SENATE BILL No. 627

DIGEST OF INTRODUCED BILL

Citations Affected: IC 6-2.5-8-7; IC 24-4; IC 34-30-2-152.3; IC 35-48.

Synopsis: Sale of low THC hemp extract products. Repeals laws concerning: (1) the distribution of low tetrahydrocannabinol (THC) hemp extract; and (2) low THC hemp extract sales. Provides that only a pharmacy or National Precursor Log Exchange (NPLEx) retailer may sell low THC hemp extract. Specifies that a person who is denied the sale of a nonprescription product containing low THC hemp extract is not prohibited from obtaining low THC hemp extract pursuant to a prescription. Provides that a pharmacist or pharmacy technician may determine that the purchaser has a relationship on record with the pharmacy, in compliance with rules adopted by the board. Allows a pharmacist to deny the sale of low THC hemp extract on the basis of the pharmacist's professional judgment, and provides the pharmacist with civil immunity for making such a denial. Provides that a purchaser who has a relationship on record with the pharmacy may purchase low THC hemp extract. Allows the pharmacist to provide certain low THC hemp extract products to a purchaser who does not have a relationship on record with the pharmacy or for whom the pharmacist has made a professional judgment that there is not a medical or pharmaceutical need. Requires the Indiana scheduled prescription electronic collection and tracking (INSPECT) program to track low THC hemp extract dispensed pursuant to a prescription. Makes conforming changes.

Effective: July 1, 2019.

Brown L

January 15, 2019, read first time and referred to Committee on Corrections and Criminal Law.



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.

SENATE BILL No. 627

A BILL FOR AN ACT to amend the Indiana Code concerning criminal law and procedure.

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

1	SECTION 1. IC 6-2.5-8-7, AS AMENDED BY P.L.153-2018,
2	SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
3	JULY 1, 2019]: Sec. 7. (a) The department may, for good cause, revoke
4	a certificate issued under section 1, 3, or 4 of this chapter. However,
5	the department must give the certificate holder at least five (5) days
6	notice before it revokes the certificate under this subsection. Good
7	cause for revocation may include the following:
8	(1) Failure to:
9	(A) file a return required under this chapter or for any tax
10	collected for the state in trust; or
11	(B) remit any tax collected for the state in trust.
12	(2) Being charged with a violation of any provision under IC 35.
13	(3) Being subject to a court order under IC 7.1-2-6-7,
14	IC 32-30-6-8, IC 32-30-7, or IC 32-30-8.
15	(4) Being charged with a violation of IC 23-15-12.
16	The department may revoke a certificate before a criminal adjudication
17	or without a criminal charge being filed. If the department gives notice



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of an intent to revoke based on an alleged violation of subdivision (2), the department shall hold a public hearing to determine whether good cause exists. If the department finds in a public hearing by a preponderance of the evidence that a person has committed a violation described in subdivision (2), the department shall proceed in accordance with subsection (i) (if the violation resulted in a criminal conviction) or subsection (j) (if the violation resulted in a judgment for an infraction).
(b) The department shall revoke a certificate issued under section
1, 3, or 4 of this chapter if, for a period of three (3) years, the certificate
holder fails to:
(1) file the returns required by IC 6-2.5-6-1; or
(2) report the collection of any state gross retail or use tax on the

(2) report the collection of any state gross retail or use tax on the returns filed under IC 6-2.5-6-1.

However, the department must give the certificate holder at least five (5) days notice before it revokes the certificate.

- (c) The department may, for good cause, revoke a certificate issued under section 1 of this chapter after at least five (5) days notice to the certificate holder if:
 - (1) the certificate holder is subject to an innkeeper's tax under IC 6-9; and
 - (2) a board, bureau, or commission established under IC 6-9 files a written statement with the department.
 - (d) The statement filed under subsection (c) must state that:
 - (1) information obtained by the board, bureau, or commission under IC 6-8.1-7-1 indicates that the certificate holder has not complied with IC 6-9; and
 - (2) the board, bureau, or commission has determined that significant harm will result to the county from the certificate holder's failure to comply with IC 6-9.
- (e) The department shall revoke or suspend a certificate issued under section 1 of this chapter after at least five (5) days notice to the certificate holder if:
 - (1) the certificate holder owes taxes, penalties, fines, interest, or costs due under IC 6-1.1 that remain unpaid at least sixty (60) days after the due date under IC 6-1.1; and
 - (2) the treasurer of the county to which the taxes are due requests the department to revoke or suspend the certificate.
- (f) The department shall reinstate a certificate suspended under subsection (e) if the taxes and any penalties due under IC 6-1.1 are paid or the county treasurer requests the department to reinstate the certificate because an agreement for the payment of taxes and any



