

ENGROSSED SENATE BILL No. 340

DIGEST OF SB 340 (Updated February 21, 2018 5:04 pm - DI 107)

Citations Affected: IC 16-18; IC 16-21; IC 16-34; IC 31-34; IC 34-30; IC 35-46; IC 35-52.

Synopsis: Regulation of abortion and newborn safety devices. Makes various changes to the abortion law concerning abortion clinic license applications, abortion clinic inspections, abortion inducing drugs, abortion complications, the provision of information to a woman seeking an abortion, and the collection of data by the state department of health. Specifies under the safe haven law that it is a defense to a claim of neglect of a dependent if the individual left the child in a newborn safety device that is located at a fire department, including a volunteer fire department, that meets specified requirements. Provides civil immunity for a fire department that operates a newborn safety device for an act or omission relating to the device: (1) if the device meets specified requirements; and (2) unless the act or omission constitutes gross negligence or willful or wanton misconduct. Makes a technical correction.

Effective: Upon passage; July 1, 2018.

Holdman, Brown L, Young M,

Zakas, Houchin, Kruse, Crane, Raatz

(HOUSE SPONSORS — MAYFIELD, WESCO, BACON, JUDY)

January 4, 2018, read first time and referred to Committee on Judiciary. January 25, 2018, amended, reported favorably — Do Pass. January 29, 2018, read second time, amended, ordered engrossed. January 30, 2018, engrossed. Read third time, passed. Yeas 38, nays 11.

HOUSE ACTION
February 6, 2018, read first time and referred to Committee on Public Policy.
February 22, 2018, amended, reported — Do Pass.



Second Regular Session 120th General Assembly (2018)

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 2017 Regular Session of the General Assembly.

ENGROSSED SENATE BILL No. 340

A BILL FOR AN ACT to amend the Indiana Code concerning health.

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

1	SECTION 1. IC 16-18-2-1.5, AS AMENDED BY THE
2	TECHNICAL CORRECTIONS BILL OF THE 2018 GENERAL
3	ASSEMBLY, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
4	JULY 1, 2018]: Sec. 1.5. (a) "Abortion clinic", for purposes of
5	IC 16-19-3-31, IC 16-21-2, IC 16-34-2-4.7, IC 16-34-3, and
6	IC 16-41-16, means a health care provider (as defined in section
7	163(d)(1) of this chapter) that:
8	(1) performs surgical abortion procedures; or
9	(2) beginning January 1, 2014, provides an abortion inducing
0	drug for the purpose of inducing an abortion.
1	(b) The term does not include the following:
2	(1) A hospital that is licensed as a hospital under IC 16-21-2.
3	(2) An ambulatory outpatient surgical center that is licensed as an
4	ambulatory outpatient surgical center under IC 16-21-2.
5	(3) A health care provider that provides, prescribes, administers,
6	or dispenses an abortion inducing drug to fewer than five (5)
7	patients per year for the purposes of inducing an abortion.



1	SECTION 2. IC 16-18-2-1.7 IS ADDED TO THE INDIANA CODE
2	AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE JULY
3	1, 2018]: Sec. 1.7. "Abortion complication", for purposes of
4	IC 16-34-2-4.7, has the meaning set forth in IC 16-34-2-4.7.
5	SECTION 3. IC 16-18-2-9.4 IS ADDED TO THE INDIANA CODE
6	AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE JULY
7	1, 2018]: Sec. 9.4. "Affiliate", for purposes of IC 16-21-2-11, means
8	any person who directly or indirectly controls, is controlled by, or
9	is under common control of another person.
10	SECTION 4. IC 16-21-2-2.5, AS AMENDED BY P.L.173-2017,
11	SECTION 2, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
12	UPON PASSAGE]: Sec. 2.5. (a) The state department shall adopt rules
13	under IC 4-22-2 to do the following concerning birthing centers and
14	abortion clinics:
15	(1) Establish minimum license qualifications.
16	(2) Establish the following requirements:
17	(A) Sanitation standards.
18	(B) Staff qualifications.
19	(C) Necessary emergency equipment.
20	(D) Procedures to provide emergency care.
21	(E) Procedures to monitor patients after the administration of
22	anesthesia.
23	(F) Procedures to provide follow-up care for patient
24 25	complications.
	(G) Quality assurance standards.
26	(H) Infection control.
27	(I) Provision of informed consent brochures, as described in
28	IC 16-34-2-1.5, in English, Spanish, and a third language
29	determined by the state department, inside abortion clinics.
30	(J) Provision of a hotline telephone number that provides
31	assistance for patients who are:
32	(i) coerced into an abortion; or
33	(ii) victims of sex trafficking.
34	(K) Annual training by law enforcement officers on identifying
35	and assisting women who are:
36	(i) coerced into an abortion; or
37	(ii) victims of sex trafficking.
38	(3) Prescribe the operating policies, supervision, and maintenance
39	of medical records, including the requirement that all forms that
40	require a patient signature be stored in the patient's medical
41	record.
12	(1) Establish procedures for the issuance renewal denial and



1	revocation of licenses under this chapter. The rules adopted under
2	this subsection must address the following:
3	(A) The form and content of the license.
4	(B) The collection of an annual license fee.
5	(5) Prescribe the procedures and standards for inspections.
6	(6) Prescribe procedures for:
7	(A) implementing a plan of correction to address any
8	violations of any provision of this chapter or any rules adopted
9	under this chapter; and
10	(B) implementing a system for the state department to follow
11	if the abortion clinic or birthing center fails to comply with the
12	plan of correction described in clause (A) and disciplinary
13	action is needed.
14	(b) A person who knowingly or intentionally:
15	(1) operates a birthing center or an abortion clinic that is not
16	licensed under this chapter; or
17	(2) advertises the operation of a birthing center or an abortion
18	clinic that is not licensed under this chapter;
19	commits a Class A misdemeanor.
20	(c) Not later than January 1, 2019, the state department shall:
21	(1) adopt separate rules under IC 4-22-2, including those required
21 22 23 24	under subsection (a), for existing and future abortion clinics that
23	perform only surgical abortions;
24	(2) adopt separate rules under IC 4-22-2, including those required
25	under subsection (a), for existing and future abortion clinics that
26	perform abortions only through the provision of an abortion
27 28	inducing drug; and
	(3) establish procedures regarding the issuance of licenses to
29	existing and future abortion clinics that:
30	(A) perform only surgical abortions;
31	(B) perform abortions only through the provision of an
32	abortion inducing drug; or
33	(C) perform both surgical abortions and abortions through the
34	provision of abortion inducing drugs.
35	(d) A rule or emergency rule adopted under subsection (c)(1).
36	(c)(2), or (c)(3) applies, respectively, to every abortion clinic of the
37	type described in subsection (c)(1), (c)(2), or (c)(3), regardless of
38	the date of adoption of the rule or emergency rule.
39	(e) Before January 1, 2019, the state department shall adopt
40	emergency rules in the manner provided under IC 4-22-2-37.1 to
41	carry out the duties established in this section under the following:
42	(1) Subsection (a)(2)(E).



