

1 AN ACT concerning regulation.

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

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

6 (5 ILCS 375/6.11)

7 Sec. 6.11. Required health benefits; Illinois Insurance
8 Code requirements. The program of health benefits shall provide
9 the post-mastectomy care benefits required to be covered by a
10 policy of accident and health insurance under Section 356t of
11 the Illinois Insurance Code. The program of health benefits
12 shall provide the coverage required under Sections 356g,
13 356g.5, 356g.5-1, 356m, 356u, 356w, 356x, 356z.2, 356z.4,
14 356z.6, 356z.8, 356z.9, 356z.10, 356z.11, 356z.12, 356z.13,
15 356z.14, 356z.15, 356z.17, 356z.22, ~~and~~ 356z.25, 356z.26, and
16 356z.29 of the Illinois Insurance Code. The program of health
17 benefits must comply with Sections 155.22a, 155.37, 355b,
18 356z.19, 370c, and 370c.1 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

1 whatever reason, is unauthorized.

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

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

7 (20 ILCS 2310/2310-700 new)

8 Sec. 2310-700. Contraceptive drugs and products; Director
9 standing order.

10 (a) As used in this Section:

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

15 "Standing order" has the meaning given to that term in the
16 Pharmacy Practice Act.

17 (b) If the Director of Public Health is a physician
18 licensed to practice medicine in all its branches in Illinois,
19 the Director shall establish a standing order complete with the
20 issuance of a prescription for a hormonal contraceptive in
21 accordance with this Section. If the Director is not a
22 physician licensed to practice medicine in all its branches in
23 Illinois, then the Medical Director of the Department of Public
24 Health shall establish a standing order in accordance with this

1 Section.

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

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

6 (2) A pharmacist shall have the patient complete the
7 self-screening risk assessment tool. The self-screening
8 risk assessment tool is to be based on the most current
9 version of the United States Medical Eligibility Criteria
10 for Contraceptive Use published by the federal Centers for
11 Disease Control and Prevention.

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

17 (4) The pharmacist shall provide, during the patient
18 assessment and consultation, counseling and education
19 about all methods of contraception, including methods not
20 covered under the standing order, and their proper use and
21 effectiveness.

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

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

1 (A) complete an educational training program
2 accredited by the Accreditation Council for Pharmacy
3 Education and approved by the Department that is
4 related to the patient self-screening risk assessment,
5 patient assessment, contraceptive counseling and
6 education, and dispensation of hormonal
7 contraceptives; and

8 (B) dispense the hormonal contraceptive to the
9 patient as soon as practicable after meeting the
10 requirements of paragraph (2).

11 (7) All State and federal laws governing insurance
12 coverage of contraceptive drugs shall apply to hormonal
13 contraceptives dispensed by a pharmacist under this
14 Section.

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

17 (55 ILCS 5/5-1069.3)

18 Sec. 5-1069.3. Required health benefits. If a county,
19 including a home rule county, is a self-insurer for purposes of
20 providing health insurance coverage for its employees, the
21 coverage shall include coverage for the post-mastectomy care
22 benefits required to be covered by a policy of accident and
23 health insurance under Section 356t and the coverage required
24 under Sections 356g, 356g.5, 356g.5-1, 356u, 356w, 356x,

1 356z.6, 356z.8, 356z.9, 356z.10, 356z.11, 356z.12, 356z.13,
2 356z.14, 356z.15, 356z.22, ~~and 356z.25,~~ 356z.26, and 356z.29 of
3 the Illinois Insurance Code. The coverage shall comply with
4 Sections 155.22a, 355b, 356z.19, and 370c of the Illinois
5 Insurance Code. The requirement that health benefits be covered
6 as provided in this Section is an exclusive power and function
7 of the State and is a denial and limitation under Article VII,
8 Section 6, subsection (h) of the Illinois Constitution. A home
9 rule county to which this Section applies must comply with
10 every provision of this Section.

11 Rulemaking authority to implement Public Act 95-1045, if
12 any, is conditioned on the rules being adopted in accordance
13 with all provisions of the Illinois Administrative Procedure
14 Act and all rules and procedures of the Joint Committee on
15 Administrative Rules; any purported rule not so adopted, for
16 whatever reason, is unauthorized.

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

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

21 (65 ILCS 5/10-4-2.3)

22 Sec. 10-4-2.3. Required health benefits. If a
23 municipality, including a home rule municipality, is a
24 self-insurer for purposes of providing health insurance

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

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

21 (Source: P.A. 99-480, eff. 9-9-15; 100-24, eff. 7-18-17;
22 100-138, eff. 8-18-17; revised 10-5-17.)

