to the dispensing of a prescription drug under this section;

(c) The procedure for reviewing the prescription drug inventory and drug records maintained by the remote dispensing site pharmacy;

(d) The policy and procedure for providing appropriate security to protect the confidentiality and integrity of patient information;

(e) The written plan for recovery from an event that interrupts or prevents a pharmacist from supervising the operation of the remote dispensing site pharmacy;

(f) The specific duties, tasks, and functions that a registered pharmacy technician or intern pharmacist is authorized to perform at the remote dispensing site pharmacy under the remote supervision of a licensed pharmacist at the supervising pharmacy; and

(g) The procedure for maintaining an up-to-date inventory
of all controlled substances.

3. A remote dispensing site pharmacy shall be under the supervision and control of a supervising pharmacist employed by the supervising pharmacy. The supervising pharmacist shall not be required to be immediately physically present to supervise activities at the remote dispensing site pharmacy, but shall make monthly visits to the remote dispensing site pharmacy in order to ensure compliance with this section.

4. A supervising pharmacist and a remote dispensing site pharmacy shall share common ownership. A pharmacist shall neither be designated nor act as a supervising pharmacist for more than two remote dispensing site pharmacies at one time.

5. A pharmacist at the supervising pharmacy shall verify each prescription before it leaves the remote dispensing site pharmacy. Verification shall occur through the use of technology that includes bar coding and visual review via remote video. As applicable, a pharmacist, intern pharmacist, and pharmacy technician's initials or unique identifier shall appear in the prescription record to identify the name and specific activities of each pharmacist, intern pharmacist, or pharmacy technician involved in the dispensing process.

6. Unless a pharmacist is onsite at the remote dispensing site pharmacy, counseling shall be done by a supervising pharmacist at the supervising pharmacy via a HIPAA-compliant continuous real-time video and audio link before a drug or medical device is released to the patient. The system being used to perform the consultation shall retain the initials or unique identifier of the pharmacist who performs the consultation. The
pharmacist providing counseling under this subsection shall be employed by and located at the supervising pharmacy and have access to all relevant patient information maintained by the remote dispensing site pharmacy.

7. A remote dispensing site pharmacy shall be located at least ten miles from an existing retail pharmacy unless:
   (1) The remote dispensing site pharmacy is part of a community mental health center, federally qualified health center, hospital, rural health clinic, or outpatient clinic setting; or
   (2) An applicant of a proposed remote dispensing site pharmacy demonstrates to the board how the proposed remote dispensing site pharmacy will promote public health.

8. The remote dispensing pharmacy shall be staffed by a pharmacist at least eight hours a month and shall reconcile the up-to-date controlled substance inventory twice a month. The supervising pharmacist may provide services as allowed in section 338.010 and as provided by policies and procedures.

9. If the average number of prescriptions dispensed per day by the remote dispensing site pharmacy exceeds one hundred fifty prescriptions, the remote dispensing site pharmacy shall, within ten days, apply to the board for licensure as a class A, B, or C pharmacy, as applicable. The average number of prescriptions dispensed per day shall be determined by averaging the number of prescriptions dispensed per day over the previous ninety-day period.

10. Unless otherwise approved by the board, the supervising pharmacy shall be located in this state and within fifty road
miles of a remote dispensing site pharmacy to ensure that the remote dispensing site pharmacy is sufficiently supported by the supervising pharmacy and that necessary personnel or supplies may be delivered to the remote dispensing site pharmacy within a reasonable period of time of an identified need.

11. The board of pharmacy may promulgate all necessary rules and regulations for the implementation of this section, provided that no such rules and regulations shall restrict the practice of pharmacy at a remote dispensing site pharmacy. Any rule or portion of a rule, as that term is defined in section 536.010, that is created under the authority delegated in this section shall become effective only if it complies with and is subject to all of the provisions of chapter 536 and, if applicable, section 536.028. This section and chapter 536 are nonseverable, and if any of the powers vested with the general assembly pursuant to chapter 536 to review, to delay the effective date, or to disapprove and annul a rule are subsequently held unconstitutional, then the grant of rulemaking authority and any rule proposed or adopted after August 28, 2020, shall be invalid and void.