1	(2) Subsection (a)(2)(F).
2	(3) Subsection (a)(2)(I).
3	(4) Subsection (a)(2)(J).
4	(5) Subsection (a)(2)(K).
5	(6) Subsection (a)(3).
6	(7) Subsection (a)(5).
7	(8) Subsection (a)(6).
8	This subsection expires July 1, 2019.
9	SECTION 5. IC 16-21-2-2.6, AS ADDED BY P.L.98-2014,
10	SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
11	JULY 1, 2018]: Sec. 2.6. The state department may shall inspect an
12	abortion clinic at least one (1) time per calendar year and may conduct
13	a complaint inspection as needed.
14	SECTION 6. IC 16-21-2-11, AS AMENDED BY P.L.172-2011,
15	SECTION 114, IS AMENDED TO READ AS FOLLOWS
16	[EFFECTIVE JULY 1, 2018]: Sec. 11. (a) An applicant must submit
17	an application for a license on a form prepared by the state department
18	showing that:
19	(1) the applicant is of reputable and responsible character;
20	(2) the applicant is able to comply with the minimum standards
21	for a hospital, an ambulatory outpatient surgical center, an
22	abortion clinic, or a birthing center, and with rules adopted under
23	this chapter; and
22 23 24	(3) the applicant has complied with section 15.4 of this chapter.
25	(b) The application must contain the following additional
26	information:
27	(1) The name of the applicant.
28	(2) The type of institution to be operated.
29	(3) The location of the institution.
30	(4) The name of the person to be in charge of the institution.
31	(5) If the applicant is a hospital, the range and types of services to
32	be provided under the general hospital license, including any
33	service that would otherwise require licensure by the state
34	department under the authority of IC 16-19.
35	(6) Other information the state department requires.
36	(c) If the department of state revenue notifies the department that a
37	person is on the most recent tax warrant list, the department shall not
38	issue or renew the person's license until:
39	(1) the person provides to the department a statement from the
40	department of state revenue that the person's tax warrant has been
41	satisfied; or
42	(2) the department receives a notice from the commissioner of the



1	department of state revenue under IC 6-8.1-8-2(k).
2	(d) An application for an abortion clinic license must require the
3	applicant to do the following:
4	(1) Disclose whether the applicant, or an owner or affiliate of
5	the applicant, operated an abortion clinic that was closed as
6	a direct result of patient health and safety concerns.
7	(2) Disclose whether a principal or clinic staff member was
8	convicted of a felony.
9	(3) Disclose whether a principal or clinic staff member was
10	ever employed by a facility owned or operated by the
11	applicant that closed as a result of administrative or legal
12	action.
13	(4) Provide copies of:
14	(A) administrative and legal documentation relating to the
15	information required under subdivisions (1) and (2);
16	(B) inspection reports; and
17	(C) violation remediation contracts;
18	if any.
19	SECTION 7. IC 16-34-2-1, AS AMENDED BY P.L.213-2016,
20	SECTION 13, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
21	JULY 1, 2018]: Sec. 1. (a) Abortion shall in all instances be a criminal
22	act, except when performed under the following circumstances:
23	(1) Except as prohibited in IC 16-34-4, during the first trimester
24	of pregnancy for reasons based upon the professional, medical
25	judgment of the pregnant woman's physician if:
26	(A) the abortion is performed by the physician;
27	(B) the woman submitting to the abortion has filed her consent
28	with her physician. However, if in the judgment of the
29	physician the abortion is necessary to preserve the life of the
30	woman, her consent is not required; and
31	(C) the woman submitting to the abortion has filed with her
32	physician the written consent of her parent or legal guardian
33	if required under section 4 of this chapter.
34	However, an abortion inducing drug may not be dispensed,
35	prescribed, administered, or otherwise given to a pregnant woman
36	after nine (9) weeks of postfertilization age unless the Food and
37	Drug Administration has approved the abortion inducing drug to
38	be used for abortions later than nine (9) weeks of postfertilization
39	age. A physician shall examine a pregnant woman in person
40	before prescribing or dispensing an abortion inducing drug. In
41	accordance with FDA guidelines, the physician shall provide

the pregnant woman with a copy of the manufacturer's



1	instruction sheets and require that the pregnant woman sign
2	the manufacturer's patient agreement form. The physician
3	shall retain a copy of the signed patient agreement form, and
4	the signed physician's agreement form required by the
5	manufacturer, in the patient's file. As used in this subdivision,
6	"in person" does not include the use of telehealth or telemedicine
7 8	services.
	(2) Except as prohibited by IC 16-34-4, for an abortion performed
9	by a surgical procedure, after the first trimester of pregnancy and
10	before the earlier of viability of the fetus or twenty (20) weeks of
11	postfertilization age, for reasons based upon the professional,
12	medical judgment of the pregnant woman's physician if:
13 14	(A) all the circumstances and provisions required for legal abortion during the first trimester are present and adhered to;
15	and
16	(B) the abortion is performed in a hospital or ambulatory
17	outpatient surgical center (as defined in IC 16-18-2-14).
18	(3) Except as provided in subsection (b) or as prohibited by
19	IC 16-34-4, and for an abortion performed by a surgical
20	procedure, at the earlier of viability of the fetus or twenty (20)
21	weeks of postfertilization age and any time after, for reasons
22	based upon the professional, medical judgment of the pregnant
23	woman's physician if:
24	(A) all the circumstances and provisions required for legal
25	abortion before the earlier of viability of the fetus or twenty
26	(20) weeks of postfertilization age are present and adhered to;
27	(B) the abortion is performed in compliance with section 3 of
28	this chapter; and
29	(C) before the abortion the attending physician shall certify in
30	writing to the hospital in which the abortion is to be
31	performed, that in the attending physician's professional,
32	medical judgment, after proper examination and review of the
33	woman's history, the abortion is necessary to prevent a
34	substantial permanent impairment of the life or physical health
35	of the pregnant woman. All facts and reasons supporting the
36	certification shall be set forth by the physician in writing and
37	attached to the certificate.
38	(b) A person may not knowingly or intentionally perform a partial
39	birth abortion unless a physician reasonably believes that:
40	(1) performing the partial birth abortion is necessary to save the
41	mother's life; and

(2) no other medical procedure is sufficient to save the mother's



1	life.
2	SECTION 8. IC 16-34-2-1.5, AS AMENDED BY P.L.156-2017,
3	SECTION 14, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
4	JULY 1, 2018]: Sec. 1.5. (a) The state department shall develop an
5	informed consent brochure and post the informed consent brochure on
6	the state department's Internet web site.
7	(b) The state department shall develop an informed consent
8	brochure that includes the following:
9	(1) Objective scientific information concerning the probable
10	anatomical and physiological characteristics of a fetus every two
11	(2) weeks of gestational age, including the following:
12	(A) Realistic pictures in color for each age of the fetus,
13	including the dimensions of the fetus.
14	(B) Whether there is any possibility of the fetus surviving
15	outside the womb.
16	(2) Objective scientific information concerning the medical risks
17	associated with each abortion procedure or the use of an abortion
18	inducing drug, including the following:
19	(A) The risks of infection and hemorrhaging.
20	(B) The potential danger:
21	(i) to a subsequent pregnancy; or
22	(ii) of infertility.
22 23 24	(3) Information concerning the medical risks associated with
	carrying the child to term.
25	(4) Information that medical assistance benefits may be available
26	for prenatal care, childbirth, and neonatal care.
27	(5) Information that the biological father is liable for assistance in
28	support of the child, regardless of whether the biological father
29	has offered to pay for an abortion.
30	(6) Information regarding telephone 211 dialing code services for
31	accessing human services as described in IC 8-1-19.5, and the
32	types of services that are available through this service.
33	(7) Information concerning Indiana's safe haven law under
34	IC 31-34-2.5-1.
35	(8) Information that, under certain conditions, a pregnant
36	woman may relinquish a child who is, or who appears to be,
37	not more than thirty (30) days of age:
38	(A) to an emergency medical services provider (as defined
39	in IC 16-41-10-1); or
10	(B) in a newborn safety device (described in
1 1	IC 31-34-2.5-1) at a participating fire department or other
12	site that is staffed by an emergency medical services