23 Section 25. The School Code is amended by changing Section
24 10-22.3f as follows:

1 (105 ILCS 5/10-22.3f)

2 Sec. 10-22.3f. Required health benefits. Insurance
3 protection and benefits for employees shall provide the
4 post-mastectomy care benefits required to be covered by a
5 policy of accident and health insurance under Section 356t and
6 the coverage required under Sections 356g, 356g.5, 356g.5-1,
7 356u, 356w, 356x, 356z.6, 356z.8, 356z.9, 356z.11, 356z.12,
8 356z.13, 356z.14, 356z.15, 356z.22, ~~and~~ 356z.25, 356z.26, and
9 356z.29 of the Illinois Insurance Code. Insurance policies
10 shall comply with Section 356z.19 of the Illinois Insurance
11 Code. The coverage shall comply with Sections 155.22a and 355b
12 of the Illinois Insurance Code.

13 Rulemaking authority to implement Public Act 95-1045, if
14 any, is conditioned on the rules being adopted in accordance
15 with all provisions of the Illinois Administrative Procedure
16 Act and all rules and procedures of the Joint Committee on
17 Administrative Rules; any purported rule not so adopted, for
18 whatever reason, is unauthorized.

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

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

23 (215 ILCS 5/356z.29 new)

24 Sec. 356z.29. Coverage for patient care services for

1 hormonal contraceptives provided by a pharmacist. A group or
2 individual policy of accident and health insurance or a managed
3 care plan that is amended, delivered, issued, or renewed after
4 the effective date of this amendatory Act of the 100th General
5 Assembly shall provide coverage for patient care services
6 provided by a pharmacist for hormonal contraceptives
7 assessment and consultation.

8 Section 35. The Pharmacy Practice Act is amended by
9 changing Section 3 as follows:

10 (225 ILCS 85/3)

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

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

14 (a) "Pharmacy" or "drugstore" means and includes every
15 store, shop, pharmacy department, or other place where
16 pharmacist care is provided by a pharmacist (1) where drugs,
17 medicines, or poisons are dispensed, sold or offered for sale
18 at retail, or displayed for sale at retail; or (2) where
19 prescriptions of physicians, dentists, advanced practice
20 registered nurses, physician assistants, veterinarians,
21 podiatric physicians, or optometrists, within the limits of
22 their licenses, are compounded, filled, or dispensed; or (3)
23 which has upon it or displayed within it, or affixed to or used
24 in connection with it, a sign bearing the word or words

1 "Pharmacist", "Druggist", "Pharmacy", "Pharmaceutical Care",
2 "Apothecary", "Drugstore", "Medicine Store", "Prescriptions",
3 "Drugs", "Dispensary", "Medicines", or any word or words of
4 similar or like import, either in the English language or any
5 other language; or (4) where the characteristic prescription
6 sign (Rx) or similar design is exhibited; or (5) any store, or
7 shop, or other place with respect to which any of the above
8 words, objects, signs or designs are used in any advertisement.

9 (b) "Drugs" means and includes (1) articles recognized in
10 the official United States Pharmacopoeia/National Formulary
11 (USP/NF), or any supplement thereto and being intended for and
12 having for their main use the diagnosis, cure, mitigation,
13 treatment or prevention of disease in man or other animals, as
14 approved by the United States Food and Drug Administration, but
15 does not include devices or their components, parts, or
16 accessories; and (2) all other articles intended for and having
17 for their main use the diagnosis, cure, mitigation, treatment
18 or prevention of disease in man or other animals, as approved
19 by the United States Food and Drug Administration, but does not
20 include devices or their components, parts, or accessories; and
21 (3) articles (other than food) having for their main use and
22 intended to affect the structure or any function of the body of
23 man or other animals; and (4) articles having for their main
24 use and intended for use as a component or any articles
25 specified in clause (1), (2) or (3); but does not include
26 devices or their components, parts or accessories.

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

4 (d) "Practice of pharmacy" means:

5 (1) the interpretation and the provision of assistance
6 in the monitoring, evaluation, and implementation of
7 prescription drug orders;

8 (2) the dispensing of prescription drug orders;

9 (3) participation in drug and device selection;

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

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

14 (B) vaccination of patients 14 years of age and
15 older pursuant to a valid prescription or standing
16 order, by a physician licensed to practice medicine in
17 all its branches, upon completion of appropriate
18 training, including how to address contraindications
19 and adverse reactions set forth by rule, with
20 notification to the patient's physician and
21 appropriate record retention, or pursuant to hospital
22 pharmacy and therapeutics committee policies and
23 procedures; and

24 (C) administration of injections of
25 alpha-hydroxyprogesterone caproate, pursuant to a
26 valid prescription, by a physician licensed to

1 practice medicine in all its branches, upon completion
2 of appropriate training, including how to address
3 contraindications and adverse reactions set forth by
4 rule, with notification to the patient's physician and
5 appropriate record retention, or pursuant to hospital
6 pharmacy and therapeutics committee policies and
7 procedures;

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

20 (6) drug regimen review;

21 (7) drug or drug-related research;

22 (8) the provision of patient counseling;

23 (9) the practice of telepharmacy;

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

26 (11) medication therapy management; and

1 (12) the responsibility for compounding and labeling
2 of drugs and devices (except labeling by a manufacturer,
3 repackager, or distributor of non-prescription drugs and
4 commercially packaged legend drugs and devices), proper
5 and safe storage of drugs and devices, and maintenance of
6 required records; and-

7 (13) the assessment and consultation of patients and
8 dispensing of hormonal contraceptives pursuant to the
9 standing order under Section 2310-700 of the Department of
10 Public Health Powers and Duties Law of the Civil
11 Administrative Code of Illinois.

