



Rep. Michelle Mussman

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1 AMENDMENT TO HOUSE BILL 274

2 AMENDMENT NO. _____. Amend House Bill 274, AS AMENDED, by
3 replacing everything after the enacting clause with the
4 following:

5 "Section 5. The State Employees Group Insurance Act of 1971
6 is amended by changing Section 6.11 as follows:

7 (5 ILCS 375/6.11)

8 Sec. 6.11. Required health benefits; Illinois Insurance
9 Code requirements. The program of health benefits shall provide
10 the post-mastectomy care benefits required to be covered by a
11 policy of accident and health insurance under Section 356t of
12 the Illinois Insurance Code. The program of health benefits
13 shall provide the coverage required under Sections 356g,
14 356g.5, 356g.5-1, 356m, 356u, 356w, 356x, 356z.2, 356z.4,
15 356z.6, 356z.8, 356z.9, 356z.10, 356z.11, 356z.12, 356z.13,
16 356z.14, 356z.15, 356z.17, 356z.22, ~~and~~ 356z.25, 356z.26, and

1 356z.29 of the Illinois Insurance Code. The program of health
2 benefits must comply with Sections 155.22a, 155.37, 355b,
3 356z.19, 370c, and 370c.1 of the Illinois Insurance Code.

4 Rulemaking authority to implement Public Act 95-1045, if
5 any, is conditioned on the rules being adopted in accordance
6 with all provisions of the Illinois Administrative Procedure
7 Act and all rules and procedures of the Joint Committee on
8 Administrative Rules; any purported rule not so adopted, for
9 whatever reason, is unauthorized.

10 (Source: P.A. 99-480, eff. 9-9-15; 100-24, eff. 7-18-17;
11 100-138, eff. 8-18-17; revised 10-3-17.)

12 Section 10. The Department of Public Health Powers and
13 Duties Law of the Civil Administrative Code of Illinois is
14 amended by adding Section 2310-700 as follows:

15 (20 ILCS 2310/2310-700 new)

16 Sec. 2310-700. Contraceptive drugs and products; Director
17 standing order.

18 (a) As used in this Section:

19 "Hormonal contraceptive" means a prescribed
20 medically-acceptable oral drug, transdermal patch, or vaginal
21 ring that is approved by the United States Food and Drug
22 Administration to prevent pregnancy.

23 "Standing order" has the meaning given to that term in the
24 Pharmacy Practice Act.

1 (b) If the Director of Public Health is a physician
2 licensed to practice medicine in all its branches in Illinois,
3 the Director shall establish a standing order complete with the
4 issuance of a prescription for a hormonal contraceptive in
5 accordance with this Section. If the Director is not a
6 physician licensed to practice medicine in all its branches in
7 Illinois, then the Medical Director of the Department of Public
8 Health shall establish a standing order in accordance with this
9 Section.

10 (c) The standing order, at a minimum, shall comply with the
11 following:

12 (1) A pharmacist may dispense a 12-month supply of
13 hormonal contraceptives to a patient.

14 (2) A pharmacist shall have the patient complete the
15 self-screening risk assessment tool. The self-screening
16 risk assessment tool is to be based on the most current
17 version of the United States Medical Eligibility Criteria
18 for Contraceptive Use published by the federal Centers for
19 Disease Control and Prevention.

20 (3) Based upon the results of the self-screening risk
21 assessment and the patient assessment, the pharmacist
22 shall use his or her professional and clinical judgment as
23 to when a patient should be referred to the patient's
24 physician or another health care provider.

25 (4) The pharmacist shall provide, during the patient
26 assessment and consultation, counseling and education

1 about all methods of contraception, including methods not
2 covered under the standing order, and their proper use and
3 effectiveness.

4 (5) The patient consultation shall take place in a
5 private manner consistent with rules adopted by the
6 Department of Financial and Professional Regulation.

7 (6) The Department shall adopt rules under this Section
8 that require a pharmacist to:

9 (A) complete an educational training program
10 accredited by the Accreditation Council for Pharmacy
11 Education and approved by the Department that is
12 related to the patient self-screening risk assessment,
13 patient assessment, contraceptive counseling and
14 education, and dispensation of hormonal
15 contraceptives; and

16 (B) dispense the hormonal contraceptive to the
17 patient as soon as practicable after meeting the
18 requirements of paragraph (2).