1	provider.
2	(c) In complying with subsection (b)(6), the state department shall
3	consult with the recognized 211 service providers and the Indiana
4	housing and community development authority as required by
5	IC 8-1-19.5-9.
6	(d) In the development of the informed consent brochure described
7	in this section, the state department shall use information and pictures
8	that are available at no cost or nominal cost to the state department.
9	(e) The informed consent brochure must include the requirements
10	specified in this chapter.
11	SECTION 9. IC 16-34-2-4.7 IS ADDED TO THE INDIANA CODE
12	AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE JULY
13	1, 2018]: Sec. 4.7. (a) As used in this section, "abortion
14	complication" means any adverse physical or psychological
15	condition arising from the induction or performance of an
16	abortion. The term includes the following:
17	(1) Uterine perforation.
18	(2) Cervical perforation.
19	(3) Infection.
20	(4) Hemorrhaging.
21	(5) Blood clots.
22	(6) Failure to terminate the pregnancy.
23	(7) Incomplete abortion (retained tissue).
24	(8) Pelvic inflammatory disease.
25	(9) Missed ectopic pregnancy.
26	(10) Cardiac arrest.
27	(11) Respiratory arrest.
28	(12) Renal failure.
29	(13) Metabolic disorder.
30	(14) Shock.
31	(15) Embolism.
32	(16) Coma.
33	(17) Placenta previa in subsequent pregnancies.
34	(18) Pre-term delivery in subsequent pregnancies.
35	(19) Free fluid in the abdomen.
36	(20) Hemolytic reaction due to the administration of
37	ABO-incompatible blood or blood products.
38	(21) Hypoglycemia occurring while the patient is being
39	treated at the abortion facility.
40	(22) Physical injury associated with treatment performed at
41	the abortion facility.
42	(23) Adverse reaction to anesthesia or other drugs.



1	(24) Psychological or emotional complications, including
2 3	depression, suicidal ideation, anxiety, and sleeping disorders.
	(25) Death.
4	(26) Any other adverse event as defined by criteria provided
5	in the Food and Drug Administration Safety Information and
6	Adverse Event Reporting Program.
7	(b) The following persons shall report to the state department
8	each case in which the person treated a patient suffering from an
9	abortion complication:
10	(1) A physician licensed under IC 25-22.5.
11	(2) A hospital licensed under IC 16-21.
12	(3) An abortion clinic licensed under IC 16-21-2-2.5.
13	(c) The state department shall develop a process for the
14	submission of a report under this section.
15	(d) A report under this section shall be submitted to the state
16	department in the manner prescribed by the state department.
17	(e) The report under this section must include the following
18	information concerning the abortion complication:
19	(1) The date the patient presented for treatment for the
20	abortion complication.
21	(2) The age of the patient.
22	(3) The race of the patient.
23	(4) The county and state of the patient's residence.
24	(5) The type of abortion obtained by the patient.
25	(6) The date of abortion obtained by the patient.
26	(7) The name of the:
27	(A) abortion clinic;
28	(B) medical facility; or
29	(C) hospital;
30	where the patient obtained the abortion.
31	(8) Whether the patient obtained abortion medication via mail
32	order or Internet web site, and if so, information identifying
33	the source of the medication.
34	(9) Whether the complication was previously managed by the
35	abortion provider or the abortion provider's required
36	back-up physician.
37	(10) The name of the medications taken by the patient as part
38	of the pharmaceutical abortion regimen, if any.
39	(11) A list of each diagnosed complication.
40	(12) A list of each treated complication, with a description of
41	the treatment provided.
42	(13) Whether the patient's visit to treat the complications was



1	the original visit or a follow-up v	isit.
2	(14) The date of each follow-up v	isit.

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- (14) The date of each follow-up visit, if any.
- (15) A list of each complication diagnosed at a follow-up visit,
- (16) A list of each complication treated at a follow-up visit, if
- (f) Before February 1, 2019, the state department shall inform in writing all providers described in subsection (b) of the new reporting requirements for abortion complications. This subsection expires December 31, 2019.
- (g) Not later than June 30 of each year, the state department shall compile a public report summarizing the information collected under this section. The report must include statistics for the previous calendar year, with updated information for the most recent calendar year.
- (h) The state department shall summarize the aggregate data from the data submitted under this section and submit the data, on or before June 30 of each year, to the United States Centers for Disease Control and Prevention for its inclusion in the annual Vital Statistics Report.
- (i) The state department shall ensure that no identifying information of a pregnant woman is included in the report described in subsection (g).
- (j) This subsection applies after August 31, 2019. Each failure to report an abortion complication as required under this section is a Class B misdemeanor.
- (k) Before January 1, 2019, the state department shall adopt rules under IC 4-22-2 to implement this section.

SECTION 10. IC 16-34-2-5, AS AMENDED BY P.L.173-2017, SECTION 6, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2018]: Sec. 5. (a) Every health care provider who performs a surgical abortion or provides, prescribes, administers, or dispenses an abortion inducing drug for the purposes of inducing an abortion shall report the performance of the abortion or the provision, prescribing, administration, or dispensing of an abortion inducing drug on a form drafted by the state department, the purpose and function of which shall be the improvement of maternal health and life through the compilation of relevant maternal life and health factors and data, and a further purpose and function shall be to monitor all abortions performed in Indiana to assure the abortions are done only under the authorized provisions of the law. For each abortion performed and abortion inducing drug provided, prescribed, administered, or dispensed, the



1	report shall include, among other things, the following:
2	(1) The age of the patient.
3	(2) Whether a waiver of consent under section 4 of this
4	chapter was obtained.
5	(3) Whether a waiver of notification under section 4 of this
6	chapter was obtained.
7	(2) (4) The date and location the abortion was performed or the
8	abortion inducing drug was provided, prescribed, administered,
9	or dispensed.
10	(3) (5) The health care provider's full name and address, including
11	the name of the physicians performing the abortion or providing,
12	prescribing, administering, or dispensing the abortion inducing
13	drug.
14	(4) The name of the father if known.
15	(6) The city and county where the pregnancy termination
16	occurred.
17	(5) (7) The age of the father, or the approximate age of the father
18	if the father's age is unknown.
19	(8) The patient's county and state of residence.
20	(9) The marital status of the patient.
21	(10) The educational level of the patient.
22	(11) The race of the patient.
23	(12) The ethnicity of the patient.
24	(13) The number of the patient's previous live births.
25	(14) The number of the patient's deceased children.
26	(15) The number of the patient's spontaneous pregnancy
27	terminations.
28	(16) The number of the patient's previous induced
29	terminations.
30	(17) The date of the patient's last menses.
31	(18) The physician's determination of the gestation of the fetus
32	in weeks.
33	(19) Whether the patient indicated that the patient was
34	seeking an abortion as a result of being:
35	(A) abused;
36	(B) coerced;
37	(C) harassed; or
38	(D) trafficked.
39 40	(6) (20) The following information concerning the abortion or the
40 41	provision, prescribing, administration, or dispensing of the
41 42	abortion inducing drug:
4/	(A) The postfertilization age of the fetus (in weeks)