1	penalties due under IC 6-1.1 has been reached to the satisfaction of the
2	county treasurer.
3	(g) The department shall revoke a certificate issued under section
4	1 of this chapter after at least five (5) days notice to the certificate
5	holder if the department finds in a public hearing by a preponderance
6	of the evidence that the certificate holder has violated IC 35-45-5-3,
7	IC 35-45-5-3.5, or IC 35-45-5-4.
8	(h) If a person makes a payment for the certificate under section 1
9	or 3 of this chapter with a check, credit card, debit card, or electronic
10	funds transfer, and the department is unable to obtain payment of the
11	check, credit card, debit card, or electronic funds transfer for its full
12	face amount when the check, credit card, debit card, or electronic funds
13	transfer is presented for payment through normal banking channels, the
14	department shall notify the person by mail that the check, credit card,
15	debit card, or electronic funds transfer was not honored and that the
16	person has five (5) days after the notice is mailed to pay the fee in cash,
17	by certified check, or other guaranteed payment. If the person fails to
18	make the payment within the five (5) day period, the department shall
19	revoke the certificate.
20	(i) If the department finds in a public hearing by a preponderance of
21	the evidence that a person has a conviction for a violation of
22	IC 35-48-4-10.5 and the conviction involved the sale of or the offer to
23	sell, in the normal course of business, a synthetic drug or a synthetic
24	drug lookalike substance by a retail merchant in a place of business for
25	which the retail merchant has been issued a registered retail merchant
26	certificate under section 1 of this chapter, the department:
27	(1) shall suspend the registered retail merchant certificate for the
28	place of business for one (1) year; and
29	(2) may not issue another retail merchant certificate under section
30	1 of this chapter for one (1) year to any person:
31	(A) that:
32	(i) applied for; or
33	(ii) made a retail transaction under;
34	the retail merchant certificate suspended under subdivision
35	(1); or
36	(B) that:
37	(i) owned or co-owned, directly or indirectly; or
38	(ii) was an officer, a director, a manager, or a partner of;
39	the retail merchant that was issued the retail merchant
40	certificate suspended under subdivision (1).
41	(j) If the department finds in a public hearing by a preponderance of

(j) If the department finds in a public hearing by a preponderance of

the evidence that a person has a judgment for a violation of



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1	IC 35-48-4-10.5 as an infraction and the violation involved the sale of
2	or the offer to sell, in the normal course of business, a synthetic drug
3	or a synthetic drug lookalike substance by a retail merchant in a place
4	of business for which the retail merchant has been issued a registered
5	retail merchant certificate under section 1 of this chapter, the
6	department:
7	(1) may suspend the registered retail merchant certificate for the
8	place of business for six (6) months; and
9	(2) may withhold issuance of another retail merchant certificate
10	under section 1 of this chapter for six (6) months to any person:
11	(A) that:
12	(i) applied for; or
13	(ii) made a retail transaction under;
14	the retail merchant certificate suspended under subdivision
15	(1); or
16	(B) that:
17	(i) owned or co-owned, directly or indirectly; or
18	(ii) was an officer, a director, a manager, or a partner of;
19	the retail merchant that was issued the retail merchant
20	certificate suspended under subdivision (1).
21	(k) If the department finds in a public hearing by a preponderance
22	of the evidence that a person has a conviction for a violation of
23 24 25	IC 35-48-4-10(d)(3) and the conviction involved an offense committed
24	by a retail merchant in a place of business for which the retail merchant
25	has been issued a registered retail merchant certificate under section 1
26	of this chapter, the department:
27	(1) shall suspend the registered retail merchant certificate for the
28	place of business for one (1) year; and
29	(2) may not issue another retail merchant certificate under section
30	1 of this chapter for one (1) year to any person:
31	(A) that:
32	(i) applied for; or
33	(ii) made a retail transaction under;
34	the retail merchant certificate suspended under subdivision
35	(1); or
36	(B) that:
37	(i) owned or co-owned, directly or indirectly; or
38	(ii) was an officer, a director, a manager, or a partner of;
39	the retail merchant that was issued the retail merchant
40	certificate suspended under subdivision (1).
41	SECTION 2. IC 24-4-21 IS REPEALED [EFFECTIVE JULY 1,
42	2019]. (Distribution of Low THC Hemp Extract).



1	SECTION 3. IC 24-4-22 IS REPEALED [EFFECTIVE JULY 1
2	2019]. (Low THC Hemp Extract Sales).
3	SECTION 4. IC 34-30-2-152.3, AS AMENDED BY P.L.5-2016
4	SECTION 4, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
5	JULY 1, 2019]: Sec. 152.3. (a) IC 35-48-4-14.7(d) and
6	IC 35-48-4-14.7(k) (Concerning a pharmacy or NPLEx retailer tha
7	discloses information concerning the sale of a product containing
8	ephedrine, or pseudoephedrine, or low THC hemp extract).
9	(b) IC 35-48-4-14.7(d)(3) (Concerning a pharmacist's professiona
10	judgment not to sell ephedrine, or pseudoephedrine, or low THC
1	hemp extract to an individual).
12	SECTION 5. IC 35-48-4-10, AS AMENDED BY P.L.153-2018
13	SECTION 25, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
14	JULY 1, 2019]: Sec. 10. (a) A person who:
15	(1) knowingly or intentionally:
16	(A) manufactures;
17	(B) finances the manufacture of;
18	(C) delivers; or
19	(D) finances the delivery of;
20	marijuana, hash oil, hashish, or salvia, pure or adulterated; or
21	(2) possesses, with intent to:
22	(A) manufacture;
23	(B) finance the manufacture of;
24 25	(C) deliver; or
25	(D) finance the delivery of;
26	marijuana, hash oil, hashish, or salvia, pure or adulterated;
27	commits dealing in marijuana, hash oil, hashish, or salvia, a Class A
28	misdemeanor, except as provided in subsections (b) through (d).
29	(b) A person may be convicted of an offense under subsection (a)(2)
30	only if:
31	(1) there is evidence in addition to the weight of the drug that the
32	person intended to manufacture, finance the manufacture of
33	deliver, or finance the delivery of the drug; or
34	(2) the amount of the drug involved is at least:
35	(A) ten (10) pounds, if the drug is marijuana; or
36	(B) three hundred (300) grams, if the drug is hash oil, hashish
37	or salvia.
38	(c) The offense is a Level 6 felony if:
39	(1) the person has a prior conviction for a drug offense and the
10	amount of the drug involved is:
11	(A) less than thirty (30) grams of marijuana; or
12	(B) less than five (5) grams of hash oil, hashish, or salvia; or



