

**SUBSTITUTE FOR  
SENATE BILL NO. 1082**

A bill to regulate the collection, processing, and selling of reproductive health data; to regulate the disclosure of reproductive health data; to require individual consent to collect, process, and sell reproductive health data; to prohibit the use of geofences around facilities that provide reproductive health services; to provide remedies and prescribe civil sanctions; and to provide for the powers and duties of certain state governmental officers and entities.

**THE PEOPLE OF THE STATE OF MICHIGAN ENACT:**

1       Sec. 1. This act may be cited as the "reproductive health data  
2 privacy act".

3       Sec. 3. As used in this act:

4       (a) "Affiliate" means a legal entity that is controlled by or

1 is under common control with another legal entity. For the purposes  
2 of this subdivision, an entity is controlled by another entity or  
3 under common control if the controlling entity has any of the  
4 following:

5 (i) The majority voting or ownership interest of the  
6 outstanding shares of any class of voting security of the  
7 controlled entity.

8 (ii) Control in any manner over the election of a majority of  
9 the directors or of individuals exercising similar functions of the  
10 controlled entity.

11 (iii) The power to exercise controlling influence over the  
12 management of the controlled entity.

13 (b) "Collect" means to buy, rent, gather, obtain, receive, or  
14 access any reproductive health data about an individual in any  
15 manner, including, but not limited to, by receiving data from the  
16 individual, actively or passively, or by observing or tracking the  
17 individual's online activity.

18 (c) "Consent" means a clear affirmative act that signifies an  
19 individual's freely given, specific, informed, opt-in, voluntary,  
20 and unambiguous agreement, that may be given electronically, and is  
21 provided in response to a specific request from a regulated entity  
22 or a service provider. Consent does not include an agreement  
23 obtained by any of the following:

24 (i) A general or broad terms-of-use agreement or a similar  
25 document that contains descriptions of reproductive health data  
26 processing along with other unrelated information.

27 (ii) An individual hovering over, muting, pausing, or closing a  
28 given piece of consent.

29 (iii) Through the use of a deceptive design.

1 (d) "Deceptive design" means an interface design or choice  
2 architecture to obtain required consent that has been designed or  
3 manipulated with the substantial effect of subverting or impairing  
4 user autonomy, decision making, or choice, or unfairly,  
5 fraudulently, or deceptively manipulating or coercing an individual  
6 into providing consent.

7 (e) "Geofence" means technology that uses global positioning  
8 coordinates, cell tower connectivity, cellular data, radio  
9 frequency identification, Wi-Fi data, or any other form of spatial  
10 or location detection to establish a virtual boundary around a  
11 specific physical location, or to locate an individual within a  
12 virtual boundary, where the virtual boundary is not more than 1,850  
13 feet from the perimeter of the physical location.

14 (f) "Mobile application" means a software program that runs on  
15 the operating system of a cellular telephone, a tablet computer, or  
16 a similar portable computing device that transmits data over a  
17 wireless connection and includes a service or application offered  
18 via a connected device.

19 (g) "Person" means an individual or a partnership,  
20 corporation, limited liability company, association, governmental  
21 entity, or other legal entity.

22 (h) "Process" means any use of data provided under this act.

23 (i) "Publicly available information" means information that  
24 has been made lawfully available by federal, state, or municipal  
25 government records, widely distributed media, or a disclosure to  
26 the general public as required under federal, state, or local law.  
27 Publicly available information does not include any of the  
28 following:

29 (i) An obscene visual depiction as that term is defined in 18

1 USC 1460.

2 (ii) An inference made exclusively from multiple independent  
3 sources of publicly available information that reveals an  
4 individual's reproductive health data.

5 (iii) Biometric data.

6 (iv) Reproductive health data that is created through the  
7 combination of information that identifies the individual's past,  
8 present, or future reproductive health status with publicly  
9 available information.

10 (v) Genetic data, unless the data is otherwise made publicly  
11 available by the individual to whom the information pertains.

12 (vi) Information made available by an individual on a website  
13 or online service made available to all members of the public, for  
14 free or for a fee, where the consumer has maintained a reasonable  
15 expectation of privacy by restricting the information to a specific  
16 audience.

17 (vii) Intimate images, authentic or computer generated, known  
18 to be nonconsensual.