338.220. 1. It shall be unlawful for any person, copartnership, association, corporation or any other business entity to open, establish, operate, or maintain any pharmacy as defined by statute without first obtaining a permit or license to do so from the Missouri board of pharmacy. A permit shall not be required for an individual licensed pharmacist to perform nondispensing activities outside of a pharmacy, as provided by the rules of the board. A permit shall not be required for an
1 individual licensed pharmacist to administer drugs, vaccines, and
2 biologicals by protocol, as permitted by law, outside of a
3 pharmacy. The following classes of pharmacy permits or licenses
4 are hereby established:

5    (1) Class A: Community/ambulatory;
6    (2) Class B: Hospital pharmacy;
7    (3) Class C: Long-term care;
8    (4) Class D: Nonsterile compounding;
9    (5) Class E: Radio pharmaceutical;
10    (6) Class F: Renal dialysis;
11    (7) Class G: Medical gas;
12    (8) Class H: Sterile product compounding;
13    (9) Class I: Consultant services;
14    (10) Class J: Shared service;
15    (11) Class K: Internet;
16    (12) Class L: Veterinary;
17    (13) Class M: Specialty (bleeding disorder);
18    (14) Class N: Automated dispensing system (health care
19        facility);
20    (15) Class O: Automated dispensing system (ambulatory
21        care);
22    (16) Class P: Practitioner office/clinic;
23    (17) Class Q: Charitable pharmacy; and
24    (18) Class R: Remote dispensing site pharmacy.
25 2. Application for such permit or license shall be made
26 upon a form furnished to the applicant; shall contain a statement
27 that it is made under oath or affirmation and that its
28 representations are true and correct to the best knowledge and
belief of the person signing same, subject to the penalties of
making a false affidavit or declaration; and shall be accompanied
by a permit or license fee. The permit or license issued shall
be renewable upon payment of a renewal fee. Separate
applications shall be made and separate permits or licenses
required for each pharmacy opened, established, operated, or
maintained by the same owner.

3. All permits, licenses or renewal fees collected pursuant
to the provisions of sections 338.210 to 338.370 shall be
deposited in the state treasury to the credit of the Missouri
board of pharmacy fund, to be used by the Missouri board of
pharmacy in the enforcement of the provisions of sections 338.210
to 338.370, when appropriated for that purpose by the general
assembly.

4. Class L: veterinary permit shall not be construed to
prohibit or interfere with any legally registered practitioner of
veterinary medicine in the compounding, administering,
prescribing, or dispensing of their own prescriptions, or
medicine, drug, or pharmaceutical product to be used for animals.

5. Except for any legend drugs under 21 U.S.C. Section 353,
the provisions of this section shall not apply to the sale,
dispensing, or filling of a pharmaceutical product or drug used
for treating animals.

6. A "class B hospital pharmacy" shall be defined as a
pharmacy owned, managed, or operated by a hospital as defined by
section 197.020 or a clinic or facility under common control,
management or ownership of the same hospital or hospital system.
This section shall not be construed to require a class B hospital
pharmacy permit or license for hospitals solely providing services within the practice of pharmacy under the jurisdiction of, and the licensure granted by, the department of health and senior services under and pursuant to chapter 197.

7. Upon application to the board, any hospital that holds a pharmacy permit or license on August 28, 2014, shall be entitled to obtain a class B pharmacy permit or license without fee, provided such application shall be submitted to the board on or before January 1, 2015.

338.260. 1. No person shall carry on, conduct or transact a business under a name which contains as part of the name the words "pharmacist", "pharmacy", "apothecary", "apothecary shop", "chemist shop", "drug store", "druggist", "drugs", "consultant pharmacist", or any word of similar or like import, unless the place of business is supervised by a licensed pharmacist.

2. Nothing in this chapter shall be construed to prevent any person from using a historical name in reference to any building, structure, or business so long as the person is not engaged in the practice of pharmacy as defined in section 338.010.

3. Notwithstanding the provisions of subsection 2 of this section, the board of pharmacy shall retain authority to enforce the provisions of subsection 1 of this section against any person offering for sale any naturopathic or homeopathic service or any herbal, nutritional, vitamin, dietary, mineral, or other supplement intended for human application, absorption, or consumption.

4. Supervision of a licensed remote dispensing site
pharmacy shall not require a pharmacist to be physically present at the remote dispensing site pharmacy location, provided that dispensing activities are supervised by a supervising pharmacist located at a Missouri-licensed supervising pharmacy through the use of a continuous real-time audio and video link.