19 (7) All State and federal laws governing insurance
20 coverage of contraceptive drugs shall apply to hormonal
21 contraceptives dispensed by a pharmacist under this
22 Section.

23 Section 15. The Counties Code is amended by changing
24 Section 5-1069.3 as follows:

1 (55 ILCS 5/5-1069.3)

2 Sec. 5-1069.3. Required health benefits. If a county,
3 including a home rule county, is a self-insurer for purposes of
4 providing health insurance coverage for its employees, the
5 coverage shall include coverage for the post-mastectomy care
6 benefits required to be covered by a policy of accident and
7 health insurance under Section 356t and the coverage required
8 under Sections 356g, 356g.5, 356g.5-1, 356u, 356w, 356x,
9 356z.6, 356z.8, 356z.9, 356z.10, 356z.11, 356z.12, 356z.13,
10 356z.14, 356z.15, 356z.22, ~~and~~ 356z.25, 356z.26, and 356z.29 of
11 the Illinois Insurance Code. The coverage shall comply with
12 Sections 155.22a, 355b, 356z.19, and 370c of the Illinois
13 Insurance Code. The requirement that health benefits be covered
14 as provided in this Section is an exclusive power and function
15 of the State and is a denial and limitation under Article VII,
16 Section 6, subsection (h) of the Illinois Constitution. A home
17 rule county to which this Section applies must comply with
18 every provision of this Section.

19 Rulemaking authority to implement Public Act 95-1045, if
20 any, is conditioned on the rules being adopted in accordance
21 with all provisions of the Illinois Administrative Procedure
22 Act and all rules and procedures of the Joint Committee on
23 Administrative Rules; any purported rule not so adopted, for
24 whatever reason, is unauthorized.

25 (Source: P.A. 99-480, eff. 9-9-15; 100-24, eff. 7-18-17;
26 100-138, eff. 8-18-17; revised 10-5-17.)

1 Section 20. The Illinois Municipal Code is amended by
2 changing Section 10-4-2.3 as follows:

3 (65 ILCS 5/10-4-2.3)

4 Sec. 10-4-2.3. Required health benefits. If a
5 municipality, including a home rule municipality, is a
6 self-insurer for purposes of providing health insurance
7 coverage for its employees, the coverage shall include coverage
8 for the post-mastectomy care benefits required to be covered by
9 a policy of accident and health insurance under Section 356t
10 and the coverage required under Sections 356g, 356g.5,
11 356g.5-1, 356u, 356w, 356x, 356z.6, 356z.8, 356z.9, 356z.10,
12 356z.11, 356z.12, 356z.13, 356z.14, 356z.15, 356z.22, ~~and~~
13 356z.25, 356z.26, and 356z.29 of the Illinois Insurance Code.
14 The coverage shall comply with Sections 155.22a, 355b, 356z.19,
15 and 370c of the Illinois Insurance Code. The requirement that
16 health benefits be covered as provided in this is an exclusive
17 power and function of the State and is a denial and limitation
18 under Article VII, Section 6, subsection (h) of the Illinois
19 Constitution. A home rule municipality to which this Section
20 applies must comply with every provision of this Section.

21 Rulemaking authority to implement Public Act 95-1045, if
22 any, is conditioned on the rules being adopted in accordance
23 with all provisions of the Illinois Administrative Procedure
24 Act and all rules and procedures of the Joint Committee on

1 Administrative Rules; any purported rule not so adopted, for
2 whatever reason, is unauthorized.

3 (Source: P.A. 99-480, eff. 9-9-15; 100-24, eff. 7-18-17;
4 100-138, eff. 8-18-17; revised 10-5-17.)