19 (j) "Regulated entity" means a public, private, operated for  
20 profit, or not operated for profit business or organization that  
21 provides reproductive health care or services and collects  
22 reproductive health data from an individual. Regulated entity  
23 includes a business or organization that licenses or certifies  
24 other persons to provide reproductive health care or services.

25 (k) "Reproductive health data" means information that is  
26 linked or reasonably linkable to an individual and that identifies  
27 the individual's past, present, or future reproductive health  
28 status. Reproductive health data does not include aggregated and  
29 de-identified data or information that is used to engage in public

1 or peer-reviewed scientific, historical, or statistical research in  
2 the public interest, including information described under 1967 PA  
3 270, MCL 331.531 to 331.534, that adheres to all other applicable  
4 ethics and privacy laws and is approved, monitored, or governed by  
5 an institutional review board, human subjects research ethics  
6 board, or a similar independent oversight entity that determines  
7 that the regulated entity has implemented reasonable safeguards to  
8 reduce privacy risks associated with research, including risks  
9 associated with reidentification.

10 (l) "Reproductive health services" means health care services  
11 or products that support an individual's reproductive system,  
12 pregnancy status, or sexual well-being, including, but not limited  
13 to, any of the following:

14 (i) Individual health conditions, status, diseases, or  
15 diagnoses.

16 (ii) Social, psychological, behavioral, and medical  
17 interventions.

18 (iii) Health-related surgeries or procedures, including, but not  
19 limited to, abortions.

20 (iv) Bodily functions, vital signs, symptoms, or measurements  
21 of the information described in this subdivision.

22 (v) Diagnoses or diagnostic testing, treatment, or medication.

23 (vi) Medical or nonmedical services related to and provided in  
24 conjunction with an abortion, including, but not limited to,  
25 associated diagnostics, counseling, supplies, and follow-up  
26 services.

27 (m) "Reproductive health status" means any of the following as  
28 it relates to an individual's reproductive health, menstrual cycle,  
29 fertility, pregnancy, pregnancy outcome, plans to conceive, or type

1 of sexual activity:

2 (i) Individual health conditions, treatment, or diseases.

3 (ii) Diagnoses done by a medical professional.

4 (iii) Social, psychological, behavioral, and medical  
5 interventions.

6 (iv) Health-related surgeries or procedures.

7 (v) Use or purchase of medications.

8 (vi) Bodily functions, vital signs, symptoms, or measurements  
9 of the information described in this subdivision.

10 (vii) Diagnoses or diagnostic testing, treatment, or medication  
11 done or prescribed by a medical professional.

12 (viii) Data concerning medical or nonmedical services related to  
13 and provided in conjunction with an abortion, including, but not  
14 limited to, associated diagnostics, counseling, supplies, and  
15 follow-up services done by a medical professional.

16 (ix) Biometric data. As used in this subparagraph, "biometric  
17 data" means data generated by automatic measurements of an  
18 individual's biological characteristics, including, but not limited  
19 to, a fingerprint, a voiceprint, an eye retina, an iris, or any  
20 other biological pattern or characteristic used to identify a  
21 specific individual. Biometric data does not include any of the  
22 following:

23 (A) A physical or digital photograph.

24 (B) A video or audio recording.

25 (C) Any data generated from a physical or digital photograph  
26 or a video or audio recording, unless the data is generated to  
27 identify a specific individual.

28 (x) Genetic data.

29 (xi) Precise location information that could reasonably

1 indicate an individual's attempt to acquire or receive reproductive  
2 health services or supplies.

3 (xii) Data that identifies an individual seeking reproductive  
4 health services or supplies.

5 (xiii) Any information that a regulated entity, or a regulated  
6 entity's respective service provider, processes to associate or  
7 identify an individual with the data described in subparagraphs (i)  
8 to (xi) that is derived or extrapolated from other information, such  
9 as proxy, derivative, inferred, or emergent data, by any means,  
10 including algorithms and machine learning.