1	(2) the amount of the drug involved is:
2	(A) at least thirty (30) grams but less than ten (10) pounds of
3	marijuana; or
4	(B) at least five (5) grams but less than three hundred (300)
5	grams of hash oil, hashish, or salvia.
6	(d) The offense is a Level 5 felony if:
7	(1) the person has a prior conviction for a drug dealing offense
8	and the amount of the drug involved is:
9	(A) at least thirty (30) grams but less than ten (10) pounds of
0	marijuana; or
1	(B) at least five (5) grams but less than three hundred (300)
2	grams of hash oil, hashish, or salvia;
3	(2) the:
4	(A) amount of the drug involved is:
5	(i) at least ten (10) pounds of marijuana; or
6	(ii) at least three hundred (300) grams of hash oil, hashish
7	or salvia; or
8	(B) offense involved a sale to a minor; or
9	(3) the:
0.	(A) person is a pharmacy or NPLEx retailer (as defined in
21	section 14.7(b) of this chapter;
.2	(B) marijuana, hash oil, hashish, or salvia is packaged in a
23	manner that appears to be low THC hemp extract; and
12 13 14 15 16	(C) person knew or reasonably should have known that the
25	product was marijuana, hash oil, hashish, or salvia.
	SECTION 6. IC 35-48-4-14.7, AS AMENDED BY P.L.252-2017
27	SECTION 25, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
8	JULY 1, 2019]: Sec. 14.7. (a) This section does not apply to the
9	following:
0	(1) Ephedrine, or pseudoephedrine, or low THC hemp extract
1	dispensed pursuant to a prescription. Nothing in this section
2	prohibits a person who is denied the sale of a nonprescription
3	product containing pseudoephedrine, or ephedrine, or low THC
4	hemp extract from obtaining pseudoephedrine, or ephedrine, or
5	low THC hemp extract pursuant to a prescription.
6	(2) The sale of a drug containing ephedrine, or pseudoephedrine
7	or low THC hemp extract to a licensed health care provider
8	pharmacist, retail distributor, wholesaler, manufacturer, or ar
9	agent of any of these persons if the sale occurs in the regular
0	course of lawful business activities. However, a retail distributor
-1	wholesaler, or manufacturer is required to report a suspicious
.2	order to the state police department in accordance with subsection



1	(g).
2	(3) The sale of a drug containing ephedrine, or pseudoephedrine,
3	or low THC hemp extract by a person who does not sell
4	exclusively to walk-in customers for the personal use of the
5	walk-in customers. However, if the person described in this
6	subdivision is a retail distributor, wholesaler, or manufacturer, the
7	person is required to report a suspicious order to the state police
8	department in accordance with subsection (g).
9	(b) The following definitions apply throughout this section:
10	(1) "Constant video monitoring" means the surveillance by an
11	automated camera that:
12	(A) records at least one (1) photograph or digital image every
13	ten (10) seconds;
14	(B) retains a photograph or digital image for at least
15	seventy-two (72) hours;
16	(C) has sufficient resolution and magnification to permit the
17	identification of a person in the area under surveillance; and
18	(D) stores a recorded photograph or digital image at a location
19	that is immediately accessible to a law enforcement officer.
20	(2) "Convenience package" means a package that contains a drug
21	having as an active ingredient not more than sixty (60) milligrams
22	of ephedrine or pseudoephedrine, or both.
23	(3) "Ephedrine" means pure or adulterated ephedrine.
24	(4) "Low THC hemp extract" has the meaning set forth in
25	IC 35-48-1-17.5.
26	(4) (5) "Pharmacy or NPLEx retailer" means:
27	(A) a pharmacy, as defined in IC 25-26-13-2;
28	(B) a retailer containing a pharmacy, as defined in
29	IC 25-26-13-2; or
30	(C) a retailer that electronically submits the required
31	information to the National Precursor Log Exchange (NPLEx).
32	(5) (6) "Pseudoephedrine" means pure or adulterated
33	pseudoephedrine.
34	(6) (7) "Retailer" means a grocery store, general merchandise
35	store, or other similar establishment. The term does not include a
36	pharmacy or NPLEx retailer.
37	(7) (8) "Suspicious order" means a sale or transfer of a drug
38	containing ephedrine or pseudoephedrine if the sale or transfer:
39	(A) is a sale or transfer that the retail distributor, wholesaler,
40	or manufacturer is required to report to the United States Drug
41	Enforcement Administration;
42	(B) appears suspicious to the retail distributor, wholesaler, or



