



## 103RD GENERAL ASSEMBLY

### State of Illinois

2023 and 2024

HB3585

Introduced 2/17/2023, by Rep. Tom Weber

#### SYNOPSIS AS INTRODUCED:

New Act  
215 ILCS 5/356z.4  
225 ILCS 85/45 new  
775 ILCS 55/1-40 new

Creates the Long-Acting Reversible Contraception Information Act. Provides that the Department of Public Health shall create and allocate funding for an online learning module to promote postpartum and postabortion long-acting reversible contraception insertion. Provides that long-acting reversible contraception services and information may be provided by physicians to any minor over the age of 12 who meets specified qualifications. Provides that the Department shall provide printed materials, guidance, and information on how to obtain low-cost and no-cost contraceptives. Provides that the Department shall develop a long-acting reversible contraception promotion plan intended to reduce cases of neonatal abstinence syndrome and fetal substance exposure. Provides that the Department shall produce an annual report on the program. Provides that the Department shall adopt rules necessary to carry out the Act. Amends the Illinois Insurance Code. Provides that an individual or group policy of accident and health insurance shall also cover long-acting reversible contraception on the day of the abortion as long as the procedure is medically feasible. Amends the Pharmacy Practice Act. Provides that a pharmacist licensed under the Act who dispenses self-administered hormonal contraceptives shall provide the patient with information on the effectiveness and availability of intrauterine devices and implants. Amends the Reproductive Health Act. Provides that a health care professional shall provide information about intrauterine devices at the time that a health care professional performs an abortion.

LRB103 29391 CPF 55782 b

1 AN ACT concerning health.

2 **Be it enacted by the People of the State of Illinois,**  
3 **represented in the General Assembly:**

4 Section 1. Short title. This Act may be cited as the  
5 Long-Acting Reversible Contraception Information Act.

6 Section 5. Definitions. As used in this Act:

7 "Department" means the Department of Public Health.

8 "Health care professional" means a person who is  
9 licensed as a physician, advanced practice registered  
10 nurse, or physician assistant.

11 Section 10. Internet learning module for long-acting  
12 reversible contraception. The Department shall create and  
13 allocate funding for an online learning module, for health  
14 care professionals, to promote postpartum and postabortion  
15 long-acting reversible contraception insertion. A health care  
16 professional who participates in this module may apply the  
17 hours toward some of the continuing education requirements for  
18 the health care professionals.

19 Section 15. Long-acting reversible contraception for  
20 minors. Long-acting reversible contraception services and  
21 information may be rendered by physicians licensed in the

1 State to practice medicine in all of its branches to any minor  
2 over the age of 12:

3 (1) who is married;

4 (2) who is a parent;

5 (3) who has the consent of the minor's parents or  
6 legal guardian;

7 (4) who would face a serious health hazard if those  
8 services were not provided; or

9 (5) who is referred for such services by a physician,  
10 clergyman, or a planned parenthood agency.

11 Section 20. Guidance on low and no cost options of  
12 long-acting reversible contraception; long-acting reversible  
13 contraception promotion plan.

14 (a) The Department shall provide printed materials,  
15 guidance, and information on how to obtain low-cost and  
16 no-cost options for contraceptive promotion and long-acting  
17 reversible contraception to health care professionals.

18 (b) The Department shall develop a long-acting reversible  
19 contraception promotion plan to reduce cases of neonatal  
20 abstinence syndrome and fetal substance exposure. The  
21 Department shall produce an annual report on the program.

22 Section 25. Rulemaking. The Department shall adopt rules  
23 necessary to carry out this Act.

1           Section 30. The Illinois Insurance Code is amended by  
2 changing Section 356z.4 as follows:

3           (215 ILCS 5/356z.4)

4           Sec. 356z.4. Coverage for contraceptives.

5           (a) (1) The General Assembly hereby finds and declares all  
6 of the following:

7           (A) Illinois has a long history of expanding timely  
8 access to birth control to prevent unintended pregnancy.