11 (n) "Sell" or "sale" means the exchange of reproductive health  
12 data for monetary or other valuable consideration by a regulated  
13 entity to a third party. Sell or sale does not include any of the  
14 following:

15 (i) The exchange of reproductive health data for monetary or  
16 other valuable consideration to a third party as an asset that is  
17 part of a merger, acquisition, bankruptcy, or other transaction, or  
18 a proposed merger, acquisition, bankruptcy, or other transaction,  
19 in which the third party assumes control of all or part of the  
20 regulated entity's assets, only if the regulated entity, in a  
21 reasonable time before the exchange, provides the affected  
22 individual with both of the following:

23 (A) A notice describing the transfer, including the name of  
24 the entity receiving the individual's reproductive health data and  
25 the applicable privacy policies of the entity.

26 (B) A reasonable opportunity to withdraw previously provided  
27 consent related to the individual's reproductive health data and  
28 request the deletion of the individual's reproductive health data.

29 (ii) The disclosure of reproductive health data to a service

1 provider that processes reproductive health data on behalf of a  
2 regulated entity.

3 (iii) The disclosure or transfer of reproductive health data to  
4 an affiliate of a regulated entity.

5 (iv) The disclosure of publicly available information.

6 (o) "Service provider" means a person that collects,  
7 processes, retains, transfers, or sells reproductive health data on  
8 behalf of, and at the direction of, a regulated entity.

9 (p) "Third party" means a person other than a party to a  
10 transaction or a party's representative for the purposes specified  
11 under this act.

12 (q) "Trade secrets" means that term as defined in the uniform  
13 trade secrets act, 1998 PA 448, MCL 445.1902.

14 Sec. 5. (1) A regulated entity shall not collect or process  
15 reproductive health data unless the regulated entity does all of  
16 the following:

17 (a) Provides the individual whose reproductive health data is  
18 being collected with a copy of the regulated entity's privacy  
19 policy.

20 (b) Obtains consent from the individual to whom the  
21 reproductive health data pertains, or the individual's authorized  
22 representative.

23 (c) Collects or processes the reproductive health data only  
24 for 1 or more purposes described under subsection (3).

25 (2) This section does not apply to reproductive health data  
26 that is considered protected health information or to information  
27 originating from, and intermingled to be indistinguishable with,  
28 protected health information that is maintained by a covered entity  
29 or business associate as those terms are defined by the health



1 insurance portability and accountability act of 1996, Public Law  
2 104-191, and the regulations promulgated under that act, 45 CFR  
3 parts 160 and 164. As used in this subsection, "protected health  
4 information" means that term as defined in the health insurance  
5 portability and accountability act of 1996, Public Law 104-191.

6 (3) A regulated entity may process reproductive health data  
7 only for the following purposes:

8 (a) As strictly necessary to provide a product, service, or  
9 service feature to the individual to whom the reproductive health  
10 data pertains when requested by that individual.

11 (b) To initiate, manage, execute, or complete a financial or  
12 commercial transaction or to fulfill an order for a specific  
13 product or service requested by an individual to whom the  
14 reproductive health data pertains, including, but not limited to,  
15 associated routine administrative, operational, and account  
16 servicing activity such as billing, shipping, storage, and  
17 accounting.

18 (c) To comply with an obligation under a law of this state or  
19 federal law.

20 (d) To protect public safety or public health.

21 (e) To prevent, detect, protect against, or respond to a  
22 security incident, identity theft, fraud, harassment, malicious or  
23 deceptive activities, or activities that are illegal under the laws  
24 of this state.

25 (f) To preserve the integrity or security of systems.

26 (g) To investigate, report, or prosecute persons responsible  
27 for activities that are illegal under the laws of this state.

28 (4) A regulated entity that collects or processes reproductive  
29 health data shall not do any of the following:

1 (a) Collect more precise reproductive health data than is  
2 necessary to perform a purpose described in subsection (3).

3 (b) Retain reproductive health data for longer than is  
4 necessary to perform a purpose described in subsection (3).

5 (c) Derive or infer from reproductive health data any  
6 information that is not necessary to perform a purpose described in  
7 subsection (3).

8 (d) Disclose, cause to disclose, assist with the disclosure  
9 of, or facilitate the disclosure of an individual's reproductive  
10 health data to a third party, unless the disclosure is either of  
11 the following:

12 (i) Necessary to perform a purpose described under subsection  
13 (3).

14 (ii) Subject to the requirements of section 6, disclosed to a  
15 service provider.