376.383. 1. For purposes of this section and section 376.384, the following terms shall mean:

(1) "Claimant", any individual, corporation, association, partnership or other legal entity asserting a right to payment arising out of a contract or a contingency or loss covered under a health benefit plan as defined in section 376.1350;

(2) "Clean claim", a claim that has no defect, impropriety, lack of any required substantiating documentation, or particular circumstance requiring special treatment that prevents timely payment;

(3) "Deny" or "denial", when the health carrier refuses to reimburse all or part of the claim;

(4) "Health care provider", health care provider as defined in section 376.1350;

(5) "Health care services", health care services as defined in section 376.1350;

(6) "Health carrier", health carrier as defined in section 376.1350 and any self-insured health plan, to the extent allowed by federal law; except that health carrier shall not include a workers' compensation carrier providing benefits to an employee pursuant to chapter 287. For the purposes of this section and section 376.384, third-party contractors are health carriers;

(7) "Processing days", number of days the health carrier or
any of its agents, subsidiaries, contractors, subcontractors, or third-party contractors has the claim in its possession. Processing days shall not include days in which the health carrier is waiting for a response to a request for additional information from the claimant;

(8) "Request for additional information", a health carrier's electronic or facsimile request for additional information from the claimant specifying all of the documentation or information necessary to process all of the claim, or all of the claim on a multi-claim form, as a clean claim for payment;

(9) "Third-party contractor", a third party contracted with the health carrier to receive or process claims for reimbursement of health care services.

2. Within forty-eight hours after receipt of an electronically filed claim by a health carrier or a third-party contractor, a health carrier shall send an electronic acknowledgment of the date of receipt.

3. Within thirty processing days after receipt of a filed claim by a health carrier or a third-party contractor, a health carrier shall send an electronic or facsimile notice of the status of the claim that notifies the claimant:

(1) Whether the claim is a clean claim as defined under this section; or

(2) The claim requires additional information from the claimant.

If the claim is a clean claim, then the health carrier shall pay or deny the claim. If the claim requires additional information,
the health carrier shall include in the notice a request for additional information. If a health carrier pays the claim, this subsection shall not apply.

4. Within ten processing days after receipt of additional information by a health carrier or a third-party contractor, a health carrier shall pay the claim or any undisputed part of the claim in accordance with this section or send an electronic or facsimile notice of receipt and status of the claim:
   (1) That denies all or part of the claim and specifies each reason for denial; or
   (2) That makes a final request for additional information.

5. Within five processing days after the day on which the health carrier or a third-party contractor receives the additional requested information in response to a final request for information, it shall pay the claim or any undisputed part of the claim or deny the claim.

6. (1) If the health carrier has not paid the claimant on or before the forty-fifth processing day from the date of receipt of the claim, the health carrier shall pay the claimant one percent interest per month and a penalty in an amount equal to one percent of the claim per day. **On claims where the amount owed by a health carrier exceeds thirty-five thousand dollars on the unpaid balance of a claim, the health carrier shall pay the claimant one percent interest per month and a penalty in an amount equal to one percent of the claim per day for a maximum of one hundred days, and thereafter shall pay the claimant two percent interest per month.** The interest and penalty shall be calculated based upon the unpaid balance of the claim as of the
forty-fifth processing day. The interest and penalty paid pursuant to this subsection shall be included in any late reimbursement without the necessity for the person that filed the original claim to make an additional claim for that interest and penalty. A health carrier may combine interest payments and make payment once the aggregate amount reaches one hundred dollars.

(2) Any claim or portion of a claim which has been properly denied before the forty-fifth processing day under this section and section 376.384 shall not be subject to interest or penalties. For a claim or any portion of such claim that was denied before the forty-fifth processing day, interest and penalties shall begin to accrue beginning on the date the first appeal is filed by the claimant with the health carrier until such claim is paid, if the claim or portion of the claim is approved. If any appeal filed with the health carrier does not result in the disputed claim or portion of such claim being approved for payment to the claimant, and a petition is filed in a court of competent jurisdiction to recover payment of all or part of such claim, interest and penalties shall continue to accrue for no more than one hundred days from the day the first appeal was filed by the claimant with the health carrier, and such interest and penalties shall [cease] continue to accrue [on the day] ten days after [a petition is filed in] a court of competent jurisdiction [to recover payment of such claim] finds that the claim or portion of the claim shall be paid to the claimant. Upon a finding by a court of competent jurisdiction that the health carrier failed to pay a claim, interest, or penalty without good cause, the court shall enter judgment for
reasonable attorney fees for services necessary for recovery.
Upon a finding that a health care provider filed suit without
reasonable grounds to recover a claim, the court shall award the
health carrier reasonable attorney fees necessary to the defense.

7. The department of commerce and insurance shall monitor
denials and determine whether the health carrier acted
reasonably.

8. If a health carrier or third-party contractor has
reasonable grounds to believe that a fraudulent claim is being
made, the health carrier or third-party contractor shall notify
the department of commerce and insurance of the fraudulent claim
pursuant to sections 375.991 to 375.994.

9. Denial of a claim shall be communicated to the claimant
and shall include the specific reason why the claim was denied.
Any claim for which the health carrier has not communicated a
specific reason for the denial shall not be considered denied
under this section or section 376.384.