1	manufacturer in light of the recommendations contained in
2	Appendix A of the report to the United States attorney general
3	by the suspicious orders task force under the federal
4	Comprehensive Methamphetamine Control Act of 1996; or
5	(C) is for cash or a money order in a total amount of at least
6	two hundred dollars (\$200).
7	(8) (9) "Unusual theft" means the theft or unexplained
8	disappearance from a particular pharmacy or NPLEx retailer of
9	drugs containing ten (10) grams or more of ephedrine,
10	pseudoephedrine, or both in a twenty-four (24) hour period.
11	(c) A drug containing ephedrine, or pseudoephedrine, or low THC
12	hemp extract may be sold only by a pharmacy or NPLEx retailer.
13	(d) A pharmacy or NPLEx retailer may sell a drug that contains the
14	active ingredient of ephedrine, pseudoephedrine, low THC hemp
15	extract, or both any combination of the ingredients only if the
16	pharmacy or NPLEx retailer complies with the following conditions:
17	(1) The pharmacy or NPLEx retailer does not sell the drug to a
18	person less than eighteen (18) years of age.
19	(2) The pharmacy or NPLEx retailer does not sell drugs
20	containing more than:
21	(A) three and six-tenths (3.6) grams of ephedrine or
22	pseudoephedrine, or both, to one (1) individual on one (1) day;
23	(B) seven and two-tenths (7.2) grams of ephedrine or
24	pseudoephedrine, or both, to one (1) individual in a thirty (30)
25	day period; or
26	(C) sixty-one and two-tenths (61.2) grams of ephedrine or
27	pseudoephedrine, or both, to one (1) individual in a three
28	hundred sixty-five (365) day period.
29	(3) Except as provided in subsection (f), before the sale occurs the
30	pharmacist or the pharmacy technician (as defined by
31	IC 25-26-19-2) has determined that the purchaser has a
32	relationship on record with the pharmacy, in compliance with
33	rules adopted by the board under IC 25-26-13-4. If it has been
34	determined that the purchaser does not have a relationship on
35	record with the pharmacy, the pharmacist shall make a
36	professional determination as to whether there is a legitimate
37	medical or pharmaceutical need for ephedrine, or
38	pseudoephedrine, or low THC hemp extract before selling
39	ephedrine, or pseudoephedrine, or low THC hemp extract to an
40	individual. The pharmacist's professional determination must
41	comply with the rules adopted under IC 25-26-13-4 and may
42	include the following:



1	(A) Prior medication filling history of the individual.
2	(B) Consulting with the individual.
3	(C) Other tools that provide professional reassurance to the
4	pharmacist that a legitimate medical or pharmaceutical need
5	for ephedrine, or pseudoephedrine, or low THC hemp extract
6	exists.
7	A pharmacist who in good faith does not sell ephedrine, or
8	pseudoephedrine, or low THC hemp extract to an individual
9	under this subdivision is immune from civil liability unless the
10	refusal to sell constitutes gross negligence or intentional, wanton,
11	or willful misconduct.
12	(4) The pharmacy or NPLEx retailer requires:
13	(A) the purchaser to produce a valid government issued photo
14	identification card showing the date of birth of the person;
15	(B) the purchaser to sign a written or electronic log attesting
16	to the validity of the information; and
17	(C) the clerk who is conducting the transaction to initial or
18	electronically record the clerk's identification on the log.
19	Records from the completion of a log must be retained for at least
20	two (2) years. A law enforcement officer has the right to inspect
21	and copy a log or the records from the completion of a log in
22	accordance with state and federal law. A pharmacy or NPLEx
23	retailer may not sell or release a log or the records from the
24	completion of a log for a commercial purpose. The Indiana
25	criminal justice institute may obtain information concerning a log
26	or the records from the completion of a log from a law
27	enforcement officer if the information may not be used to identify
28	a specific individual and is used only for statistical purposes. A
29	pharmacy or NPLEx retailer that in good faith releases
30	information maintained under this subsection is immune from
31	civil liability unless the release constitutes gross negligence or
32	intentional, wanton, or willful misconduct.
33	(5) The pharmacy or NPLEx retailer maintains a record of
34	information for each sale of a nonprescription product containing
35	pseudoephedrine, or ephedrine, or low THC hemp extract.
36	Required information includes:
37	(A) the name and address of each purchaser;
38	(B) the type of identification presented;
39	(C) the governmental entity that issued the identification;
40	(D) the identification number; and
41	(E) the ephedrine, or pseudoephedrine, or low THC hemp
42	extract product purchased, including the number of grams the
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1	product contains and the date and time of the transaction.
2	(6) A pharmacy or NPLEx retailer shall, except as provided in
3	subdivision (7), before completing a sale of an over-the-counter
4	product containing pseudoephedrine, or ephedrine, or low THC
5	hemp extract, electronically submit the required information to
6	the National Precursor Log Exchange (NPLEx), if the NPLEx
7	system is available to pharmacies or NPLEx retailers in the state
8	without a charge for accessing the system. The pharmacy or
9	NPLEx retailer may not complete the sale if the system generates
10	a stop sale alert, including a stop sale alert for a person convicted
11	of a drug related felony reported under IC 33-24-6-3.
12	(7) If a pharmacy or NPLEx retailer selling an over-the-counter
13	product containing ephedrine, or pseudoephedrine, or low THC
14	hemp extract experiences mechanical or electronic failure of the
15	electronic sales tracking system and is unable to comply with the
16	electronic sales tracking requirement, the pharmacy or NPLEx
17	retailer shall maintain a written log or an alternative electronic
18	record keeping mechanism until the pharmacy or NPLEx retailer
19	is able to comply with the electronic sales tracking requirement.
20	(8) The pharmacy or NPLEx retailer stores the drug behind a
21	counter in an area inaccessible to a customer or in a locked
22	display case that makes the drug unavailable to a customer
23	without the assistance of an employee.
24	(e) A person may not purchase drugs containing more than:
25	(1) three and six-tenths (3.6) grams of ephedrine or
26	pseudoephedrine, or both, on one (1) day;
27	(2) seven and two-tenths (7.2) grams of ephedrine or
28	pseudoephedrine, or both, in a thirty (30) day period; or
29	(3) sixty-one and two-tenths (61.2) grams of ephedrine or
30	pseudoephedrine, or both, in a three hundred sixty-five (365) day
31	period.
32	These limits apply to the total amount of base ephedrine and
33	pseudoephedrine contained in the products and not to the overall
34	weight of the products.
35	(f) If a purchaser does not have a relationship on record with the
36	pharmacy, as determined by rules adopted by the board under
37	IC 25-26-13-4, or the pharmacist has made a professional
38	determination that there is not a legitimate medical or pharmaceutical
39	need for ephedrine, or pseudoephedrine, or low THC hemp extract
40	under subsection (d), the purchaser may, at the pharmacist's discretion,
41	purchase only the following:
42	(1) A product that has been determined under section 14.3 of this