5 Section 25. The School Code is amended by changing Section
6 10-22.3f as follows:

7 (105 ILCS 5/10-22.3f)

8 Sec. 10-22.3f. Required health benefits. Insurance
9 protection and benefits for employees shall provide the
10 post-mastectomy care benefits required to be covered by a
11 policy of accident and health insurance under Section 356t and
12 the coverage required under Sections 356g, 356g.5, 356g.5-1,
13 356u, 356w, 356x, 356z.6, 356z.8, 356z.9, 356z.11, 356z.12,
14 356z.13, 356z.14, 356z.15, 356z.22, ~~and~~ 356z.25, 356z.26, and
15 356z.29 of the Illinois Insurance Code. Insurance policies
16 shall comply with Section 356z.19 of the Illinois Insurance
17 Code. The coverage shall comply with Sections 155.22a and 355b
18 of the Illinois Insurance Code.

19 Rulemaking authority to implement Public Act 95-1045, if
20 any, is conditioned on the rules being adopted in accordance
21 with all provisions of the Illinois Administrative Procedure
22 Act and all rules and procedures of the Joint Committee on
23 Administrative Rules; any purported rule not so adopted, for
24 whatever reason, is unauthorized.

1 (Source: P.A. 100-24, eff. 7-18-17; 100-138, eff. 8-18-17;
2 revised 9-25-17.)

3 Section 30. The Illinois Insurance Code is amended by
4 adding Section 356z.29 as follows:

5 (215 ILCS 5/356z.29 new)

6 Sec. 356z.29. Coverage for patient care services for
7 hormonal contraceptives provided by a pharmacist. A group or
8 individual policy of accident and health insurance or a managed
9 care plan that is amended, delivered, issued, or renewed after
10 the effective date of this amendatory Act of the 100th General
11 Assembly shall provide coverage for patient care services
12 provided by a pharmacist for hormonal contraceptives
13 assessment and consultation.

14 Section 35. The Pharmacy Practice Act is amended by
15 changing Section 3 as follows:

16 (225 ILCS 85/3)

17 (Section scheduled to be repealed on January 1, 2020)

18 Sec. 3. Definitions. For the purpose of this Act, except
19 where otherwise limited therein:

20 (a) "Pharmacy" or "drugstore" means and includes every
21 store, shop, pharmacy department, or other place where
22 pharmacist care is provided by a pharmacist (1) where drugs,

1 medicines, or poisons are dispensed, sold or offered for sale
2 at retail, or displayed for sale at retail; or (2) where
3 prescriptions of physicians, dentists, advanced practice
4 registered nurses, physician assistants, veterinarians,
5 podiatric physicians, or optometrists, within the limits of
6 their licenses, are compounded, filled, or dispensed; or (3)
7 which has upon it or displayed within it, or affixed to or used
8 in connection with it, a sign bearing the word or words
9 "Pharmacist", "Druggist", "Pharmacy", "Pharmaceutical Care",
10 "Apothecary", "Drugstore", "Medicine Store", "Prescriptions",
11 "Drugs", "Dispensary", "Medicines", or any word or words of
12 similar or like import, either in the English language or any
13 other language; or (4) where the characteristic prescription
14 sign (Rx) or similar design is exhibited; or (5) any store, or
15 shop, or other place with respect to which any of the above
16 words, objects, signs or designs are used in any advertisement.

17 (b) "Drugs" means and includes (1) articles recognized in
18 the official United States Pharmacopoeia/National Formulary
19 (USP/NF), or any supplement thereto and being intended for and
20 having for their main use the diagnosis, cure, mitigation,
21 treatment or prevention of disease in man or other animals, as
22 approved by the United States Food and Drug Administration, but
23 does not include devices or their components, parts, or
24 accessories; and (2) all other articles intended for and having
25 for their main use the diagnosis, cure, mitigation, treatment
26 or prevention of disease in man or other animals, as approved

1 by the United States Food and Drug Administration, but does not
2 include devices or their components, parts, or accessories; and
3 (3) articles (other than food) having for their main use and
4 intended to affect the structure or any function of the body of
5 man or other animals; and (4) articles having for their main
6 use and intended for use as a component or any articles
7 specified in clause (1), (2) or (3); but does not include
8 devices or their components, parts or accessories.

9 (c) "Medicines" means and includes all drugs intended for
10 human or veterinary use approved by the United States Food and
11 Drug Administration.