10. Requests for additional information shall specify all
of the documentation and additional information that is necessary
to process all of the claim, or all of the claims on a
multi-claim form, as a clean claim for payment. Information
requested shall be reasonable and pertain solely to the health
carrier's liability. The health carrier shall acknowledge
receipt of the requested additional information to the claimant
within five calendar days or pay the claim.

376.387. 1. For purposes of this section, the following
terms shall mean:

(1) "Covered person", the same meaning as such term is
defined in section 376.1257;

(2) "Health benefit plan", the same meaning as such term is defined in section 376.1350;

(3) "Health carrier" or "carrier", the same meaning as such term is defined in section 376.1350;

(4) "Pharmacy", the same meaning as such term is defined in chapter 338;

(5) "Pharmacy benefits manager", the same meaning as such term is defined in section 376.388.

2. No pharmacy benefits manager shall include a provision in a contract entered into or modified on or after August 28, 2018, with a pharmacy or pharmacist that requires a covered person to make a payment for a prescription drug at the point of sale in an amount that exceeds the lesser of:

(1) The copayment amount as required under the health benefit plan; or

(2) The amount an individual would pay for a prescription if that individual paid with cash.

3. A pharmacy or pharmacist shall have the right to provide to a covered person information regarding the amount of the covered person's cost share for a prescription drug, the covered person's cost of an alternative drug, and the covered person's cost of the drug without adjudicating the claim through the pharmacy benefits manager. Neither a pharmacy nor a pharmacist shall be proscribed by a pharmacy benefits manager from discussing any such information or from selling a more affordable alternative to the covered person.

4. No pharmacy benefits manager shall, directly or
indirectly, charge or hold a pharmacist or pharmacy responsible for any fee amount related to a claim that is not known at the time of the claim's adjudication, unless the amount is a result of improperly paid claims or charges for administering a health benefit plan.

5. This section shall not apply with respect to claims under Medicare Part D, or any other plan administered or regulated solely under federal law, and to the extent this section may be preempted under the Employee Retirement Income Security Act of 1974 for self-funded employer-sponsored health benefit plans.

6. A pharmacy benefits manager shall notify in writing any health carrier with which it contracts if the pharmacy benefits manager has a conflict of interest, any commonality of ownership, or any other relationship, financial or otherwise, between the pharmacy benefits manager and any other health carrier with which the pharmacy benefits manager contracts.

7. The department of commerce and insurance shall enforce this section.

376.393. 1. As used in this section, the following terms shall mean:

(1) "Health carrier" or "carrier", the same meaning as is ascribed to such term in section 376.1350;

(2) "Pharmacy benefits manager", the same meaning as is ascribed to such term in section 376.388.

2. No entity subject to the jurisdiction of this state shall act as a pharmacy benefits manager without a license issued by the department. The department shall establish by rule the
application process and license fee for pharmacy benefits managers.

3. The department may cause a complaint to be filed with the administrative hearing commission as provided in chapter 621 against any holder of a license issued under this section for:

   (1) Violation of the laws or regulations of any state or of the United States, where the offense is reasonably related to the qualifications, functions, or duties of a pharmacy benefit manager, including, but not limited to, where an essential element of the offense is fraud, dishonesty, or an act of violence, or where the offense involves moral turpitude, or where the offense involves failure to comply with a requirement of this chapter, whether or not sentence or penalty is imposed;

   (2) Use of fraud, deception, misrepresentation, or bribery for any reason;

   (3) Obtaining or attempting to obtain any fee, charge, tuition, or other compensation by fraud, deception, or misrepresentation;

   (4) Incompetence, misconduct, gross negligence, or dishonesty in the performance of the functions or duties of a pharmacy benefits manager or other regulated profession or activity; or

   (5) Disciplinary action taken against the holder of a license or other right to practice as a pharmacy benefits manager or other regulated profession.

After the filing of such complaint, the proceedings shall be conducted in accordance with the provisions of chapter 621. Upon
a finding by the administrative hearing commission that grounds provided in this subsection for disciplinary action are met, the department may, singly or in combination, censure or place the person named in the complaint on probation with such terms and conditions as the department deems appropriate for a period not to exceed five years, or may suspend, for a period not to exceed three years, or revoke the license, certificate, or permit. An individual whose license has been revoked shall wait at least one year from the date of revocation to apply for relicensure. Relicensure shall be at the discretion of the department.