1	(B) The manner in which the postfertilization age was
2	determined.
3	(C) The gender of the fetus, if detectable.
4	(D) Whether the fetus has been diagnosed with or has a
5	potential diagnosis of having Down syndrome or any other
6	disability.
7	(E) If after the earlier of the time the fetus obtains viability or
8	the time the postfertilization age of the fetus is at least twenty
9	(20) weeks, the medical reason for the performance of the
10	abortion or the provision, prescribing, administration, or
11	dispensing of the abortion inducing drug.
12	(7) (21) For a surgical abortion, the medical procedure used for
13	the abortion and, if the fetus was viable or had a postfertilization
14	age of at least twenty (20) weeks:
15	(A) whether the procedure, in the reasonable judgment of the
16	health care provider, gave the fetus the best opportunity to
17	survive; and
18	(B) the basis for the determination that the pregnant woman
19	had a condition described in this chapter that required the
20	abortion to avert the death of or serious impairment to the
21	pregnant woman; and
22	(C) the name of the second doctor present, as required
22 23	(C) the name of the second doctor present, as required under IC 16-34-2-3(a)(3).
23	under IC 16-34-2-3(a)(3).
23 24	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided,
23 24 25	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery
23 24 25 26	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient.
23 24 25 26 27	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's
23 24 25 26 27 28	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient
23 24 25 26 27 28 29	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement.
23 24 25 26 27 28 29 30	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical
23 24 25 26 27 28 29 30 31	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother.
23 24 25 26 27 28 29 30 31 32	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother. (10) (25) The mother's obstetrical history, including dates of other
23 24 25 26 27 28 29 30 31 32 33	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother. (10) (25) The mother's obstetrical history, including dates of other abortions, if any.
23 24 25 26 27 28 29 30 31 32 33 34	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother. (10) (25) The mother's obstetrical history, including dates of other abortions, if any. (26) Any preexisting medical conditions of the patient that
23 24 25 26 27 28 29 30 31 32 33 34 35	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother. (10) (25) The mother's obstetrical history, including dates of other abortions, if any. (26) Any preexisting medical conditions of the patient that may complicate the abortion.
23 24 25 26 27 28 29 30 31 32 33 34 35 36	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother. (10) (25) The mother's obstetrical history, including dates of other abortions, if any. (26) Any preexisting medical conditions of the patient that may complicate the abortion. (11) (27) The results of pathological examinations if performed.
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother. (10) (25) The mother's obstetrical history, including dates of other abortions, if any. (26) Any preexisting medical conditions of the patient that may complicate the abortion. (11) (27) The results of pathological examinations if performed. (12) (28) For a surgical abortion, whether the fetus was delivered
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother. (10) (25) The mother's obstetrical history, including dates of other abortions, if any. (26) Any preexisting medical conditions of the patient that may complicate the abortion. (11) (27) The results of pathological examinations if performed. (12) (28) For a surgical abortion, whether the fetus was delivered alive, and if so, how long the fetus lived.
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39	under IC 16-34-2-3(a)(3). (8) (22) For a nonsurgical abortion, the precise drugs provided, prescribed, administered, or dispensed, and the means of delivery of the drugs to the patient. (23) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient signed the patient agreement. (9) (24) For an early pre-viability termination, the medical indication by diagnosis code for the fetus and the mother. (10) (25) The mother's obstetrical history, including dates of other abortions, if any. (26) Any preexisting medical conditions of the patient that may complicate the abortion. (11) (27) The results of pathological examinations if performed. (12) (28) For a surgical abortion, whether the fetus was delivered alive, and if so, how long the fetus lived. (13) (29) Records of all maternal deaths occurring at the location



1	department and, if applicable, separately to the department of
2	child services.
3	(b) The health care provider shall complete the form provided for in
4	subsection (a) and shall transmit the completed form to the state
5	department, in the manner specified on the form, within thirty (30) days
6	after the date of each abortion. However, if an abortion is for a female
7	who is less than sixteen (16) years of age, the health care provider shall
8	transmit the form to the state department of health and separately to the
9	department of child services within three (3) days after the abortion is
10	performed.
11	(c) The dates supplied on the form may not be redacted for any
12	reason before the form is transmitted as provided in this section.
13	(d) Each failure to complete or timely transmit a form, as required
14	under this section, for each abortion performed or abortion inducing
15	drug that was provided, prescribed, administered, or dispensed, is a
16	Class B misdemeanor.
17	(e) Not later than June 30 of each year, the state department shall
18	compile a public report providing the following:
19	(1) Statistics for the previous calendar year from the information
20	submitted under this section.
21	(2) Statistics for previous calendar years compiled by the state
22	department under this subsection, with updated information for
23	the calendar year that was submitted to the state department after
24	the compilation of the statistics.
25	The state department shall ensure that no identifying information of a
26	pregnant woman is contained in the report.
27	(f) The state department shall:
28	(1) summarize aggregate data from all data submitted under
29	this section; and
30	(2) submit the data, before July 1 of each year, to the United
31	States Centers for Disease Control and Prevention for its
32	inclusion in the annual Vital Statistics Report.
33	SECTION 11. IC 31-34-2.5-1, AS AMENDED BY P.L.186-2017,
34	SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
35	JULY 1, 2018]: Sec. 1. (a) An emergency medical services provider (as
36	defined in IC 16-41-10-1) shall, without a court order, take custody of
37	a child who is, or who appears to be, not more than thirty (30) days of
38	age if:
39	(1) the child is voluntarily left:
40	(A) with the provider by the child's parent;
41	(B) in a newborn safety device that:

(i) has been approved by a hospital licensed under IC 16-21;



1	(ii) is physically located inside a hospital that is staffed
2	continuously on a twenty-four (24) hour basis every day to
3	provide care to patients in an emergency; and
4	(iii) is located in an area that is conspicuous and visible to
5	hospital staff; or
6	(C) in a newborn safety device that was installed on or before
7	January 1, 2017, and is located at a site that is staffed by an
8	emergency medical services provider (as defined in
9	IC 16-41-10-1); or
10	(D) in a newborn safety device that:
11	(i) is located at a fire department, including a volunteer
12	fire department, that is staffed by an emergency medical
13	services provider (as defined in IC 16-41-10-1) on a
14	twenty-four (24) hour seven (7) day a week basis;
15	(ii) is located in an area that is conspicuous and visible to
16	staff; and
17	(iii) includes an adequate dual alarm system connected
18	to the site that is tested at least one (1) time per month to
19	ensure the alarm system is in working order; and
20	(2) the parent does not express an intent to return for the child.
21	(b) An emergency medical services provider who takes custody of
22	a child under this section shall perform any act necessary to protect the
23	child's physical health or safety.
24	(c) Any person who in good faith voluntarily leaves a child:
25	(1) with an emergency medical services provider; or
26	(2) in a newborn safety device described in subsection (a)(1)(B);
27	is not obligated to disclose the parent's name or the person's name.
28	(d) A hospital that approves the operation of a newborn safety
29	device that meets the requirements set forth in subsection (a)(1)(B) is
30	immune from civil liability for an act or omission relating to the
31	operation of the newborn safety device unless the act or omission
32	constitutes gross negligence or willful or wanton misconduct.
33	(e) A newborn safety device described in subsection (a)(1)(C) may
34	continue to operate without meeting the conditions set forth in
35	subsection (a) (1) (B).
36	(f) A fire department, including a volunteer fire department,
37	that meets the requirements set forth in subsection (a)(1)(D) is
38	immune from civil liability for an act or omission relating to the
39	operation of the newborn safety device unless the act or omission
40	constitutes gross negligence or willful or wanton misconduct.
41	SECTION 12. IC 34-30-2-134.5, AS ADDED BY P.L.263-2017,