12 (d) "Practice of pharmacy" means:

13 (1) the interpretation and the provision of assistance
14 in the monitoring, evaluation, and implementation of
15 prescription drug orders;

16 (2) the dispensing of prescription drug orders;

17 (3) participation in drug and device selection;

18 (4) drug administration limited to the administration
19 of oral, topical, injectable, and inhalation as follows:

20 (A) in the context of patient education on the
21 proper use or delivery of medications;

22 (B) vaccination of patients 14 years of age and
23 older pursuant to a valid prescription or standing
24 order, by a physician licensed to practice medicine in
25 all its branches, upon completion of appropriate
26 training, including how to address contraindications

1 and adverse reactions set forth by rule, with
2 notification to the patient's physician and
3 appropriate record retention, or pursuant to hospital
4 pharmacy and therapeutics committee policies and
5 procedures; and

6 (C) administration of injections of
7 alpha-hydroxyprogesterone caproate, pursuant to a
8 valid prescription, by a physician licensed to
9 practice medicine in all its branches, upon completion
10 of appropriate training, including how to address
11 contraindications and adverse reactions set forth by
12 rule, with notification to the patient's physician and
13 appropriate record retention, or pursuant to hospital
14 pharmacy and therapeutics committee policies and
15 procedures;

16 (5) vaccination of patients ages 10 through 13 limited
17 to the Influenza (inactivated influenza vaccine and live
18 attenuated influenza intranasal vaccine) and Tdap (defined
19 as tetanus, diphtheria, acellular pertussis) vaccines,
20 pursuant to a valid prescription or standing order, by a
21 physician licensed to practice medicine in all its
22 branches, upon completion of appropriate training,
23 including how to address contraindications and adverse
24 reactions set forth by rule, with notification to the
25 patient's physician and appropriate record retention, or
26 pursuant to hospital pharmacy and therapeutics committee

1 policies and procedures;

2 (6) drug regimen review;

3 (7) drug or drug-related research;

4 (8) the provision of patient counseling;

5 (9) the practice of telepharmacy;

6 (10) the provision of those acts or services necessary
7 to provide pharmacist care;

8 (11) medication therapy management; and

9 (12) the responsibility for compounding and labeling
10 of drugs and devices (except labeling by a manufacturer,
11 repackager, or distributor of non-prescription drugs and
12 commercially packaged legend drugs and devices), proper
13 and safe storage of drugs and devices, and maintenance of
14 required records; ~~and-~~

15 (13) the assessment and consultation of patients and
16 dispensing of hormonal contraceptives pursuant to the
17 standing order under Section 2310-700 of the Department of
18 Public Health Powers and Duties Law of the Civil
19 Administrative Code of Illinois.

20 A pharmacist who performs any of the acts defined as the
21 practice of pharmacy in this State must be actively licensed as
22 a pharmacist under this Act.

23 (e) "Prescription" means and includes any written, oral,
24 facsimile, or electronically transmitted order for drugs or
25 medical devices, issued by a physician licensed to practice
26 medicine in all its branches, dentist, veterinarian, podiatric

1 physician, or optometrist, within the limits of his or her
2 license ~~their licenses~~, by a physician assistant in accordance
3 with subsection (f) of Section 4, or by an advanced practice
4 registered nurse in accordance with subsection (g) of Section
5 4, containing the following: (1) name of the patient; (2) date
6 when prescription was issued; (3) name and strength of drug or
7 description of the medical device prescribed; and (4) quantity;
8 (5) directions for use; (6) prescriber's name, address, and
9 signature; and (7) DEA registration number where required, for
10 controlled substances. The prescription may, but is not
11 required to, list the illness, disease, or condition for which
12 the drug or device is being prescribed. DEA registration
13 numbers shall not be required on inpatient drug orders.

14 (f) "Person" means and includes a natural person,
15 partnership, association, corporation, government entity, or
16 any other legal entity.

17 (g) "Department" means the Department of Financial and
18 Professional Regulation.

19 (h) "Board of Pharmacy" or "Board" means the State Board of
20 Pharmacy of the Department of Financial and Professional
21 Regulation.