1	chapter to be an extraction resistant or a conversion resistant form
2	of ephedrine or pseudoephedrine.
3	(2) A product that contains not more than:
4	(A) a total of seven hundred twenty (720) milligrams of
5	ephedrine or pseudoephedrine per package; and
6	(B) thirty (30) milligrams of ephedrine or pseudoephedrine per
7	tablet.
8	(3) A product that contains not more than:
9	(A) sixty (60) milligrams of cannabidiol per milliliter; and
10	(B) thirty (30) milliliters per container.
11	The pharmacist may not sell more than one (1) package of ephedrine
12	or pseudoephedrine or one (1) container of low THC hemp extract
13	to a purchaser under this subdivision subsection per day. However, if
14	the pharmacist believes that the ephedrine or pseudoephedrine
15	purchase will be used to manufacture methamphetamine, the
16	pharmacist may refuse to sell ephedrine or pseudoephedrine to the
17	purchaser.
18	(g) A retail distributor, wholesaler, or manufacturer shall report a
19	suspicious order to the state police department in writing.
20	(h) Not later than three (3) days after the discovery of an unusual
21	theft at a particular retail store, the pharmacy or NPLEx retailer shall
22	report the unusual theft to the state police department in writing. If
23	three (3) unusual thefts occur in a thirty (30) day period at a particular
24	pharmacy or NPLEx retailer, the pharmacy or NPLEx retailer shall, for
25	at least one hundred eighty (180) days after the date of the last unusual
26	theft, locate all drugs containing ephedrine, or pseudoephedrine, or low
27	THC hemp extract at that particular pharmacy or NPLEx retailer
28	behind a counter in an area inaccessible to a customer or in a locked
29	display case that makes the drug unavailable to customers without the
30	assistance of an employee.
31	(i) A unit (as defined in IC 36-1-2-23) may not adopt an ordinance
32	after February 1, 2005, that is more stringent than this section.
33	(j) A person who knowingly or intentionally violates this section
34	commits a Class C misdemeanor. However, the offense is a Class A
35	misdemeanor if the person has a prior unrelated conviction under this
36	section.
37	(k) A pharmacy or NPLEx retailer that uses the electronic sales
38	tracking system in accordance with this section is immune from civil
39	liability for any act or omission committed in carrying out the duties
40	required by this section, unless the act or omission was due to
41	recklessness or deliberate or wanton misconduct. A pharmacy or

NPLEx retailer is immune from liability to a third party unless the



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1	pharmacy or NPLEx retailer has violated a provision of this section and
2	the third party brings an action based on the pharmacy's or NPLEx
3	retailer's violation of this section.
4	(1) The following requirements apply to the NPLEx:
5	(1) Information contained in the NPLEx may be shared only with
6	law enforcement officials.
7	(2) A law enforcement official may access Indiana transaction
8	information maintained in the NPLEx for investigative purposes.
9	(3) NADDI may not modify sales transaction data that is shared
0	with law enforcement officials.
1	(4) At least one (1) time per day, Indiana data contained in the
2	NPLEx for the previous calendar day shall be forwarded to the
.3	state police department.
4	(m) A person or corporate entity may not mandate a protocol or
.5	procedure that interferes with the pharmacist's ability to exercise the
.6	pharmacist's independent professional judgment under this section,
.7	including whether to deny the sale of ephedrine, or pseudoephedrine,
8	or low THC hemp extract under subsection (f).
9	SECTION 7. IC 35-48-7-8.1, AS AMENDED BY P.L.194-2018,
20	SECTION 2, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
21	JULY 1, 2019]: Sec. 8.1. (a) The board shall provide for an ephedrine,
22	pseudoephedrine, low THC hemp extract, and controlled substance
23	prescription monitoring program that includes the following
24	components:
25	(1) Each time ephedrine, pseudoephedrine, low THC hemp
26	extract, or a controlled substance designated by the board under
27	IC 35-48-2-5 through IC 35-48-2-10 is dispensed, the dispenser
28	shall transmit to the INSPECT program the following
29	information:
80	(A) The ephedrine, pseudoephedrine, low THC hemp extract,
31	or controlled substance recipient's name.
32	(B) The ephedrine, pseudoephedrine, low THC hemp extract,
33	or controlled substance recipient's or the recipient
34	representative's identification number or the identification
35	number or phrase designated by the INSPECT program.
86	(C) The ephedrine, pseudoephedrine, low THC hemp extract,
37	or controlled substance recipient's date of birth.
88	(D) The national drug code number of the ephedrine,
39	pseudoephedrine, low THC hemp extract, or controlled
10	substance dispensed.
1	(E) The date the ephedrine, pseudoephedrine, low THC hemp
12	extract or controlled substance is dispensed