SECTION 2, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE



1	JULY 1, 2018]: Sec. 134.5. IC 31-34-2.5-1 (Concerning a hospital or
2	fire department operating a newborn safety device).
3	SECTION 13. IC 35-46-1-4, AS AMENDED BY P.L.252-2017,
4	SECTION 17, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
5	JULY 1, 2018]: Sec. 4. (a) A person having the care of a dependent,
6	whether assumed voluntarily or because of a legal obligation, who
7	knowingly or intentionally:
8	(1) places the dependent in a situation that endangers the
9	dependent's life or health;
10	(2) abandons or cruelly confines the dependent;
11	(3) deprives the dependent of necessary support; or
12	(4) deprives the dependent of education as required by law;
13	commits neglect of a dependent, a Level 6 felony.
14	(b) However, the offense is:
15	(1) a Level 5 felony if it is committed under subsection (a)(1),
16	(a)(2), or (a)(3) and:
17	(A) results in bodily injury; or
18	(B) is:
19	(i) committed in a location where a person is violating
20	IC 35-48-4-1 (dealing in cocaine or a narcotic drug),
21	IC 35-48-4-1.1 (dealing in methamphetamine), or
22	IC 35-48-4-1.2 (manufacturing methamphetamine); or
23	(ii) the result of a violation of IC 35-48-4-1 (dealing in
24	cocaine or a narcotic drug), IC 35-48-4-1.1 (dealing in
25	methamphetamine), or IC 35-48-4-1.2 (manufacturing
26	methamphetamine);
27	(2) a Level 3 felony if it is committed under subsection (a)(1),
28	(a)(2), or (a)(3) and results in serious bodily injury;
29	(3) a Level 1 felony if it is committed under subsection (a)(1),
30	(a)(2), or (a)(3) by a person at least eighteen (18) years of age and
31	results in the death of a dependent who is less than fourteen (14)
32	years of age or in the death of a dependent of any age who has a
33	mental or physical disability; and
34	(4) a Level 5 felony if it is committed under subsection (a)(2) and
35	consists of cruel confinement or abandonment that:
36	(A) deprives a dependent of necessary food, water, or sanitary
37	facilities;
38	(B) consists of confinement in an area not intended for human
39	habitation; or
40	(C) involves the unlawful use of handcuffs, a rope, a cord,
41	tape, or a similar device to physically restrain a dependent.
42	(c) It is a defense to a prosecution based on an alleged act under this



1	section that:
2	(1) the accused person left a dependent child who was, at the time
3	the alleged act occurred, not more than thirty (30) days of age:
4	(A) in a newborn safety device described in
5	IC $31-34-2.5-1(a)(1)(B)$, or IC $31-34-2.5-1(a)(1)(C)$, or
6	IC 31-34-2.5-1(a)(1)(D); or
7	(B) with a person who is an emergency medical services
8	provider (as defined in IC 16-41-10-1) who took custody of the
9	child under IC 31-34-2.5;
10	when the prosecution is based solely on the alleged act of leaving
11	the child in the newborn safety device or with the emergency
12	medical services provider and the alleged act did not result in
13	bodily injury or serious bodily injury to the child; or
14	(2) the accused person, in the legitimate practice of the accused
15	person's religious belief, provided treatment by spiritual means
16	through prayer, in lieu of medical care, to the accused person's
17	dependent.
18	(d) Except for property transferred or received:
19	(1) under a court order made in connection with a proceeding
20	under IC 31-15, IC 31-16, IC 31-17, or IC 31-35 (or IC 31-1-11.5
21	or IC 31-6-5 before their repeal); or
22	(2) under section 9(d) of this chapter;
23	a person who transfers or receives any property in consideration for the
24	termination of the care, custody, or control of a person's dependent
25	child commits child selling, a Level 6 felony.
26	SECTION 14. IC 35-52-16-20.9 IS ADDED TO THE INDIANA
27	CODE AS A NEW SECTION TO READ AS FOLLOWS
28	[EFFECTIVE JULY 1, 2018]: Sec. 20.9. IC 16-34-2-4.7 defines a
29	crime concerning abortion.
30	SECTION 15. An emergency is declared for this act.



COMMITTEE REPORT

Madam President: The Senate Committee on Judiciary, to which was referred Senate Bill No. 340, has had the same under consideration and begs leave to report the same back to the Senate with the recommendation that said bill be AMENDED as follows:

Page 2, between lines 4 and 5, begin a new paragraph and insert: "SECTION 3. IC 16-18-2-9.4 IS ADDED TO THE INDIANA CODE AS A NEW SECTION TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2018]: Sec. 9.4. "Affiliate", for purposes of IC 16-21-2-11, means any person who directly or indirectly controls, is controlled by, or is under common control of another person."

Page 3, line 34, delete "July 1, 2018," and insert "**January 1, 2019,**".

Page 3, between lines 40 and 41, begin a new line block indented and insert:

"(5) Subsection (a)(2)(K).".

Page 3, line 41, delete "(5)" and insert "(6)".

Page 3, line 42, delete "(6)" and insert "(7)".

Page 4, line 1, delete "(7)" and insert "(8)".

Page 4, line 40, delete "applicant" and insert "applicant, or an owner or affiliate of the applicant,".

Page 5, between lines 1 and 2, begin a new line block indented and insert:

"(3) Disclose whether a principal or clinic staff member was ever employed by a facility owned or operated by the applicant that closed as a result of administrative or legal action."

Page 5, line 2, delete "(3)" and insert "(4)".

Page 5, line 29, delete "The" and insert "In accordance with FDA guidelines, the".

Page 5, line 31, delete "request" and insert "require".

Page 5, line 32, delete "a" and insert "the manufacturer's".

Page 5, line 34, delete "form" and insert "form, and the signed physician's agreement form required by the manufacturer,".

Page 5, line 34, after "file." insert "The department shall develop a form attesting that the patient was given the manufacturer's instruction materials and signed the patient agreement form before the medication was dispensed to her, in accordance with FDA requirements. The patient shall sign the form and the form shall be retained in the patient's file."



Page 6, between lines 29 and 30, begin a new paragraph and insert: "SECTION 8. IC 16-34-2-1.1, AS AMENDED BY P.L.213-2016, SECTION 14, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2018]: Sec. 1.1. (a) An abortion shall not be performed except with the voluntary and informed consent of the pregnant woman upon whom the abortion is to be performed. Except in the case of a medical emergency, consent to an abortion is voluntary and informed only if the following conditions are met:

- (1) At least eighteen (18) hours before the abortion and in the private, not group, presence of the pregnant woman, the physician who is to perform the abortion, the referring physician or a physician assistant (as defined in IC 25-27.5-2-10), an advanced practice nurse (as defined in IC 25-23-1-1(b)), or a certified nurse midwife (as defined in IC 34-18-2-6.5) to whom the responsibility has been delegated by the physician who is to perform the abortion or the referring physician has informed the pregnant woman orally and in writing of the following:
 - (A) The name of the physician performing the abortion, the physician's medical license number, and an emergency telephone number where the physician or the physician's designee may be contacted on a twenty-four (24) hour a day, seven (7) day a week basis.
 - (B) That follow-up care by the physician or the physician's designee (if the designee is licensed under IC 25-22.5) is available on an appropriate and timely basis when clinically necessary.
 - (C) The nature of the proposed procedure or information concerning the abortion inducing drug.
 - (D) Objective scientific information of the risks of and alternatives to the procedure or the use of an abortion inducing drug, including:
 - (i) the risk of infection and hemorrhage;
 - (ii) the potential danger to a subsequent pregnancy; and
 - (iii) the potential danger of infertility.
 - (E) That human physical life begins when a human ovum is fertilized by a human sperm.
 - (F) The probable gestational age of the fetus at the time the abortion is to be performed, including:
 - (i) a picture of a fetus;
 - (ii) the dimensions of a fetus; and
 - (iii) relevant information on the potential survival of an unborn fetus;



at this stage of development.