22 (i) "Secretary" means the Secretary of Financial and
23 Professional Regulation.

24 (j) "Drug product selection" means the interchange for a
25 prescribed pharmaceutical product in accordance with Section
26 25 of this Act and Section 3.14 of the Illinois Food, Drug and

1 Cosmetic Act.

2 (k) "Inpatient drug order" means an order issued by an
3 authorized prescriber for a resident or patient of a facility
4 licensed under the Nursing Home Care Act, the ID/DD Community
5 Care Act, the MC/DD Act, the Specialized Mental Health
6 Rehabilitation Act of 2013, the Hospital Licensing Act, or the
7 University of Illinois Hospital Act ~~"An Act in relation to the~~
8 ~~founding and operation of the University of Illinois Hospital~~
9 ~~and the conduct of University of Illinois health care~~
10 ~~programs", approved July 3, 1931, as amended,~~ or a facility
11 which is operated by the Department of Human Services (as
12 successor to the Department of Mental Health and Developmental
13 Disabilities) or the Department of Corrections.

14 (k-5) "Pharmacist" means an individual health care
15 professional and provider currently licensed by this State to
16 engage in the practice of pharmacy.

17 (l) "Pharmacist in charge" means the licensed pharmacist
18 whose name appears on a pharmacy license and who is responsible
19 for all aspects of the operation related to the practice of
20 pharmacy.

21 (m) "Dispense" or "dispensing" means the interpretation,
22 evaluation, and implementation of a prescription drug order,
23 including the preparation and delivery of a drug or device to a
24 patient or patient's agent in a suitable container
25 appropriately labeled for subsequent administration to or use
26 by a patient in accordance with applicable State and federal

1 laws and regulations. "Dispense" or "dispensing" does not mean
2 the physical delivery to a patient or a patient's
3 representative in a home or institution by a designee of a
4 pharmacist or by common carrier. "Dispense" or "dispensing"
5 also does not mean the physical delivery of a drug or medical
6 device to a patient or patient's representative by a
7 pharmacist's designee within a pharmacy or drugstore while the
8 pharmacist is on duty and the pharmacy is open.

9 (n) "Nonresident pharmacy" means a pharmacy that is located
10 in a state, commonwealth, or territory of the United States,
11 other than Illinois, that delivers, dispenses, or distributes,
12 through the United States Postal Service, commercially
13 acceptable parcel delivery service, or other common carrier, to
14 Illinois residents, any substance which requires a
15 prescription.

16 (o) "Compounding" means the preparation and mixing of
17 components, excluding flavorings, (1) as the result of a
18 prescriber's prescription drug order or initiative based on the
19 prescriber-patient-pharmacist relationship in the course of
20 professional practice or (2) for the purpose of, or incident
21 to, research, teaching, or chemical analysis and not for sale
22 or dispensing. "Compounding" includes the preparation of drugs
23 or devices in anticipation of receiving prescription drug
24 orders based on routine, regularly observed dispensing
25 patterns. Commercially available products may be compounded
26 for dispensing to individual patients only if all of the

1 following conditions are met: (i) the commercial product is not
2 reasonably available from normal distribution channels in a
3 timely manner to meet the patient's needs and (ii) the
4 prescribing practitioner has requested that the drug be
5 compounded.

6 (p) (Blank).

7 (q) (Blank).

8 (r) "Patient counseling" means the communication between a
9 pharmacist or a student pharmacist under the supervision of a
10 pharmacist and a patient or the patient's representative about
11 the patient's medication or device for the purpose of
12 optimizing proper use of prescription medications or devices.
13 "Patient counseling" may include without limitation (1)
14 obtaining a medication history; (2) acquiring a patient's
15 allergies and health conditions; (3) facilitation of the
16 patient's understanding of the intended use of the medication;
17 (4) proper directions for use; (5) significant potential
18 adverse events; (6) potential food-drug interactions; and (7)
19 the need to be compliant with the medication therapy. A
20 pharmacy technician may only participate in the following
21 aspects of patient counseling under the supervision of a
22 pharmacist: (1) obtaining medication history; (2) providing
23 the offer for counseling by a pharmacist or student pharmacist;
24 and (3) acquiring a patient's allergies and health conditions.

