

First Regular Session of the 121st General Assembly (2019)

PRINTING CODE. Amendments: Whenever an existing statute (or a section of the Indiana Constitution) is being amended, the text of the existing provision will appear in this style type, additions will appear in **this style type**, and deletions will appear in ~~this style type~~.

Additions: Whenever a new statutory provision is being enacted (or a new constitutional provision adopted), the text of the new provision will appear in **this style type**. Also, the word **NEW** will appear in that style type in the introductory clause of each SECTION that adds a new provision to the Indiana Code or the Indiana Constitution.

Conflict reconciliation: Text in a statute in *this style type* or ~~this style type~~ reconciles conflicts between statutes enacted by the 2018 Regular and Special Session of the General Assembly.

## HOUSE ENROLLED ACT No. 1542

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AN ACT to amend the Indiana Code concerning human services.

*Be it enacted by the General Assembly of the State of Indiana:*

SECTION 1. IC 12-8-1.5-17.5 IS ADDED TO THE INDIANA CODE AS A **NEW** SECTION TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: **Sec. 17.5. (a) Before October 1, 2019, the office of the secretary of family and social services shall prepare and submit a report as described in subsection (b) to the legislative council in an electronic format under IC 5-14-6.**

**(b) The office of the secretary shall conduct a comprehensive study of the health programs that the office of the secretary administers or oversees, including programs administered by managed care programs under IC 12-15-12 and programs contracted with the office of Medicaid policy and planning. The report must:**

- (1) identify administrative and reporting requirements by health providers under contract with the office of the secretary that are unnecessary or overly burdensome; and**
- (2) include recommendations for reductions in administrative burdens related to the administration and oversight described in this subsection.**

**(c) This section expires July 1, 2020.**

SECTION 2. IC 12-12.7-2-22 IS ADDED TO THE INDIANA CODE AS A **NEW** SECTION TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: **Sec. 22. Notwithstanding any other**

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law, any appropriation made to a program established under this chapter and 20 U.S.C. 1431 through 1444 (first steps program) that exceeds eleven million three hundred thirty-nine thousand sixty-three dollars (\$11,339,063) in a state fiscal year must be distributed by the office of the secretary of family and social services as follows:

- (1) Not more than ten percent (10%) to the division of disability and rehabilitative services for infrastructure expenses.
- (2) Not less than forty percent (40%) to systems point of entry contracts.
- (3) Not less than fifty percent (50%) to rates of providers who provide services under this chapter and 20 U.S.C. 1431 through 1444.

SECTION 3. IC 12-15-12-23 IS ADDED TO THE INDIANA CODE AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: **Sec. 23. A managed care organization may not require a psychiatrist who is licensed under IC 25-22.5 to be certified by the American Board of Psychiatry and Neurology for purposes of credentialing or contracting with the psychiatrist while the psychiatrist is practicing at a community mental health center.**

SECTION 4. IC 16-21-1-7, AS AMENDED BY SEA 575-2019, SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: **Sec. 7. (a) The executive board may adopt rules under IC 4-22-2 necessary to protect the health, safety, rights, and welfare of patients, including the following:**

- (1) Rules pertaining to the operation and management of hospitals, ambulatory outpatient surgical centers, abortion clinics, and birthing centers.
- (2) Rules establishing standards for equipment, facilities, and staffing required for efficient and quality care of patients.

(b) Notwithstanding 410 IAC 15-1.7-1 and 410 IAC 15-2.7-1, the following apply to a publication that is referred to in 410 IAC 15:

- (1) The Guidelines for Construction and Equipment of Hospital and Medical Facilities refers to the **following**:
  - (A) **The 2018 edition or most recent publication of the Guidelines for Design and Construction of Hospitals.**
  - (B) **The 2018 edition or most recent publication of the Guidelines for Design and Construction of Outpatient Facilities.**
- (2) **The National Fire Protection Association (NFPA) 101, Life**



**Safety Code Handbook publication refers to the 2018 edition or most recent publication.**

~~(2)~~ **(3)** The National Fire Protection Association 99, Health Care Facilities publication refers to the 2018 edition or most recent publication.

~~(3)~~ **(4)** A publication incorporated by reference is not effective until one hundred eighty (180) days after the date of publication. The executive board shall amend 410 IAC 15-1.7-1 and 410 IAC 15-2.7-1 to reflect the requirements in this subsection. This subsection expires July 1, 2021.

SECTION 5. IC 16-27-2.5-2, AS ADDED BY P.L.224-2017, SECTION 8, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: Sec. 2. (a) A home health agency must:

- (1) have a written drug testing policy that is distributed to all employees; and
- (2) require each employee to acknowledge receipt of the policy.