- (G) That objective scientific information shows that a fetus can feel pain at or before twenty (20) weeks of postfertilization age.
- (H) The medical risks associated with carrying the fetus to term.
- (I) The availability of fetal ultrasound imaging and auscultation of fetal heart tone services to enable the pregnant woman to view the image and hear the heartbeat of the fetus and how to obtain access to these services.
- (J) That the pregnancy of a child less than fifteen (15) years of age may constitute child abuse under Indiana law if the act included an adult and must be reported to the department of child services or the local law enforcement agency under IC 31-33-5.
- (K) That Indiana does not allow a fetus to be aborted solely because of the fetus's race, color, national origin, ancestry, sex, or diagnosis or potential diagnosis of the fetus having Down syndrome or any other disability.
- (2) At least eighteen (18) hours before the abortion, the pregnant woman will be informed orally and in writing of the following:
 - (A) That medical assistance benefits may be available for prenatal care, childbirth, and neonatal care from the county office of the division of family resources.
 - (B) That the father of the unborn fetus is legally required to assist in the support of the child. In the case of rape, the information required under this clause may be omitted.
 - (C) That adoption alternatives are available and that adoptive parents may legally pay the costs of prenatal care, childbirth, and neonatal care.
 - (D) That there are physical risks to the pregnant woman in having an abortion, both during the abortion procedure and after.
 - (E) That Indiana has enacted the safe haven law under IC 31-34-2.5.
 - (F) The:
 - (i) Internet web site address of the state department of health's web site; and
 - (ii) description of the information that will be provided on the web site and that are;

described in section 1.5 of this chapter.

(G) For the facility in which the abortion is to be performed,



- an emergency telephone number that is available and answered on a twenty-four (24) hour a day, seven (7) day a week basis.
- (H) On a form developed by the state department and as described in IC 16-34-3, that the pregnant woman has a right to determine the final disposition of the remains of the aborted fetus
- (I) On a form developed by the state department, information concerning the available options for disposition of the aborted fetus.
- (J) On a form developed by the state department, information concerning any counseling that is available to a pregnant woman after having an abortion.

The state department shall develop and distribute the forms required by clauses (H) through (J).

- (3) The pregnant woman certifies in writing, on a form developed by the state department, before the abortion is performed, that:
 - (A) the information required by subdivisions (1) and (2) has been provided to the pregnant woman;
 - (B) the pregnant woman has been offered by the provider the opportunity to view the fetal ultrasound imaging and hear the auscultation of the fetal heart tone if the fetal heart tone is audible and that the woman has:
 - (i) viewed or refused to view the offered fetal ultrasound imaging; and
 - (ii) listened to or refused to listen to the offered auscultation of the fetal heart tone if the fetal heart tone is audible; and
 - (C) the pregnant woman has been given a written copy of the printed materials described in section 1.5 of this chapter.
- (4) At least eighteen (18) hours before the abortion and in the presence of the pregnant woman, the physician who is to perform the abortion, the referring physician or a physician assistant (as defined in IC 25-27.5-2-10), an advanced practice nurse (as defined in IC 25-23-1-1(b)), or a midwife (as defined in IC 34-18-2-19) to whom the responsibility has been delegated by the physician who is to perform the abortion or the referring physician has provided the pregnant woman with a color copy of the informed consent brochure described in section 1.5 of this chapter by printing the informed consent brochure from the state department's Internet web site and including the following information on the back cover of the brochure:
 - (A) The name of the physician performing the abortion and the



physician's medical license number.

- (B) An emergency telephone number where the physician or the physician's designee may be contacted twenty-four (24) hours a day, seven (7) days a week.
- (C) A statement that follow-up care by the physician or the physician's designee who is licensed under IC 25-22.5 is available on an appropriate and timely basis when clinically necessary.
- (5) At least eighteen (18) hours before an abortion is performed and at the same time that the pregnant woman receives the information required by subdivision (1), the provider shall perform, and the pregnant woman shall view, the fetal ultrasound imaging and hear the auscultation of the fetal heart tone if the fetal heart tone is audible unless the pregnant woman certifies in writing, on a form developed by the state department, before the abortion is performed, that the pregnant woman:
 - (A) does not want to view the fetal ultrasound imaging; and
 - (B) does not want to listen to the auscultation of the fetal heart tone if the fetal heart tone is audible.
- (6) The pregnant woman certifies in writing, on a form developed by the state department with a copy kept in her patient file, that she has been informed:
 - (A) of Indiana's safe haven law under IC 31-34-2.5-1; and
 - (B) that, under certain conditions, she may relinquish a child who is, or who appears to be, not more than thirty (30) days of age:
 - (i) to an emergency medical services provider (as defined in IC 16-41-10-1); or
 - (ii) in a newborn safety device (described in IC 31-34-2.5-1) at a participating fire department or other site that is staffed by an emergency medical services provider.
- (b) This subsection applies to a pregnant woman whose unborn child has been diagnosed with a lethal fetal anomaly. The requirements of this subsection are in addition to the other requirements of this section. At least eighteen (18) hours before an abortion is performed on the pregnant woman, the physician who will perform the abortion shall:
 - (1) orally and in person, inform the pregnant woman of the availability of perinatal hospice services; and
 - (2) provide the pregnant woman copies of the perinatal hospice brochure developed by the state department under IC 16-25-4.5-4 and the list of perinatal hospice providers and programs



developed under IC 16-25-4.5-5, by printing the perinatal hospice brochure and list of perinatal hospice providers from the state department's Internet web site.

(c) If a pregnant woman described in subsection (b) chooses to have an abortion rather than continuing the pregnancy in perinatal hospice care, the pregnant woman shall certify in writing, on a form developed by the state department under IC 16-25-4.5-6, at least eighteen (18) hours before the abortion is performed, that the pregnant woman has been provided the information described in subsection (b) in the manner required by subsection (b)."

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Page 7, delete line 2.
   Page 7, line 3, delete "(10)" and insert "(9)".
   Page 7, line 4, delete "(11)" and insert "(10)".
   Page 7, line 5, delete "(12)" and insert "(11)".
   Page 7, line 6, delete "(13)" and insert "(12)".
   Page 7, line 7, delete "(14)" and insert "(13)".
   Page 7, line 8, delete "(15)" and insert "(14)".
   Page 7, line 9, delete "(16)" and insert "(15)".
   Page 7, line 10, delete "(17)" and insert "(16)".
   Page 7, line 11, delete "(18)" and insert "(17)".
   Page 7, line 12, delete "(19)" and insert "(18)".
   Page 7, line 13, delete "(20)" and insert "(19)".
   Page 7, line 14, delete "(21)" and insert "(20)".
   Page 7, line 16, delete "(22)" and insert "(21)".
   Page 7, line 18, delete "(23)" and insert "(22)".
   Page 7, line 20, delete "(24)" and insert "(23)".
   Page 7, line 21, delete "(25)" and insert "(24)".
   Page 7, line 23, delete "(26)" and insert "(25)".
   Page 7, line 24, delete "(27)" and insert "(26)".
   Page 7, line 28, delete "involving" and insert "in which the person
treated".
   Page 7, delete line 41.
   Page 7, line 42, delete "(3)" and insert "(2)".
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Page 8, line 1, delete "(4)" and insert "(3)".