25 (s) "Patient profiles" or "patient drug therapy record"
26 means the obtaining, recording, and maintenance of patient

1 prescription information, including prescriptions for
2 controlled substances, and personal information.

3 (t) (Blank).

4 (u) "Medical device" or "device" means an instrument,
5 apparatus, implement, machine, contrivance, implant, in vitro
6 reagent, or other similar or related article, including any
7 component part or accessory, required under federal law to bear
8 the label "Caution: Federal law requires dispensing by or on
9 the order of a physician". A seller of goods and services who,
10 only for the purpose of retail sales, compounds, sells, rents,
11 or leases medical devices shall not, by reasons thereof, be
12 required to be a licensed pharmacy.

13 (v) "Unique identifier" means an electronic signature,
14 handwritten signature or initials, thumb print, or other
15 acceptable biometric or electronic identification process as
16 approved by the Department.

17 (w) "Current usual and customary retail price" means the
18 price that a pharmacy charges to a non-third-party payor.

19 (x) "Automated pharmacy system" means a mechanical system
20 located within the confines of the pharmacy or remote location
21 that performs operations or activities, other than compounding
22 or administration, relative to storage, packaging, dispensing,
23 or distribution of medication, and which collects, controls,
24 and maintains all transaction information.

25 (y) "Drug regimen review" means and includes the evaluation
26 of prescription drug orders and patient records for (1) known

1 allergies; (2) drug or potential therapy contraindications;
2 (3) reasonable dose, duration of use, and route of
3 administration, taking into consideration factors such as age,
4 gender, and contraindications; (4) reasonable directions for
5 use; (5) potential or actual adverse drug reactions; (6)
6 drug-drug interactions; (7) drug-food interactions; (8)
7 drug-disease contraindications; (9) therapeutic duplication;
8 (10) patient laboratory values when authorized and available;
9 (11) proper utilization (including over or under utilization)
10 and optimum therapeutic outcomes; and (12) abuse and misuse.

11 (z) "Electronically transmitted prescription" means a
12 prescription that is created, recorded, or stored by electronic
13 means; issued and validated with an electronic signature; and
14 transmitted by electronic means directly from the prescriber to
15 a pharmacy. An electronic prescription is not an image of a
16 physical prescription that is transferred by electronic means
17 from computer to computer, facsimile to facsimile, or facsimile
18 to computer.

19 (aa) "Medication therapy management services" means a
20 distinct service or group of services offered by licensed
21 pharmacists, physicians licensed to practice medicine in all
22 its branches, advanced practice registered nurses authorized
23 in a written agreement with a physician licensed to practice
24 medicine in all its branches, or physician assistants
25 authorized in guidelines by a supervising physician that
26 optimize therapeutic outcomes for individual patients through

1 improved medication use. In a retail or other non-hospital
2 pharmacy, medication therapy management services shall consist
3 of the evaluation of prescription drug orders and patient
4 medication records to resolve conflicts with the following:

5 (1) known allergies;

6 (2) drug or potential therapy contraindications;

7 (3) reasonable dose, duration of use, and route of
8 administration, taking into consideration factors such as
9 age, gender, and contraindications;

10 (4) reasonable directions for use;

11 (5) potential or actual adverse drug reactions;

12 (6) drug-drug interactions;

13 (7) drug-food interactions;

14 (8) drug-disease contraindications;

15 (9) identification of therapeutic duplication;

16 (10) patient laboratory values when authorized and
17 available;

18 (11) proper utilization (including over or under
19 utilization) and optimum therapeutic outcomes; and

20 (12) drug abuse and misuse.

21 "Medication therapy management services" includes the
22 following:

23 (1) documenting the services delivered and
24 communicating the information provided to patients'
25 prescribers within an appropriate time frame, not to exceed
26 48 hours;

1 (2) providing patient counseling designed to enhance a
2 patient's understanding and the appropriate use of his or
3 her medications; and

4 (3) providing information, support services, and
5 resources designed to enhance a patient's adherence with
6 his or her prescribed therapeutic regimens.