376.945. 1. The department shall, as a condition of the issuance of a certificate of authority pursuant to section 376.935, require that the provider establish a reserve of an amount equal to at least fifty percent of any entrance fee paid by the first occupant of a living unit under a life care contract. The reserve shall be maintained by the provider on a current basis, in escrow with a bank, trust company, or other escrow agent approved by the department. [Such] The entire amount of such reserve shall be amortized and earned by and available for release to the provider at the rate of one percent per month on the balance of the reserve, provided, however, that at no time shall the entrance fee reserve together with all interest earned thereon total less than an amount equal to one [and one-half times the percentage] hundred percent of the annual long-term debt principal and interest payments of the provider applicable only to living units occupied under life care contracts. Such portion of each entrance fee as is necessary to maintain the entrance fee reserve as set forth herein shall be
paid to the reserve fund for the second and all subsequent
occupancies of a living unit occupied under a life care contract.
The requirements of this subsection may be met in whole or in
part by other reserve funds held for the purpose of meeting loan
obligations, provided that the total amount equals or exceeds the
amount required under this subsection.

2. In addition, each provider shall establish and maintain
separately for each facility, a reserve equal to not less than
five percent of the facility's total outstanding balance of
contractually obligated move-out refunds at the close of each
fiscal year. [All reserves required hereunder for move-out
refunds]

3. All reserve funds held under subsections 1 or 2 of this
section shall be held in liquid assets consisting of federal
government or other marketable securities, deposits, or accounts
insured by the federal government.

4. This section shall be applicable only to life care
contracts executed for occupancy of living units constructed
after September 28, 1981.

376.1578. 1. Within two working days after receipt of a
[faxed or mailed completed] credentialing application, the health
carrier shall send a notice of receipt to the practitioner. A
health carrier shall provide access to a provider web portal that
allows the practitioner to receive notice of the status of an
electronically submitted application.

2. If a health carrier determines the application is not a
completed application, the health carrier shall have ten days
from the date the notice of receipt was sent as required in
subsection 1 of this section to request any additional information from the practitioner. The application shall be considered a completed application upon receipt of the requested additional information from the practitioner. Within two working days of receipt of the requested additional information, the health carrier shall send a notice to the practitioner informing him or her that he or she has submitted a completed application. If the health carrier does not request additional information, the application shall be deemed completed as of the date the notice of receipt was sent as required under subsection 1 of this section.

3. A health carrier shall assess a health care practitioner's completed credentialing application and make a decision as to whether to approve or deny the practitioner's credentialing application and notify the practitioner of such decision within sixty [business] days of the date of receipt of the completed application. The sixty-day deadline established in this section shall not apply if the application or subsequent verification of information indicates that the practitioner has:

(1) A history of behavioral disorders or other impairments affecting the practitioner's ability to practice, including but not limited to substance abuse;

(2) Licensure disciplinary actions against the practitioner's license to practice imposed by any state or territory or foreign jurisdiction;

(3) Had the practitioner's hospital admitting or surgical privileges or other organizational credentials or authority to
practice revoked, restricted, or suspended based on the practitioner's clinical performance; or

(4) A judgment or judicial award against the practitioner arising from a medical malpractice liability lawsuit.

4. If a practitioner's application is approved, the health carrier shall provide payments for covered health services performed by the practitioner during the credentialing period if the provision of services was on behalf of an entity that had a contract with such health carrier during the credentialing period. The contracted entity for which the practitioner is providing services shall submit to the health carrier all claims for services provided by such practitioner during the credentialing period within six months after the health carrier has approved that practitioner's credentialing application. Claims submitted for reimbursement under this section shall be sent to the carrier by the provider in a single request or as few requests as practical subject to any technical constraints or other issues out of the contracted provider's control.

"Credentialing period" shall mean the time between the date the practitioner submits a completed application to the health carrier to be credentialed and the date the practitioner's credentialing is approved by the health carrier.

5. A health carrier shall not require a practitioner to be credentialed in order to receive payments for covered patient care services if the practitioner is providing coverage for an absent credentialed practitioner during a temporary period of time not to exceed sixty days. Any practitioner authorized to receive payments for covered services under this section shall
provide notice to the health carrier, including, but not limited
to, the absent practitioner's name, medical license information,
and estimated duration of absence and the name and medical
license information of the practitioner providing coverage for
such absent credentialed practitioner. A health carrier may deny
payments if the practitioner providing services in lieu of the
credentialed provider meets one of the conditions in subdivisions
(1) to (4) of subsection 3 of this section.

6. All claims eligible for payment under subsection 4 or 5
of this section shall be subject to section 376.383.

7. For the purposes of this section, "covered health
services" shall mean any services provided by a practitioner that
would otherwise be covered if provided by a credentialed
provider.

[3.] 8. The department of commerce and insurance shall
establish a mechanism for reporting alleged violations of this
section to the department.