16 (5) A regulated entity that collects or processes reproductive  
17 health data shall provide a clear and conspicuous link, that is  
18 secure and reliable, on the regulated entity's internet homepage or  
19 mobile application that enables an individual, or a person  
20 authorized by the individual, to request access to and deletion of  
21 the individual's reproductive health data. Access provided under  
22 this subsection must not require the disclosure of trade secrets.

23 (6) A regulated entity shall respond to a request under this  
24 section without undue delay, but not later than 45 days after the  
25 receipt of the individual's request. The response period may be  
26 extended by an additional 45 days if reasonably necessary,  
27 considering the complexity and volume of the individual's requests.  
28 The individual must be informed of an extension and the reason for  
29 the extension within the initial 45-day response period.

1           Sec. 6. (1) A service provider shall process reproductive  
2 health data only under a contract with a regulated entity that sets  
3 forth the processing instructions and limits the actions that the  
4 service provider may take with respect to the reproductive health  
5 data that the service provider processes on behalf of the regulated  
6 entity.

7           (2) A service provider shall process reproductive health data  
8 in a manner that is consistent with the instructions set forth in  
9 the contract under subsection (1).

10          (3) If a service provider knowingly fails to comply with the  
11 instructions in the contract under subsection (1) or processes  
12 reproductive health data in a manner inconsistent with the contract  
13 under subsection (1), the service provider is considered a  
14 regulated entity regarding that reproductive health data and is  
15 subject to all of the requirements of this act.

16          (4) A service provider shall assist the regulated entity by  
17 appropriate technical or organizational measures, if possible, in  
18 fulfilling the regulated entity's obligations under this act.

19          Sec. 7. A regulated entity or service provider shall not  
20 disclose an individual's reproductive health data to a federal,  
21 state, or local governmental agency or official unless 1 or more of  
22 the following applies:

23          (a) The governmental agency or official serves the regulated  
24 entity or service provider with a valid warrant or establishes the  
25 existence of exigent circumstances that make it impracticable to  
26 obtain a warrant, except as prohibited by the laws of this state.

27          (b) Disclosure is mandated under the laws of this state or  
28 federal law.

29          (c) Disclosure is requested by the individual to whom the

1 reproductive health data pertains.

2 (d) Disclosure is ordered by a federal court.

3 Sec. 9. (1) Beginning on June 30, 2027, a regulated entity or  
4 service provider shall not sell or offer to sell reproductive  
5 health data unless the regulated entity or service provider obtains  
6 valid consent in accordance with subsection (4) from the individual  
7 to whom the reproductive health data pertains before selling or  
8 offering to sell the reproductive health data.

9 (2) A regulated entity or service provider shall not sell or  
10 offer to sell reproductive health data in a manner that is  
11 inconsistent with valid consent obtained under this section.

12 (3) Valid consent under this section is separate and distinct  
13 from consent obtained under section 5.

14 (4) To be valid, consent under this section must be in  
15 writing, in plain language, and contain all of the following:

16 (a) The specific reproductive health data concerning the  
17 individual that the regulated entity or service provider intends to  
18 sell.

19 (b) The name and contact information of the regulated entity  
20 or service provider collecting and selling the reproductive health  
21 data described in subdivision (a).

22 (c) The name and contact information of the person purchasing  
23 the reproductive health data described in subdivision (a).

24 (d) A description of the purpose for the sale, including how  
25 the reproductive health data will be gathered by the regulated  
26 entity or service provider and how the reproductive health data  
27 will be used by the person purchasing the reproductive health data.

28 (e) A statement that the provision of goods and services is  
29 not conditioned on the individual signing the consent.

1 (f) A statement that the individual has a right to revoke the  
2 individual's consent at any time, and a description of how to  
3 submit a revocation of the consent.

4 (g) A statement that the reproductive health data sold in  
5 accordance with valid consent may be subject to redisclosure by the  
6 person purchasing the reproductive health data and may no longer be  
7 protected under this section.

8 (h) The signature of the individual providing consent and the  
9 date on which the consent was signed by the individual.

10 (i) An expiration date for the consent, which must expire  
11 within 1 year after the individual's signature.

12 (5) Consent is not valid if it has any of the following  
13 defects:

14 (a) The expiration date has passed.

15 (b) The consent does not contain all of the information  
16 required under subsection (4).