Page 8, line 2, delete "(5)" and insert "(4)".

Page 8, line 3, delete "(6)" and insert "(5)".

Page 8, line 4, delete "(7)" and insert "(6)".

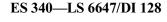
Page 8, delete lines 5 through 9.

Page 8, line 10, delete "(9)" and insert "(7)".

Page 8, line 15, delete "(10)" and insert "(8)".

Page 8, line 18, delete "(11)" and insert "(9)".

Page 8, line 21, delete "(12)" and insert "(10)".





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Page 8, line 23, delete "(13)" and insert "(11)".
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Page 8, line 24, delete "(14)" and insert "(12)".

Page 8, line 26, delete "(15)" and insert "(13)".

Page 8, line 26, after "for" insert "complication treatment".

Page 8, line 28, delete "(16)" and insert "(14)".

Page 8, line 28, delete "office".

Page 8, line 30, delete "(17)" and insert "(15)".

Page 8, line 31, delete "(18)" and insert "(16)".

Page 8, line 33, delete "(19)" and insert "(17)".

Page 9, line 10, after "(j)" insert "This subsection applies after August 31, 2019.".

Page 9, between lines 29 and 30, begin a new line block indented and insert:

"(2) Whether a waiver of consent or notification under section 4 of this chapter was obtained.".

Page 9, line 30, strike "(2)" and insert "(3)".

Page 9, line 33, strike "(3)" and insert "(4)".

Page 9, line 38, delete "(4)" and insert "(5)".

Page 9, line 40, strike "(5)" and insert "(6)".

Page 9, line 42, delete "(6)" and insert "(7)".

Page 10, line 1, delete "(7)" and insert "(8)".

Page 10, line 2, delete "(8)" and insert "(9)".

Page 10, line 3, delete "(9)" and insert "(10)".

Page 10, line 4, delete "(10)" and insert "(11)".

Page 10, line 5, delete "(11)" and insert "(12)".

Page 10, line 6, delete "(12)" and insert "(13)".

Page 10, line 7, delete "(13)" and insert "(14)".

Page 10, between lines 8 and 9, begin a new line block indented and insert:

"(15) The number of the patient's previous induced terminations."

Page 10, line 9, delete "(14)" and insert "(16)".

Page 10, line 10, delete "(15)" and insert "(17)".

Page 10, line 12, delete "(16)" and insert "(18)".

Page 10, line 18, delete "(17)" and insert "(19)".

Page 10, line 21, delete "fetus." and insert "fetus (in weeks).".

Page 10, line 33, delete "(18)" and insert "(20)".

Page 11, line 3, delete "(19)" and insert "(21)".

Page 11, between lines 5 and 6, begin a new line block indented and insert:

"(22) For a nonsurgical abortion, that the manufacturer's instructions were provided to the patient and that the patient



signed the patient agreement.".

Page 11, line 6, delete "(20)" and insert "(23)".

Page 11, line 8, delete "(21)" and insert "(24)".

Page 11, line 10, delete "(22)" and insert "(25)".

Page 11, line 12, delete "(23)" and insert "(26)".

Page 11, line 13, delete "(24)" and insert "(27)".

Page 11, line 15, delete "(25)" and insert "(28)".

Page 11, line 18, delete "(26)" and insert "(29)".

Renumber all SECTIONS consecutively.

and when so amended that said bill do pass.

(Reference is to SB 340 as introduced.)

BRAY, Chairperson

Committee Vote: Yeas 6, Nays 1.

SENATE MOTION

Madam President: I move that Senate Bill 340 be amended to read as follows:

Page 12, delete lines 39 through 41, begin a new line block indented and insert:

"(9) Whether the complication was previously managed by the abortion provider or the abortion provider's required back-up physician."

Page 13, delete lines 5 through 7.

Page 13, line 8, delete "(14)" and insert "(13)".

Page 13, line 10, delete "(15)" and insert "(14)".

Page 13, line 11, delete "(16)" and insert "(15)".

Page 13, line 13, delete "(17)" and insert "(16)".

(Reference is to SB 340 as printed January 26, 2018.)

BROWN L



COMMITTEE REPORT

Mr. Speaker: Your Committee on Public Policy, to which was referred Senate Bill 340, has had the same under consideration and begs leave to report the same back to the House with the recommendation that said bill be amended as follows:

Page 3, line 35, delete "this section" and insert "subsection (c)(1), (c)(2), or (c)(3) applies, respectively, to every abortion clinic of the type described in subsection (c)(1), (c)(2), or (c)(3),".

Page 3, delete line 36.

Page 3, line 37, delete "under this article,".

Page 5, line 5, delete "due" and insert "as a direct result of patient health and safety concerns.".

Page 5, delete line 6.

Page 6, line 5, delete "The department shall".

Page 6, delete lines 6 through 9.

Page 6, line 10, delete "the form and the form shall be retained in the patient's file.".

Page 7, delete lines 7 through 42, begin a new paragraph and insert: "SECTION 8. IC 16-34-2-1.5, AS AMENDED BY P.L.156-2017, SECTION 14, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2018]: Sec. 1.5. (a) The state department shall develop an informed consent brochure and post the informed consent brochure on the state department's Internet web site.

- (b) The state department shall develop an informed consent brochure that includes the following:
 - (1) Objective scientific information concerning the probable anatomical and physiological characteristics of a fetus every two
 - (2) weeks of gestational age, including the following:
 - (A) Realistic pictures in color for each age of the fetus, including the dimensions of the fetus.
 - (B) Whether there is any possibility of the fetus surviving outside the womb.
 - (2) Objective scientific information concerning the medical risks associated with each abortion procedure or the use of an abortion inducing drug, including the following:
 - (A) The risks of infection and hemorrhaging.
 - (B) The potential danger:
 - (i) to a subsequent pregnancy; or
 - (ii) of infertility.
 - (3) Information concerning the medical risks associated with carrying the child to term.



- (4) Information that medical assistance benefits may be available for prenatal care, childbirth, and neonatal care.
- (5) Information that the biological father is liable for assistance in support of the child, regardless of whether the biological father has offered to pay for an abortion.
- (6) Information regarding telephone 211 dialing code services for accessing human services as described in IC 8-1-19.5, and the types of services that are available through this service.
- (7) Information concerning Indiana's safe haven law under IC 31-34-2.5-1.
- (8) Information that, under certain conditions, a pregnant woman may relinquish a child who is, or who appears to be, not more than thirty (30) days of age:
 - (A) to an emergency medical services provider (as defined in IC 16-41-10-1); or
 - (B) in a newborn safety device (described in IC 31-34-2.5-1) at a participating fire department or other site that is staffed by an emergency medical services provider.
- (c) In complying with subsection (b)(6), the state department shall consult with the recognized 211 service providers and the Indiana housing and community development authority as required by IC 8-1-19.5-9.
- (d) In the development of the informed consent brochure described in this section, the state department shall use information and pictures that are available at no cost or nominal cost to the state department.
- (e) The informed consent brochure must include the requirements specified in this chapter.".

Delete pages 8 through 10.

Page 11, delete lines 1 through 15.

Page 14, line 8, delete "or notification".

Page 14, between lines 9 and 10, begin a new line block indented and insert:

"(3) Whether a waiver of notification under section 4 of this chapter was obtained.".

Page 14, line 10, delete "(3)" and insert "(4)".

Page 14, line 13, delete "(4)" and insert "(5)".

Page 14, line 18, delete "(5)" and insert "(6)".

Page 14, line 20, delete "(6)" and insert "(7)".