579.060. 1. A person commits the offense of unlawful sale,
distribution, or purchase of over-the-counter methamphetamine
precursor drugs if he or she knowingly:

(1) Sells, distributes, dispenses, or otherwise provides
any number of packages of any drug product containing detectable
amounts of ephedrine, phenylpropanolamine, or pseudoephedrine, or
any of their salts, optical isomers, or salts of optical isomers,
in a total amount greater than [nine] seven and two-tenths grams
to the same individual within a thirty-day period, unless the
amount is dispensed, sold, or distributed pursuant to a valid
prescription; or
(2) Purchases, receives, or otherwise acquires within a thirty-day period any number of packages of any drug product containing any detectable amount of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts or optical isomers, or salts of optical isomers in a total amount greater than [nine] seven and two-tenths grams, without regard to the number of transactions, unless the amount is purchased, received, or acquired pursuant to a valid prescription; or

(3) Purchases, receives, or otherwise acquires within a twenty-four-hour period any number of packages of any drug product containing any detectable amount of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts or optical isomers, or salts of optical isomers in a total amount greater than three and six-tenths grams, without regard to the number of transactions, unless the amount is purchased, received, or acquired pursuant to a valid prescription; or

(4) Sells, distributes, dispenses, or otherwise provides any number of packages of any drug product containing detectable amounts of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts, optical isomers, or salts of optical isomers, in a total amount greater than forty-three and two-tenths grams to the same individual within a twelve-month period, unless the amount is dispensed, sold, or distributed pursuant to a valid prescription; or

(5) Purchases, receives, or otherwise acquires within a twelve-month period any number of packages of any drug product containing any detectable amount of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts or
optical isomers, or salts of optical isomers in a total amount greater than forty-three and two-tenths grams, without regard to the number of transactions, unless the amount is purchased, received, or acquired pursuant to a valid prescription; or

(6) Dispenses or offers drug products that are not excluded from Schedule V in subsection 17 or 18 of section 195.017 and that contain detectable amounts of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts, optical isomers, or salts of optical isomers, without ensuring that such products are located behind a pharmacy counter where the public is not permitted and that such products are dispensed by a registered pharmacist or pharmacy technician under subsection 11 of section 195.017; or

[(5)] (7) Holds a retail sales license issued under chapter 144 and knowingly sells or dispenses packages that do not conform to the packaging requirements of section 195.418.

2. A pharmacist, intern pharmacist, or registered pharmacy technician commits the offense of unlawful sale, distribution, or purchase of over-the-counter methamphetamine precursor drugs if he or she knowingly:

(1) Sells, distributes, dispenses, or otherwise provides any number of packages of any drug product containing detectable amounts of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts or optical isomers, or salts of optical isomers, in a total amount greater than three and six-tenth grams to the same individual within a twenty-four hour period, unless the amount is dispensed, sold, or distributed pursuant to a valid prescription; or
(2) Fails to submit information under subsection 13 of section 195.017 and subsection [5] of section 195.417 about the sales of any compound, mixture, or preparation of products containing detectable amounts of ephedrine, phenylpropanolamine, or pseudoephedrine, or any of their salts, optical isomers, or salts of optical isomers, in accordance with transmission methods and frequency established by the department of health and senior services; or

(3) Fails to implement and maintain an electronic log, as required by subsection 12 of section 195.017, of each transaction involving any detectable quantity of pseudoephedrine, its salts, isomers, or salts of optical isomers or ephedrine, its salts, optical isomers, or salts of optical isomers; or

(4) Sells, distributes, dispenses or otherwise provides to an individual under eighteen years of age without a valid prescription any number of packages of any drug product containing any detectable quantity of pseudoephedrine, its salts, isomers, or salts of optical isomers, or ephedrine, its salts or optical isomers, or salts of optical isomers.

3. Any person who violates the packaging requirements of section 195.418 and is considered the general owner or operator of the outlet where ephedrine, pseudoephedrine, or phenylpropanolamine products are available for sale shall not be penalized if he or she documents that an employee training program was in place to provide the employee who made the unlawful retail sale with information on the state and federal regulations regarding ephedrine, pseudoephedrine, or phenylpropanolamine.
4. The offense of unlawful sale, distribution, or purchase of over-the-counter methamphetamine precursor drugs is a class A misdemeanor.