17 (c) The consent has been revoked by the individual.

18 (d) The consent has been combined with other documents to  
19 create a compound authorization.

20 (e) The provision of goods or services is conditioned on the  
21 individual signing the consent document.

22 (6) A copy of the valid consent must be provided to the  
23 individual by the regulated entity or service provider selling or  
24 offering to sell the reproductive health data.

25 (7) The regulated entity or service provider selling or  
26 offering to sell the reproductive health data and the purchaser of  
27 the reproductive health data shall retain a copy of the valid  
28 consent for not less than 6 years after the date that the consent  
29 is signed by the individual or the date when the consent was last

1 in effect, whichever is later.

2 (8) A regulated entity or service provider that sells  
3 reproductive health data shall provide a clear and conspicuous link  
4 on the regulated entity or service provider's internet homepage or  
5 mobile application that enables an individual, or a person  
6 authorized by the individual, to revoke the individual's consent to  
7 sell reproductive health data at any time.

8 (9) A regulated entity or service provider selling an  
9 individual's reproductive health data and the purchaser of the  
10 reproductive health data shall enter into a written agreement  
11 governing the purchaser's processing of the individual's  
12 reproductive health data. The written agreement must do all of the  
13 following:

14 (a) Legally bind the purchaser and the regulated entity or  
15 service provider selling the reproductive health data.

16 (b) Clearly set forth the nature and purpose of the sale, the  
17 type of reproductive health data subject to the sale, the duration  
18 of processing, and the rights and obligations of both parties.

19 (c) Require the purchaser to adhere to the instructions of the  
20 regulated entity or service provider.

21 (d) Set out the extent to which the purchaser may process the  
22 reproductive health data.

23 (e) Require the purchaser to process the reproductive health  
24 data that the purchaser receives from the regulated entity or  
25 service provider only to the extent provided for under subdivision  
26 (d).

27 (f) Require the purchaser to delete or return all reproductive  
28 health data to the regulated entity or service provider at the end  
29 of the provision of services or on revocation of consent by the

1 individual, unless retention of the reproductive health data is  
2 required by law.

3       Sec. 11. A person shall not implement a geofence around an  
4 entity that provides in-person reproductive health services if the  
5 geofence is used to do any of the following:

6           (a) Identify or track individuals for the purpose of  
7 determining whether the individual is seeking reproductive health  
8 services.

9           (b) Collect reproductive health data from individuals.

10          (c) Send notifications, messages, or advertisements to  
11 individuals related to the individual's reproductive health data or  
12 reproductive health services.

13       Sec. 13. (1) The attorney general may bring an action to  
14 enjoin any person from violating this act. On proper showing, a  
15 court may grant a permanent or temporary injunction, restraining  
16 order, writ of mandamus, or any other order or judgment necessary  
17 to enjoin a person from violating this act. For any action in which  
18 the attorney general prevails, the attorney general may recover the  
19 costs of the action, including reasonable attorney fees.

20          (2) The attorney general or an individual who alleges a loss  
21 as a result of a violation of this act may bring a civil action  
22 against the person that committed the violation to recover any of  
23 the following:

24           (a) Damages in an amount of not less than \$100.00 and not more  
25 than \$750.00 per incident or actual damages, whichever is greater.

26           (b) Injunctive or declaratory relief.

27           (c) Any other appropriate relief.

28          (3) The court may consider any relevant circumstances in  
29 determining the amount of damages, including, but not limited to,

1 all of the following:

2 (a) The nature and seriousness of the misconduct.

3 (b) The number of violations.

4 (c) The persistence of the misconduct.

5 (d) The length of time over which the misconduct occurred.

6 (e) The willfulness of the defendant's misconduct.

7 (f) The defendant's assets, liabilities, and net worth.

8 (4) This act does not serve as a basis for a private right of  
9 action under any other law. This subsection does not deprive or  
10 relieve a person from any rights, duties, or obligations imposed  
11 under other laws of this state or federal law.

12 Sec. 15. The attorney general may promulgate rules to  
13 implement this act under the administrative procedures act of 1969,  
14 1969 PA 306, MCL 24.201 to 24.328.

15 Enacting section 1. This act takes effect 2 years after the  
16 date it is enacted into law.