7 "Medication therapy management services" may also include
8 patient care functions authorized by a physician licensed to
9 practice medicine in all its branches for his or her identified
10 patient or groups of patients under specified conditions or
11 limitations in a standing order from the physician.

12 "Medication therapy management services" in a licensed
13 hospital may also include the following:

14 (1) reviewing assessments of the patient's health
15 status; and

16 (2) following protocols of a hospital pharmacy and
17 therapeutics committee with respect to the fulfillment of
18 medication orders.

19 (bb) "Pharmacist care" means the provision by a pharmacist
20 of medication therapy management services, with or without the
21 dispensing of drugs or devices, intended to achieve outcomes
22 that improve patient health, quality of life, and comfort and
23 enhance patient safety.

24 (cc) "Protected health information" means individually
25 identifiable health information that, except as otherwise
26 provided, is:

1 (1) transmitted by electronic media;

2 (2) maintained in any medium set forth in the
3 definition of "electronic media" in the federal Health
4 Insurance Portability and Accountability Act; or

5 (3) transmitted or maintained in any other form or
6 medium.

7 "Protected health information" does not include
8 individually identifiable health information found in:

9 (1) education records covered by the federal Family
10 Educational Right and Privacy Act; or

11 (2) employment records held by a licensee in its role
12 as an employer.

13 (dd) "Standing order" means a specific order for a patient
14 or group of patients issued by a physician licensed to practice
15 medicine in all its branches in Illinois.

16 (ee) "Address of record" means the designated address
17 recorded by the Department in the applicant's application file
18 or licensee's license file maintained by the Department's
19 licensure maintenance unit.

20 (ff) "Home pharmacy" means the location of a pharmacy's
21 primary operations.

22 (gg) "Email address of record" means the designated email
23 address recorded by the Department in the applicant's
24 application file or the licensee's license file, as maintained
25 by the Department's licensure maintenance unit.

26 (Source: P.A. 99-180, eff. 7-29-15; 100-208, eff. 1-1-18;

1 100-497, eff. 9-8-17; 100-513, eff. 1-1-18; revised 9-29-17.)

2 Section 40. The Illinois Public Aid Code is amended by
3 adding Section 5-5.12b as follows:

4 (305 ILCS 5/5-5.12b new)

5 Sec. 5-5.12b. Coverage for patient care services for
6 hormonal contraceptives provided by a pharmacist.

7 (a) Subject to approval by the federal Centers for Medicare
8 and Medicaid Services, the medical assistance program,
9 including both the fee-for-service and managed care medical
10 assistance programs established under this Article, shall
11 cover patient care services provided by a pharmacist for
12 hormonal contraceptives assessment and consultation.

13 (b) The Department shall establish a fee schedule for
14 patient care services provided by a pharmacist for hormonal
15 contraceptives assessment and consultation.

16 (c) The rate of reimbursement for patient care services
17 provided by a pharmacist for hormonal contraceptives
18 assessment and consultation shall be at 85% of the fee schedule
19 for physician services by the medical assistance program.

20 (d) A pharmacist must be enrolled in the medical assistance
21 program as an ordering and referring provider prior to
22 providing hormonal contraceptives assessment and consultation
23 that is submitted by a pharmacy or pharmacist provider for
24 reimbursement pursuant to this Section.

1 (e) The Director shall seek any necessary federal waivers
2 or approvals to implement this Section. This Section shall not
3 be implemented until the receipt of all necessary federal
4 waivers or approvals or until January 1, 2021, whichever comes
5 first. If federal approval is not obtained by January 1, 2021,
6 the provisions of this Section shall be implemented using State
7 funds.

8 (f) This Section does not restrict or prohibit any services
9 currently provided by pharmacists as authorized by law,
10 including, but not limited to, pharmacist services provided
11 under this Code or authorized under the Illinois Title XIX
12 State Plan.

13 (g) The Department shall adopt administrative rules for
14 this Section as soon as practicable but no later than May 1,
15 2019.

16 Section 99. Effective date. This Act takes effect January
17 1, 2019."