1	(F) The quantity of the ephedrine, pseudoephedrine, low THC
2	hemp extract, or controlled substance dispensed.
3	(G) The number of days of supply dispensed.
4	(H) The dispenser's United States Drug Enforcement Agency
5	registration number.
6	(I) The prescriber's United States Drug Enforcement Agency
7	registration number.
8	(J) An indication as to whether the prescription was
9	transmitted to the pharmacist orally or in writing.
10	(K) Other data required by the board.
11	(2) The information required to be transmitted under this section
12	must be transmitted not more than twenty-four (24) hours after the
13	date on which ephedrine, pseudoephedrine, low THC hemp
14	extract, or a controlled substance is dispensed. However, if the
15	dispenser's pharmacy is closed the day following the dispensing
16	the information must be transmitted by the end of the next
17	business day.
18	(3) A dispenser shall transmit the information required under this
19	section by:
20	(A) uploading to the INSPECT web site; or
21	(B) another electronic method that meets specifications
22	prescribed by the board.
23	(4) The board may require that prescriptions for ephedrine
22 23 24	pseudoephedrine, low THC hemp extract, or controlled
25	substances be written on a one (1) part form that cannot be
26	duplicated. However, the board may not apply such a requirement
27	to prescriptions filled at a pharmacy with a Category II permit (as
28	described in IC 25-26-13-17) and operated by a hospital licensed
29	under IC 16-21, or prescriptions ordered for and dispensed to
30	bona fide enrolled patients in facilities licensed under IC 16-28.
31	The board may not require multiple copy prescription forms for
32	any prescriptions written. The board may not require different
33	prescription forms for any individual drug or group of drugs.
34	Prescription forms required under this subdivision must be
35	approved by the Indiana board of pharmacy established by
36	IC 25-26-13-3.
37	(5) The costs of the program.
38	(6) As part of the information to be completed in the data base
39	and if available, an entry where a dispenser indicates that a
40	patient is participating in a pain management contract with a
41	designated practitioner.
	~ x

(b) The board shall consider the recommendations of the committee



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1	concerning the INSPECT program.
2	(c) This subsection applies only to a retail pharmacy. A pharmacist,
3	pharmacy technician, or person authorized by a pharmacist to dispense
4	ephedrine, pseudoephedrine, low THC hemp extract, or a controlled
5	substance may not dispense ephedrine, pseudoephedrine, low THC
6	hemp extract, or a controlled substance to a person who is not
7	personally known to the pharmacist, pharmacy technician, or person
8	authorized by a pharmacist to dispense a controlled substance unless
9	the person taking possession of the ephedrine, pseudoephedrine, low
10	THC hemp extract, or controlled substance provides documented
11	proof of the person's identification to the pharmacist, pharmacy
12	technician, or person authorized by a pharmacist to dispense ephedrine,
13	pseudoephedrine, low THC hemp extract, or a controlled substance.
14	SECTION 8. IC 35-48-7-10.1, AS AMENDED BY P.L.194-2018,
15	SECTION 3, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
16	JULY 1, 2019]: Sec. 10.1. (a) The INSPECT program must do the
17	following:
18	(1) Create a data base for information required to be transmitted
19	under section 8.1 of this chapter in the form required under rules
20	adopted by the board, including search capability for the
21	following:
22	(A) An ephedrine, pseudoephedrine, low THC hemp extract,
22 23	· ·
	(A) An ephedrine, pseudoephedrine, low THC hemp extract,
23	(A) An ephedrine, pseudoephedrine, low THC hemp extract , or a controlled substance recipient's name.
23 24	(A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name.(B) An ephedrine, pseudoephedrine, low THC hemp extract,
23 24 25	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient
23 24 25 26 27 28	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number.
23 24 25 26 27 28 29	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract,
23 24 25 26 27 28 29 30	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth.
23 24 25 26 27 28 29 30 31	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed.
23 24 25 26 27 28 29 30 31 32	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp
23 24 25 26 27 28 29 30 31 32 33	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed.
23 24 25 26 27 28 29 30 31 32	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp
23 24 25 26 27 28 29 30 31 32 33 34 35	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance are dispensed. (F) The quantities of ephedrine, pseudoephedrine, low THC hemp extract, or controlled substance dispensed.
23 24 25 26 27 28 29 30 31 32 33 34 35 36	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance are dispensed. (F) The quantities of ephedrine, pseudoephedrine, low THC
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance are dispensed. (F) The quantities of ephedrine, pseudoephedrine, low THC hemp extract, or controlled substance dispensed.
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance are dispensed. (F) The quantities of ephedrine, pseudoephedrine, low THC hemp extract, or controlled substance dispensed. (G) The number of days of supply dispensed. (H) A dispenser's United States Drug Enforcement Agency registration number.
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance are dispensed. (F) The quantities of ephedrine, pseudoephedrine, low THC hemp extract, or controlled substance dispensed. (G) The number of days of supply dispensed. (H) A dispenser's United States Drug Enforcement Agency registration number. (I) A prescriber's United States Drug Enforcement Agency
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance are dispensed. (F) The quantities of ephedrine, pseudoephedrine, low THC hemp extract, or controlled substance dispensed. (G) The number of days of supply dispensed. (H) A dispenser's United States Drug Enforcement Agency registration number. (I) A prescriber's United States Drug Enforcement Agency registration number.
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39	 (A) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's name. (B) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's or recipient representative's identification number. (C) An ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance recipient's date of birth. (D) The national drug code number of ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance dispensed. (E) The dates ephedrine, pseudoephedrine, low THC hemp extract, or a controlled substance are dispensed. (F) The quantities of ephedrine, pseudoephedrine, low THC hemp extract, or controlled substance dispensed. (G) The number of days of supply dispensed. (H) A dispenser's United States Drug Enforcement Agency registration number. (I) A prescriber's United States Drug Enforcement Agency