Page 14, line 22, delete "(7)" and insert "(8)".

Page 14, line 23, delete "(8)" and insert "(9)".

Page 14, line 24, delete "(9)" and insert "(10)".



- Page 14, line 25, delete "(10)" and insert "(11)".
- Page 14, line 26, delete "(11)" and insert "(12)".
- Page 14, line 27, delete "(12)" and insert "(13)".
- Page 14, line 28, delete "(13)" and insert "(14)".
- Page 14, line 29, delete "(14)" and insert "(15)".
- Page 14, line 31, delete "(15)" and insert "(16)".
- Page 14, line 33, delete "(16)" and insert "(17)".
- Page 14, line 34, delete "(17)" and insert "(18)".
- Page 14, line 36, delete "(18)" and insert "(19)".
- Page 14, line 42, delete "(19)" and insert "(20)".
- Page 15, line 15, delete "(20)" and insert "(21)".
- Page 15, line 27, delete "(21)" and insert "(22)".
- Page 15, line 30, delete "(22)" and insert "(23)".
- Page 15, line 33, delete "(23)" and insert "(24)".
- Page 15, line 35, delete "(24)" and insert "(25)".
- Page 15, line 37, delete "(25)" and insert "(26)".
- Page 15, line 39, delete "(26)" and insert "(27)".
- Page 15, line 40, delete "(27)" and insert "(28)".
- Page 15, line 42, delete "(28)" and insert "(29)".
- Page 16, line 3, delete "(29)" and insert "(30)".

Page 16, between lines 35 and 36, begin a new paragraph and insert: "SECTION 11. IC 31-34-2.5-1, AS AMENDED BY P.L.186-2017, SECTION 1, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2018]: Sec. 1. (a) An emergency medical services provider (as defined in IC 16-41-10-1) shall, without a court order, take custody of a child who is, or who appears to be, not more than thirty (30) days of age if:

- (1) the child is voluntarily left:
 - (A) with the provider by the child's parent;
 - (B) in a newborn safety device that:
 - (i) has been approved by a hospital licensed under IC 16-21;
 - (ii) is physically located inside a hospital that is staffed continuously on a twenty-four (24) hour basis every day to provide care to patients in an emergency; and
 - (iii) is located in an area that is conspicuous and visible to hospital staff; or
 - (C) in a newborn safety device that was installed on or before January 1, 2017, and is located at a site that is staffed by an emergency medical services provider (as defined in IC 16-41-10-1); or
 - (D) in a newborn safety device that:
 - (i) is located at a fire department, including a volunteer



fire department, that is staffed by an emergency medical services provider (as defined in IC 16-41-10-1) on a twenty-four (24) hour seven (7) day a week basis;

- (ii) is located in an area that is conspicuous and visible to staff; and
- (iii) includes an adequate dual alarm system connected to the site that is tested at least one (1) time per month to ensure the alarm system is in working order; and
- (2) the parent does not express an intent to return for the child.
- (b) An emergency medical services provider who takes custody of a child under this section shall perform any act necessary to protect the child's physical health or safety.
 - (c) Any person who in good faith voluntarily leaves a child:
 - (1) with an emergency medical services provider; or
- (2) in a newborn safety device described in subsection (a)(1)(B); is not obligated to disclose the parent's name or the person's name.
- (d) A hospital that approves the operation of a newborn safety device that meets the requirements set forth in subsection (a)(1)(B) is immune from civil liability for an act or omission relating to the operation of the newborn safety device unless the act or omission constitutes gross negligence or willful or wanton misconduct.
- (e) A newborn safety device described in subsection (a)(1)(C) may continue to operate without meeting the conditions set forth in subsection (a)(1)(B).
- (f) A fire department, including a volunteer fire department, that meets the requirements set forth in subsection (a)(1)(D) is immune from civil liability for an act or omission relating to the operation of the newborn safety device unless the act or omission constitutes gross negligence or willful or wanton misconduct.

SECTION 12. IC 34-30-2-134.5, AS ADDED BY P.L.263-2017, SECTION 2, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2018]: Sec. 134.5. IC 31-34-2.5-1 (Concerning a hospital **or fire department** operating a newborn safety device).

SECTION 13. IC 35-46-1-4, AS AMENDED BY P.L.252-2017, SECTION 17, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE JULY 1, 2018]: Sec. 4. (a) A person having the care of a dependent, whether assumed voluntarily or because of a legal obligation, who knowingly or intentionally:

- (1) places the dependent in a situation that endangers the dependent's life or health;
- (2) abandons or cruelly confines the dependent;
- (3) deprives the dependent of necessary support; or



- (4) deprives the dependent of education as required by law; commits neglect of a dependent, a Level 6 felony.
 - (b) However, the offense is:
 - (1) a Level 5 felony if it is committed under subsection (a)(1), (a)(2), or (a)(3) and:
 - (A) results in bodily injury; or
 - (B) is:
 - (i) committed in a location where a person is violating IC 35-48-4-1 (dealing in cocaine or a narcotic drug), IC 35-48-4-1.1 (dealing in methamphetamine), or IC 35-48-4-1.2 (manufacturing methamphetamine); or
 - (ii) the result of a violation of IC 35-48-4-1 (dealing in cocaine or a narcotic drug), IC 35-48-4-1.1 (dealing in methamphetamine), or IC 35-48-4-1.2 (manufacturing methamphetamine);
 - (2) a Level 3 felony if it is committed under subsection (a)(1), (a)(2), or (a)(3) and results in serious bodily injury;
 - (3) a Level 1 felony if it is committed under subsection (a)(1), (a)(2), or (a)(3) by a person at least eighteen (18) years of age and results in the death of a dependent who is less than fourteen (14) years of age or in the death of a dependent of any age who has a mental or physical disability; and
 - (4) a Level 5 felony if it is committed under subsection (a)(2) and consists of cruel confinement or abandonment that:
 - (A) deprives a dependent of necessary food, water, or sanitary facilities;
 - (B) consists of confinement in an area not intended for human habitation; or
 - (C) involves the unlawful use of handcuffs, a rope, a cord, tape, or a similar device to physically restrain a dependent.
- (c) It is a defense to a prosecution based on an alleged act under this section that:
 - (1) the accused person left a dependent child who was, at the time the alleged act occurred, not more than thirty (30) days of age:
 - (A) in a newborn safety device described in IC 31-34-2.5-1(a)(1)(B), or IC 31-34-2.5-1(a)(1)(C), or IC 31-34-2.5-1(a)(1)(D); or
 - (B) with a person who is an emergency medical services provider (as defined in IC 16-41-10-1) who took custody of the child under IC 31-34-2.5;

when the prosecution is based solely on the alleged act of leaving the child in the newborn safety device or with the emergency



- medical services provider and the alleged act did not result in bodily injury or serious bodily injury to the child; or
- (2) the accused person, in the legitimate practice of the accused person's religious belief, provided treatment by spiritual means through prayer, in lieu of medical care, to the accused person's dependent.
- (d) Except for property transferred or received:
 - (1) under a court order made in connection with a proceeding under IC 31-15, IC 31-16, IC 31-17, or IC 31-35 (or IC 31-1-11.5 or IC 31-6-5 before their repeal); or
 - (2) under section 9(d) of this chapter;

a person who transfers or receives any property in consideration for the termination of the care, custody, or control of a person's dependent child commits child selling, a Level 6 felony.".

Renumber all SECTIONS consecutively.

and when so amended that said bill do pass.

(Reference is to SB 340 as reprinted January 30, 2018.)

SMALTZ

Committee Vote: yeas 9, nays 4.