9           (B) The federal Patient Protection and Affordable Care  
10 Act includes a contraceptive coverage guarantee as part of  
11 a broader requirement for health insurance to cover key  
12 preventive care services without out-of-pocket costs for  
13 patients.

14           (C) The General Assembly intends to build on existing  
15 State and federal law to promote gender equity and women's  
16 health and to ensure greater contraceptive coverage equity  
17 and timely access to all federal Food and Drug  
18 Administration approved methods of birth control for all  
19 individuals covered by an individual or group health  
20 insurance policy in Illinois.

21           (D) Medical management techniques such as denials,  
22 step therapy, or prior authorization in public and private  
23 health care coverage can impede access to the most  
24 effective contraceptive methods.

25           (2) As used in this subsection (a):

1 "Contraceptive services" includes consultations,  
2 examinations, procedures, and medical services related to the  
3 use of contraceptive methods (including natural family  
4 planning) to prevent an unintended pregnancy.

5 "Medical necessity", for the purposes of this subsection  
6 (a), includes, but is not limited to, considerations such as  
7 severity of side effects, differences in permanence and  
8 reversibility of contraceptive, and ability to adhere to the  
9 appropriate use of the item or service, as determined by the  
10 attending provider.

11 "Therapeutic equivalent version" means drugs, devices, or  
12 products that can be expected to have the same clinical effect  
13 and safety profile when administered to patients under the  
14 conditions specified in the labeling and satisfy the following  
15 general criteria:

16 (i) they are approved as safe and effective;

17 (ii) they are pharmaceutical equivalents in that they  
18 (A) contain identical amounts of the same active drug  
19 ingredient in the same dosage form and route of  
20 administration and (B) meet compendial or other applicable  
21 standards of strength, quality, purity, and identity;

22 (iii) they are bioequivalent in that (A) they do not  
23 present a known or potential bioequivalence problem and  
24 they meet an acceptable in vitro standard or (B) if they do  
25 present such a known or potential problem, they are shown  
26 to meet an appropriate bioequivalence standard;

1 (iv) they are adequately labeled; and  
2 (v) they are manufactured in compliance with Current  
3 Good Manufacturing Practice regulations.

4 (3) An individual or group policy of accident and health  
5 insurance amended, delivered, issued, or renewed in this State  
6 after the effective date of this amendatory Act of the 99th  
7 General Assembly shall provide coverage for all of the  
8 following services and contraceptive methods:

9 (A) All contraceptive drugs, devices, and other  
10 products approved by the United States Food and Drug  
11 Administration. This includes all over-the-counter  
12 contraceptive drugs, devices, and products approved by the  
13 United States Food and Drug Administration, excluding male  
14 condoms. This requirement includes long-acting reversible  
15 contraception on the day of an abortion as long as the  
16 procedure is medically feasible. The following apply:

17 (i) If the United States Food and Drug  
18 Administration has approved one or more therapeutic  
19 equivalent versions of a contraceptive drug, device,  
20 or product, a policy is not required to include all  
21 such therapeutic equivalent versions in its formulary,  
22 so long as at least one is included and covered without  
23 cost-sharing and in accordance with this Section.

24 (ii) If an individual's attending provider  
25 recommends a particular service or item approved by  
26 the United States Food and Drug Administration based

1 on a determination of medical necessity with respect  
2 to that individual, the plan or issuer must cover that  
3 service or item without cost sharing. The plan or  
4 issuer must defer to the determination of the  
5 attending provider.

6 (iii) If a drug, device, or product is not  
7 covered, plans and issuers must have an easily  
8 accessible, transparent, and sufficiently expedient  
9 process that is not unduly burdensome on the  
10 individual or a provider or other individual acting as  
11 a patient's authorized representative to ensure  
12 coverage without cost sharing.

13 (iv) This coverage must provide for the dispensing  
14 of 12 months' worth of contraception at one time.

15 (B) Voluntary sterilization procedures.

16 (C) Contraceptive services, patient education, and  
17 counseling on contraception.

18 (D) Follow-up services related to the drugs, devices,  
19 products, and procedures covered under this Section,  
20 including, but not limited to, management of side effects,  
21 counseling for continued adherence, and device insertion  
22 and removal.