1	(K) An ephedrine, pseudoephedrine, low 1 HC nemp extract,
2 3	or a controlled substance recipient's method of payment for the
	ephedrine, pseudoephedrine, low THC hemp extract, or
4	controlled substance dispensed.
5	To the extent considered appropriate by the board, the data base
6	must be interoperable with other similar registries operated by
7	federal and state governments.
8	(2) Provide the board with continuing twenty-four (24) hour a day
9	online access to the data base.
10	(3) Secure the information collected and the data base maintained
11	against access by unauthorized persons.
12	(b) The board may not execute a contract with a vendor designated
13	by the board to perform any function associated with the administration
14	of the INSPECT program, unless the contract has been approved by the
15	committee.
16	(c) The INSPECT program may gather prescription data from the
17	Medicaid retrospective drug utilization review (DUR) program
18	established under IC 12-15-35.
19	(d) The board may accept and designate grants, public and private
20	financial assistance, and licensure fees to provide funding for the
21	INSPECT program.
22	SECTION 9. IC 35-48-7-11.1, AS AMENDED BY P.L.194-2018,
23	SECTION 4, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
24	JULY 1, 2019]: Sec. 11.1. (a) Information received by the INSPECT
25	program under section 8.1 of this chapter is confidential.
26	(b) The board shall carry out a program to protect the confidentiality
27	of the information described in subsection (a). The board may disclose
28	the information to another person only under subsection (c) , (d) , or (g) .
29	(c) The board may disclose confidential information described in
30	subsection (a) to any person who is authorized to engage in receiving,
31	processing, or storing the information.
32	(d) Except as provided in subsections (e) and (f), the board may
33	release confidential information described in subsection (a) to the
34	following persons:
35	(1) A member of the board or another governing body that
36	licenses practitioners and is engaged in an investigation, an
37	adjudication, or a prosecution of a violation under any state or
38	federal law that involves ephedrine, pseudoephedrine, low THC
39	hemp extract, or a controlled substance.
40	(2) An investigator for the consumer protection division of the
41	office of the attorney general, a prosecuting attorney, the attorney
42	general, a deputy attorney general, or an investigator from the



1	office of the attorney general, who is engaged in:
2	(A) an investigation;
3	(B) an adjudication; or
4	(C) a prosecution;
5	of a violation under any state or federal law that involves
6	ephedrine, pseudoephedrine, low THC hemp extract, or a
7	controlled substance.
8	(3) A law enforcement officer who is an employee of:
9	(A) a local, state, or federal law enforcement agency; or
10	(B) an entity that regulates ephedrine, pseudoephedrine, low
11	THC hemp extract, or controlled substances or enforces
12	ephedrine, pseudoephedrine, low THC hemp extract, or
13	controlled substances rules or laws in another state;
14	that is certified to receive ephedrine, pseudoephedrine, low THC
15	hemp extract, or controlled substance prescription drug
16	information from the INSPECT program.
17	(4) A practitioner or practitioner's agent certified to receive
18	information from the INSPECT program.
19	(5) An ephedrine, pseudoephedrine, low THC hemp extract, or
20	controlled substance monitoring program in another state with
21	which Indiana has established an interoperability agreement.
	(6) The state toxicologist.
22 23 24 25	(7) A certified representative of the Medicaid retrospective and
24	prospective drug utilization review program.
25	(8) A substance abuse assistance program for a licensed health
26	care provider who:
27 28	(A) has prescriptive authority under IC 25; and
28	(B) is participating in the assistance program.
29	(9) An individual who holds a valid temporary medical permit
30	issued under IC 25-22.5-5-4 or a temporary fellowship permit
31	issued under IC 25-22.5-5-4.6.
32	(10) A county coroner conducting a medical investigation of the
33	cause of death.
34	(11) The management performance hub established by Indiana
35	Executive Order 14-06 and continued by Executive Order 17-09.
36	(12) The state epidemiologist under the state department of
37	health.
38	(e) Information provided to a person under:
39	(1) subsection (d)(3) is limited to information:
40	(A) concerning an individual or proceeding involving the
41	unlawful diversion or misuse of a schedule II, III, IV, or V
12	controlled substance: and



(B) that will assist in an investigation or proceeding;
(2) subsection (d)(4) may be released only for the purpose of:
(A) providing medical or pharmaceutical treatment; or
(B) evaluating the need for providing medical or
pharmaceutical treatment to a patient; and
(3) subsection (d)(11) must be released to the extent disclosure of
the information is not prohibited by applicable federal law.
(f) Before the board releases confidential information under
subsection (d), the applicant must be approved by the INSPECT
program in a manner prescribed by the board.
(g) The board may release to:
(1) a member of the board or another governing body that licenses
practitioners;
(2) an investigator for the consumer protection division of the
office of the attorney general, a prosecuting attorney, the attorney
general, a deputy attorney general, or an investigator from the
office of the attorney general; or
(3) a law enforcement officer who is:
(A) authorized by the state police department to receive
ephedrine, pseudoephedrine, low THC hemp extract, or
controlled substance prescription drug information; and
(B) approved by the board to receive the type of information
released;
confidential information generated from computer records that
identifies practitioners who are prescribing or dispensing large
quantities of a controlled substance.
(h) The information described in subsection (g) may not be released
until it has been reviewed by:
(1) a member of the board who is licensed in the same profession
as the prescribing or dispensing practitioner identified by the data;
or
(2) the board's designee;
and until that member or the designee has certified that further
investigation is warranted. However, failure to comply with this
subsection does not invalidate the use of any evidence that is otherwise
admissible in a proceeding described in subsection (i).
(i) An investigator or a law enforcement officer receiving
confidential information under subsection (c), (d), or (g) may disclose
the information to a law enforcement officer or an attorney for the
office of the attorney general for use as evidence in the following:
(1) A proceeding under IC 16-42-20.