610.100. 1. As used in sections 610.100 to 610.150, the following words and phrases shall mean:

   (1) "Arrest", an actual restraint of the person of the defendant, or by his or her submission to the custody of the officer, under authority of a warrant or otherwise for a criminal violation which results in the issuance of a summons or the person being booked;

   (2) "Arrest report", a record of a law enforcement agency of an arrest and of any detention or confinement incident thereto together with the charge therefor;

   (3) "Inactive", an investigation in which no further action will be taken by a law enforcement agency or officer for any of the following reasons:

      (a) A decision by the law enforcement agency not to pursue the case;

      (b) Expiration of the time to file criminal charges pursuant to the applicable statute of limitations, or ten years after the commission of the offense; whichever date earliest occurs;

      (c) Finality of the convictions of all persons convicted on the basis of the information contained in the investigative report, by exhaustion of or expiration of all rights of appeal of such persons;

   (4) "Incident report", a record of a law enforcement agency consisting of the date, time, specific location, name of the
victim and immediate facts and circumstances surrounding the
initial report of a crime or incident, including any logs of
reported crimes, accidents and complaints maintained by that
agency;

(5) "Investigative report", a record, other than an arrest
or incident report, prepared by personnel of a law enforcement
agency, inquiring into a crime or suspected crime, either in
response to an incident report or in response to evidence
developed by law enforcement officers in the course of their
duties;

(6) "Mobile video recorder", any system or device that
captures visual signals that is capable of installation and being
installed in a vehicle or being worn or carried by personnel of a
law enforcement agency and that includes, at minimum, a camera
and recording capabilities;

(7) "Mobile video recording", any data captured by a mobile
video recorder, including audio, video, and any metadata;

(8) "Nonpublic location", a place where one would have a
reasonable expectation of privacy, including, but not limited to
a dwelling, school, or medical facility.

2. (1) Each law enforcement agency of this state, of any
county, and of any municipality shall maintain records of all
incidents reported to the agency, investigations and arrests made
by such law enforcement agency. All incident reports and arrest
reports shall be open records.

(2) Notwithstanding any other provision of law other than
the provisions of subsections 4, 5 and 6 of this section or
section 320.083, mobile video recordings and investigative
reports of all law enforcement agencies and any reports or
records in the possession of the department of health and senior
services' Missouri state public health laboratory, which were the
result of testing performed at the request of any municipal,
county, state, or federal law enforcement agency, are closed
records until the investigation becomes inactive.

(3) If any person is arrested and not charged with an
offense against the law within thirty days of the person's
arrest, the arrest report shall thereafter be a closed record
except that the disposition portion of the record may be accessed
and except as provided in section 610.120.

(4) Except as provided in subsections 3 and 5 of this
section, a mobile video recording that is recorded in a nonpublic
location is authorized to be closed, except that any person who
is depicted in the recording or whose voice is in the recording,
a legal guardian or parent of such person if he or she is a
minor, a family member of such person within the first degree of
consanguinity if he or she is deceased or incompetent, an
attorney for such person, or insurer of such person, upon written
request, may obtain a complete, unaltered, and unedited copy of a
recording under and pursuant to this section.

3. Except as provided in subsections 4, 5, 6 and 7 of this
section, if any portion of a record or document of a law
enforcement officer or agency, other than an arrest report, which
would otherwise be open, contains information that is reasonably
likely to pose a clear and present danger to the safety of any
victim, witness, undercover officer, or other person; or
jeopardize a criminal investigation, including records which
would disclose the identity of a source wishing to remain confidential or a suspect not in custody; or which would disclose techniques, procedures or guidelines for law enforcement investigations or prosecutions, that portion of the record shall be closed and shall be redacted from any record made available pursuant to this chapter.

4. Any person, including a legal guardian or a parent of such person if he or she is a minor, family member of such person within the first degree of consanguinity if such person is deceased or incompetent, attorney for a person, or insurer of a person involved in any incident or whose property is involved in an incident, may obtain any records closed pursuant to this section or section 610.150 for purposes of investigation of any civil claim or defense, as provided by this subsection. Any individual, legal guardian or parent of such person if he or she is a minor, his or her family member within the first degree of consanguinity if such individual is deceased or incompetent, his or her attorney or insurer, involved in an incident or whose property is involved in an incident, upon written request, may obtain a complete unaltered and unedited incident report concerning the incident, and may obtain access to other records closed by a law enforcement agency pursuant to this section. Within thirty days of such request, the agency shall provide the requested material or file a motion pursuant to this subsection with the circuit court having jurisdiction over the law enforcement agency stating that the safety of the victim, witness or other individual cannot be reasonably ensured, or that a criminal investigation is likely to be jeopardized. If, based on
such motion, the court finds for the law enforcement agency, the
court shall either order the record closed or order such portion
of the record that should be closed to be redacted from any
record made available pursuant to this subsection.