(b) A home health agency shall randomly test:

- (1) at least fifty percent (50%) of the home health agency's employees who:
  - (A) have direct contact with patients; and
  - (B) are not licensed by a board or commission under IC 25; at least annually; **or and**
- (2) when the home health agency has reasonable suspicion that an employee is engaged in the illegal use of a controlled substance.

(c) A home health agency shall either discharge or discipline with a minimum of a six (6) month suspension an employee who refuses to submit to a drug test.

SECTION 6. IC 25-23.6-2-6 IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: Sec. 6. The board shall meet at least ~~one (1) time each year:~~ **monthly.**

SECTION 7. IC 25-26-13-17, AS AMENDED BY P.L.202-2017, SECTION 4, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: Sec. 17. (a) The board shall establish classes of pharmacy permits as follows:

Category I. A retail permit for a pharmacy that provides pharmaceutical care to the general public by the dispensing of a drug or device.

Category II. An institutional permit for hospitals, clinics, health care facilities, sanitariums, nursing homes, or dispensaries that offer pharmaceutical care by dispensing a drug product to an inpatient under a drug order or to an outpatient of the institution under a prescription.



Category III. A permit for a pharmacy that provides closed door, central fill, mail order, or other processing operations that are not open to the general public but include:

- (A) traditional pharmacy functions; or
- (B) nontraditional pharmacy functions, such as infusion, nuclear pharmacy, or sterile compounding.

(b) Except for when registration as a remote dispensing facility (as defined in IC 25-26-13.5-3) is required under IC 25-26-13.5, the board may approve a remote or mobile location for Category I, II, or III permits **and any nonresident pharmacy registered with the board**. Pharmacy practice in a mobile or remote location may include, but is not limited to, telepharmacy, automated dispensing, or delivery of cognitive services.

(c) A hospital or hospital system holding a Category II permit may offer drugs or devices:

- (1) to:
  - (A) an employee, student, or volunteer of the hospital or hospital system;
  - (B) a retiree who is participating in a retirement, pension, or benefit program administered by the hospital or hospital system;
  - (C) an independent contractor who has an exclusive relationship with the hospital or hospital system;
  - (D) a member of the hospital's or hospital system's governing board; or
  - (E) a member of the hospital's or hospital system's medical staff; and

(2) to dependents of the individuals listed in subdivision (1);  
for their own use.

(d) Hospitals holding a Category II permit may operate remote locations within a reasonable distance of the licensed area, as determined by the board, after:

- (1) filing an application on a form prepared by the board;
- (2) having each location inspected by the board; and
- (3) obtaining approval from the board.

(e) Any applicable rule governing the practice of pharmacy in Indiana shall apply to all permits under this section.

(f) After June 30, 2012, a person with:

- (1) a Type I permit shall be treated as holding a Category I permit;
- (2) a Type II permit shall be treated as holding a Category II permit; and
- (3) a Type III, IV, V, or VI permit shall be treated as holding a



Category III permit.

The change in the name of the permit does not change the expiration date of the permit.

(g) After June 30, 2012, a reference in any rule or other document to:

- (1) a Type I permit shall be treated as a reference to a Category I permit;
- (2) a Type II permit shall be treated as a reference to a Category II permit; or
- (3) a Type III, IV, V, or VI permit shall be treated as a reference to a Category III permit.

**(h) A pharmacy holding a Category I permit may offer drugs or devices to the following:**

- (1) A long term care facility licensed under or subject to IC 16-28-2.**
- (2) A health facility licensed under IC 16-28.**
- (3) A housing with services establishment (as defined in IC 12-10-15-3) registered with the office of the secretary of family and social services.**

SECTION 8. IC 25-26-14-11, AS AMENDED BY P.L.212-2005, SECTION 45, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: Sec. 11. As used in this chapter, "wholesale distribution" means to distribute legend drugs to persons other than a consumer or patient. The term does not include:

- (1) a sale or transfer between a division, a subsidiary, a parent, an affiliated, or a related company under the common ownership and control of a corporate entity;
- (2) the purchase or acquisition by a hospital or other health care entity that is a member of a group purchasing organization of a drug for the hospital's or health care entity's own use from the group purchasing organization or from other hospitals or health care entities that are members of the organization;
- (3) the sale **or transfer** of a drug by a charitable organization described in Section 501(c)(3) of the Internal Revenue Code, to:
  - (A) a nonprofit affiliate of the organization; or**
  - (B) a nonprofit entity described in Section 501(c)(3) of the Internal Revenue Code that is not affiliated with the organization;**
 to the extent otherwise permitted by law;
- (4) the sale of a drug among hospitals or other health care entities that are under common control;
- (5) the sale of a drug for emergency medical reasons, including



transfers of legend drugs by a retail pharmacy to another retail pharmacy to alleviate a temporary shortage, if the gross dollar value of the transfers does not exceed five percent (5%) of the total legend drug sales revenue of either the transferor or transferee pharmacy during any twelve (12) consecutive month period;

(6) the sale of a drug or the dispensing of a drug pursuant to a prescription;

(7) the distribution of drug samples by manufacturers' representatives or distributors' representatives;

(8) the sale of blood and blood components intended for transfusion;

(9) the sale of a drug by a retail pharmacy to a practitioner (as defined in IC 25-26-13-2) for office use, if the gross dollar value of the transfers does not exceed five percent (5%) of the retail pharmacy's total legend drug sales during any twelve (12) consecutive months;

(10) the sale of a drug by a retail pharmacy that is ending its business and liquidating its inventory to another retail pharmacy;

(11) drug returns by a hospital, health care entity, or charitable institution conducted under 21 CFR 203.23;

(12) the sale of minimal quantities of drugs by retail pharmacies to licensed practitioners for office use;

(13) the distribution of prescription drugs by the original manufacturer of the finished form of the prescription drug or the distribution of the co-licensed products by a partner of the original manufacturer of the finished form of the prescription drug; or

(14) drug returns that meet criteria established by rules adopted by the board.

SECTION 9. IC 25-26-23-9 IS ADDED TO THE INDIANA CODE AS A **NEW SECTION** TO READ AS FOLLOWS [EFFECTIVE UPON PASSAGE]: **Sec. 9. (a) As used in this section, "unit" means a city, town, or county.**

**(b) A program to accept unused medication by a business or other entity that complies with applicable state and federal law is not subject to regulation by a unit.**

**(c) A unit may not do any of the following:**

**(1) Impose a tax, fee, assessment, or charge on a consumer, business, or other entity to pay for or support a program to accept unused medication in the unit's jurisdiction.**

**(2) Require a business or other entity to establish, pay for, or**



operate a program to accept unused medication in the unit's jurisdiction.

**(d) Nothing in this section prohibits a unit from using money in the unit's general fund to operate a program to accept unused medication.**

SECTION 10. IC 25-26-24-2.5 IS ADDED TO THE INDIANA CODE AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: **Sec. 2.5. As used in this chapter, "controlled substance" has the meaning set forth in IC 35-48-1-9. The term includes gabapentin.**

SECTION 11. IC 25-26-24-26 IS ADDED TO THE INDIANA CODE AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE APRIL 18, 2019 (RETROACTIVE)]: **Sec. 26. Any administrative rule adopted under IC 35-48-7-12.1 (before its repeal) is hereby considered to be adopted under section 22 of this chapter.**

SECTION 12. IC 27-1-37.4-8 IS ADDED TO THE INDIANA CODE AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2019]: **Sec. 8. (a) As used in this section, "step therapy protocol" means a protocol that specifies, as a condition of coverage under a health plan, the order in which certain prescription drugs must be used to treat a covered individual's condition.**

**(b) A health plan that denies prior authorization for a prescription drug described in subdivision (1) or (2) shall provide, in the notice of denial, an alternative list of prescription drugs or alternative treatments as follows:**

**(1) If:**

**(A) the prescription drug is not included in the health plan's formulary; and**

**(B) there is at least one (1) alternative prescription drug in the same therapeutic classification (as defined in IC 12-15-35-17.5);**

**the alternative list must specify the alternative prescription drugs described in clause (B) that are covered by the health plan.**

**(2) If the prescription drug is prescribed to treat a condition for which coverage under the health plan requires use of a step therapy protocol, the alternative list must specify the alternative prescription drugs or alternative treatments that are required by the step therapy protocol.**

SECTION 13. [EFFECTIVE UPON PASSAGE] **(a) The Indiana**



**board of veterinary medical examiners shall study the regulation of veterinary technicians and submit a report to the legislative council in an electronic format under IC 5-14-6 before November 1, 2019.**

**(b) This SECTION expires January 1, 2020.**

**SECTION 14. An emergency is declared for this act.**





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Speaker of the House of Representatives

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President of the Senate

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President Pro Tempore

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Governor of the State of Indiana

Date: \_\_\_\_\_ Time: \_\_\_\_\_

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