23 (4) Except as otherwise provided in this subsection (a), a  
24 policy subject to this subsection (a) shall not impose a  
25 deductible, coinsurance, copayment, or any other cost-sharing  
26 requirement on the coverage provided. The provisions of this

1 paragraph do not apply to coverage of voluntary male  
2 sterilization procedures to the extent such coverage would  
3 disqualify a high-deductible health plan from eligibility for  
4 a health savings account pursuant to the federal Internal  
5 Revenue Code, 26 U.S.C. 223.

6 (5) Except as otherwise authorized under this subsection  
7 (a), a policy shall not impose any restrictions or delays on  
8 the coverage required under this subsection (a).

9 (6) If, at any time, the Secretary of the United States  
10 Department of Health and Human Services, or its successor  
11 agency, promulgates rules or regulations to be published in  
12 the Federal Register or publishes a comment in the Federal  
13 Register or issues an opinion, guidance, or other action that  
14 would require the State, pursuant to any provision of the  
15 Patient Protection and Affordable Care Act (Public Law  
16 111-148), including, but not limited to, 42 U.S.C.  
17 18031(d)(3)(B) or any successor provision, to defray the cost  
18 of any coverage outlined in this subsection (a), then this  
19 subsection (a) is inoperative with respect to all coverage  
20 outlined in this subsection (a) other than that authorized  
21 under Section 1902 of the Social Security Act, 42 U.S.C.  
22 1396a, and the State shall not assume any obligation for the  
23 cost of the coverage set forth in this subsection (a).

24 (b) This subsection (b) shall become operative if and only  
25 if subsection (a) becomes inoperative.

26 An individual or group policy of accident and health



1 insurance amended, delivered, issued, or renewed in this State  
2 after the date this subsection (b) becomes operative that  
3 provides coverage for outpatient services and outpatient  
4 prescription drugs or devices must provide coverage for the  
5 insured and any dependent of the insured covered by the policy  
6 for all outpatient contraceptive services and all outpatient  
7 contraceptive drugs and devices approved by the Food and Drug  
8 Administration. Coverage required under this Section may not  
9 impose any deductible, coinsurance, waiting period, or other  
10 cost-sharing or limitation that is greater than that required  
11 for any outpatient service or outpatient prescription drug or  
12 device otherwise covered by the policy.

13 Nothing in this subsection (b) shall be construed to  
14 require an insurance company to cover services related to  
15 permanent sterilization that requires a surgical procedure.

16 As used in this subsection (b), "outpatient contraceptive  
17 service" means consultations, examinations, procedures, and  
18 medical services, provided on an outpatient basis and related  
19 to the use of contraceptive methods (including natural family  
20 planning) to prevent an unintended pregnancy.

21 (c) (Blank).

22 (d) If a plan or issuer utilizes a network of providers,  
23 nothing in this Section shall be construed to require coverage  
24 or to prohibit the plan or issuer from imposing cost-sharing  
25 for items or services described in this Section that are  
26 provided or delivered by an out-of-network provider, unless

1 the plan or issuer does not have in its network a provider who  
2 is able to or is willing to provide the applicable items or  
3 services.

4 (Source: P.A. 100-1102, eff. 1-1-19; 101-13, eff. 6-12-19.)

5 Section 35. The Pharmacy Practice Act is amended by adding  
6 Section 45 as follows:

7 (225 ILCS 85/45 new)

8 Sec. 45. Information on intrauterine devices and implants.

9 A pharmacist licensed under this Act who dispenses  
10 self-administered hormonal contraceptives shall provide the  
11 patient with information on the effectiveness and availability  
12 of intrauterine devices and implants.

13 Section 40. The Reproductive Health Act is amended by  
14 adding Section 1-40 as follows:

15 (775 ILCS 55/1-40 new)

16 Sec. 1-40. Information provided at the time of abortion. A

17 health care professional shall provide information about  
18 intrauterine devices at the time that a health care  
19 professional performs an abortion.