5. (1) Any person may bring an action pursuant to this
section in the circuit court having jurisdiction to authorize
disclosure of a mobile video recording or the information
contained in an investigative report of any law enforcement
agency, which would otherwise be closed pursuant to this section.
The court may order that all or part of a mobile video recording
or the information contained in an investigative report be
released to the person bringing the action.

(2) In making the determination as to whether information
contained in an investigative report shall be disclosed, the
court shall consider whether the benefit to the person bringing
the action or to the public outweighs any harm to the public, to
the law enforcement agency or any of its officers, or to any
person identified in the investigative report in regard to the
need for law enforcement agencies to effectively investigate and
prosecute criminal activity.

(3) In making the determination as to whether a mobile
video recording shall be disclosed, the court shall consider:

(a) Whether the benefit to the person bringing the action
or the benefit to the public outweighs any harm to the public, to
the law enforcement agency or any of its officers, or to any
person identified in the mobile video recording in regard and
with respect to the need for law enforcement agencies to
effectively investigate and prosecute criminal activity;
(b) Whether the mobile video recording contains information that is reasonably likely to disclose private matters in which the public has no legitimate concern;

(c) Whether the mobile video recording is reasonably likely to bring shame or humiliation to a person of ordinary sensibilities; and

(d) Whether the mobile video recording was taken in a place where a person recorded or depicted has a reasonable expectation of privacy.

(4) The mobile video recording or investigative report in question may be examined by the court in camera.

(5) If the disclosure is authorized in whole or in part, the court may make any order that justice requires, including one or more of the following:

(a) That the mobile video recording or investigative report may be disclosed only on specified terms and conditions, including a designation of the time or place;

(b) That the mobile video recording or investigative report may be had only by a method of disclosure other than that selected by the party seeking such disclosure and may be disclosed to the person making the request in a different manner or form as requested;

(c) That the scope of the request be limited to certain matters;

(d) That the disclosure occur with no one present except persons designated by the court;

(e) That the mobile video recording or investigative report be redacted to exclude, for example, personally identifiable
(f) That a trade secret or other confidential research, development, or commercial information not be disclosed or be disclosed only in a designated way.

(6) The court may find that the party seeking disclosure of the mobile video recording or the investigative report shall bear the reasonable and necessary costs and attorneys' fees of both parties, unless the court finds that the decision of the law enforcement agency not to open the mobile video recording or investigative report was substantially unjustified under all relevant circumstances, and in that event, the court may assess such reasonable and necessary costs and attorneys' fees to the law enforcement agency.

6. Any person may apply pursuant to this subsection to the circuit court having jurisdiction for an order requiring a law enforcement agency to open incident reports and arrest reports being unlawfully closed pursuant to this section. If the court finds by a preponderance of the evidence that the law enforcement officer or agency has knowingly violated this section, the officer or agency shall be subject to a civil penalty in an amount up to one thousand dollars. If the court finds that there is a knowing violation of this section, the court may order payment by such officer or agency of all costs and attorneys' fees, as provided by section 610.027. If the court finds by a preponderance of the evidence that the law enforcement officer or agency has purposely violated this section, the officer or agency shall be subject to a civil penalty in an amount up to five thousand dollars and the court shall order payment by such
officer or agency of all costs and attorney fees, as provided in section 610.027. The court shall determine the amount of the penalty by taking into account the size of the jurisdiction, the seriousness of the offense, and whether the law enforcement officer or agency has violated this section previously.

7. The victim of an offense as provided in chapter 566 may request that his or her identity be kept confidential until a charge relating to such incident is filed.

8. Any person who requests and receives a mobile video recording that was recorded in a nonpublic location under and pursuant to this section is prohibited from displaying or disclosing the mobile video recording, including any description or account of any or all of the mobile video recording, without first providing direct third-party notice to each person not affiliated with a law enforcement agency or each non-law enforcement agency individual whose image or sound is contained in the recording, and affording, upon receiving such notice, each person appearing and whose image or sound is contained in the mobile video recording no less than ten days to file and serve an action seeking an order from a court of competent jurisdiction to enjoin all or some of the intended display, disclosure, description, or account of the recording. Any person who fails to comply with the provisions of this subsection is subject to damages in a civil action proceeding.

Section B. Because immediate action is necessary to ensure that all owners, officers, managers, contractors, employees, and other support staff of medical marijuana facilities be subjected to state and federal fingerprint-based criminal background checks
to insure the integrity of the Missouri medical marijuana industry, the enactment of section 195.815 of this act is deemed necessary for the immediate preservation of the public health, welfare, peace, and safety, and is hereby declared to be an emergency act within the meaning of the constitution, and the enactment of section 195.815 of this act shall be in full force and effect on July 1, 2020, or upon its passage and approval, whichever occurs later.