(2) A proceeding under any state or federal law that involves



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1	ephedrine, pseudoephedrine, low THC hemp extract, or a
2	controlled substance.
3	(3) A criminal proceeding or a proceeding in juvenile court that
4	involves ephedrine, pseudoephedrine, low THC hemp extract,
5	or a controlled substance.
6	(j) The board may compile statistical reports from the information
7	described in subsection (a). The reports must not include information
8	that identifies any practitioner, ultimate user, or other person
9	administering ephedrine, pseudoephedrine, low THC hemp extract,
10	or a controlled substance. Statistical reports compiled under this
11	subsection are public records.
12	(k) Except as provided in subsection (q) and in addition to any
13	requirements provided in IC 25-22.5-13, the following practitioners
14	shall obtain information about a patient from the data base before
15	prescribing an opioid or benzodiazepine to the patient:
16	(1) A practitioner who has had the information from the data base
17	integrated into the patient's electronic health records.
18	(2) Beginning January 1, 2019, a practitioner who provides
19	services to the patient in:
20	(A) the emergency department of a hospital licensed under
21	IC 16-21; or
22	(B) a pain management clinic.
23	(3) Beginning January 1, 2020, a practitioner who provides
24	services to the patient in a hospital licensed under IC 16-21.
25	(4) Beginning January 1, 2021, all practitioners.
26	However, a practitioner is not required to obtain information about a
27	patient who is subject to a pain management contract from the data
28	base more than once every ninety (90) days.
29	(1) A practitioner who checks the INSPECT program for the
30	available data on a patient is immune from civil liability for an injury,
31	death, or loss to a person solely due to a practitioner:
32	(1) seeking information from the INSPECT program; and
33	(2) in good faith using the information for the treatment of the
34	patient.
35	The civil immunity described in this subsection does not extend to a
36	practitioner if the practitioner receives information directly from the
37	INSPECT program and then negligently misuses this information. This
38	subsection does not apply to an act or omission that is a result of gross
39	negligence or intentional misconduct.
40	(m) The board may review the records of the INSPECT program. If
41	the board determines that a violation of the law may have occurred, the
42	board shall notify the appropriate law enforcement agency or the



1	relevant government body responsible for the licensure, regulation, or
2	discipline of practitioners authorized by law to prescribe controlled
3	substances.
4	(n) A practitioner who in good faith discloses information based on
5	a report from the INSPECT program to a law enforcement agency is
6	immune from criminal or civil liability. A practitioner that discloses
7	information to a law enforcement agency under this subsection is
8	presumed to have acted in good faith.
9	(o) A practitioner's agent may act as a delegate and check INSPECT
10	program reports on behalf of the practitioner.
11	(p) A patient may access a report from the INSPECT program that
12	has been included in the patient's medical file by a practitioner.
13	(q) A practitioner is not required under subsection (k) to obtain
14	information about a patient from the data base before prescribing an
15	opioid or benzodiazepine if the practitioner has obtained a waiver from
16	the board because the practitioner does not have access to the Internet
17	at the practitioner's place of business.
18	SECTION 10. IC 35-48-7-12.1, AS AMENDED BY P.L.194-2018,
19	SECTION 6, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
20	JULY 1, 2019]: Sec. 12.1. (a) The board shall adopt rules under
21	IC 4-22-2 to implement this chapter, including the following:
22	(1) Information collection and retrieval procedures for the
23	INSPECT program, including the controlled substances to be
24	included in the program required under section 8.1 of this chapter.
25	(2) Design for the creation of the data base required under section
26	10.1 of this chapter.
27	(3) Requirements for the development and installation of online
28	electronic access by the board to information collected by the
29	INSPECT program.
30	(4) Identification of emergency situations or other circumstances
31	in which a practitioner may prescribe, dispense, and administer a
32	prescription drug specified in section 8.1 of this chapter without
33	a written prescription or on a form other than a form specified in
34	section 8.1(a)(4) of this chapter.
35	(5) Requirements for a practitioner providing treatment for a
36	patient at an opioid treatment program operating under
37	IC 12-23-18 to check the INSPECT program:
38	(A) before initially prescribing ephedrine, pseudoephedrine,
39	low THC hemp extract, or a controlled substance to a patient;
40	and
41	(B) periodically during the course of treatment that uses



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ephedrine, pseudoephedrine, low THC hemp extract, or a

1	controlled substance.
2	(b) The board may:
3	(1) set standards for education courses for individuals authorized
4	to use the INSPECT program;
5	(2) identify treatment programs for individuals addicted to
6	controlled substances monitored by the INSPECT program; and
7	(3) work with impaired practitioner associations to provide
8	intervention and treatment.
9	(c) The executive director of the Indiana professional licensing
0	agency may hire a person to serve as the director of the INSPECT
1	program, with the approval of the chairperson of the board.
2	(d) The board shall do the following:
3	(1) Establish a procedure for a practitioner to request a waiver
4	from the requirements of section 11.1(k) of this chapter if the
5	practitioner does not have access to the Internet at the
6	practitioner's place of business.
7	(2) Review a practitioner's written request for a waiver from the
8	requirements of section 11.1(k) of this chapter and determine
9	whether the practitioner should be granted a waiver.
0.	(3) Upon determination by the board under subdivision (2) that a
21	practitioner should be granted a waiver under this subsection,
2	issue the practitioner a waiver.