12 A pharmacist who performs any of the acts defined as the
13 practice of pharmacy in this State must be actively licensed as
14 a pharmacist under this Act.

15 (e) "Prescription" means and includes any written, oral,
16 facsimile, or electronically transmitted order for drugs or
17 medical devices, issued by a physician licensed to practice
18 medicine in all its branches, dentist, veterinarian, podiatric
19 physician, or optometrist, within the limits of his or her
20 license ~~their licenses~~, by a physician assistant in accordance
21 with subsection (f) of Section 4, or by an advanced practice
22 registered nurse in accordance with subsection (g) of Section
23 4, containing the following: (1) name of the patient; (2) date
24 when prescription was issued; (3) name and strength of drug or
25 description of the medical device prescribed; and (4) quantity;
26 (5) directions for use; (6) prescriber's name, address, and

1 signature; and (7) DEA registration number where required, for
2 controlled substances. The prescription may, but is not
3 required to, list the illness, disease, or condition for which
4 the drug or device is being prescribed. DEA registration
5 numbers shall not be required on inpatient drug orders.

6 (f) "Person" means and includes a natural person,
7 partnership, association, corporation, government entity, or
8 any other legal entity.

9 (g) "Department" means the Department of Financial and
10 Professional Regulation.

11 (h) "Board of Pharmacy" or "Board" means the State Board of
12 Pharmacy of the Department of Financial and Professional
13 Regulation.

14 (i) "Secretary" means the Secretary of Financial and
15 Professional Regulation.

16 (j) "Drug product selection" means the interchange for a
17 prescribed pharmaceutical product in accordance with Section
18 25 of this Act and Section 3.14 of the Illinois Food, Drug and
19 Cosmetic Act.

20 (k) "Inpatient drug order" means an order issued by an
21 authorized prescriber for a resident or patient of a facility
22 licensed under the Nursing Home Care Act, the ID/DD Community
23 Care Act, the MC/DD Act, the Specialized Mental Health
24 Rehabilitation Act of 2013, the Hospital Licensing Act, or the
25 University of Illinois Hospital Act ~~"An Act in relation to the~~
26 ~~founding and operation of the University of Illinois Hospital~~

1 ~~and the conduct of University of Illinois health care~~
2 ~~programs", approved July 3, 1931, as amended,~~ or a facility
3 which is operated by the Department of Human Services (as
4 successor to the Department of Mental Health and Developmental
5 Disabilities) or the Department of Corrections.

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

9 (l) "Pharmacist in charge" means the licensed pharmacist
10 whose name appears on a pharmacy license and who is responsible
11 for all aspects of the operation related to the practice of
12 pharmacy.

13 (m) "Dispense" or "dispensing" means the interpretation,
14 evaluation, and implementation of a prescription drug order,
15 including the preparation and delivery of a drug or device to a
16 patient or patient's agent in a suitable container
17 appropriately labeled for subsequent administration to or use
18 by a patient in accordance with applicable State and federal
19 laws and regulations. "Dispense" or "dispensing" does not mean
20 the physical delivery to a patient or a patient's
21 representative in a home or institution by a designee of a
22 pharmacist or by common carrier. "Dispense" or "dispensing"
23 also does not mean the physical delivery of a drug or medical
24 device to a patient or patient's representative by a
25 pharmacist's designee within a pharmacy or drugstore while the
26 pharmacist is on duty and the pharmacy is open.

1 (n) "Nonresident pharmacy" means a pharmacy that is located
2 in a state, commonwealth, or territory of the United States,
3 other than Illinois, that delivers, dispenses, or distributes,
4 through the United States Postal Service, commercially
5 acceptable parcel delivery service, or other common carrier, to
6 Illinois residents, any substance which requires a
7 prescription.

8 (o) "Compounding" means the preparation and mixing of
9 components, excluding flavorings, (1) as the result of a
10 prescriber's prescription drug order or initiative based on the
11 prescriber-patient-pharmacist relationship in the course of
12 professional practice or (2) for the purpose of, or incident
13 to, research, teaching, or chemical analysis and not for sale
14 or dispensing. "Compounding" includes the preparation of drugs
15 or devices in anticipation of receiving prescription drug
16 orders based on routine, regularly observed dispensing
17 patterns. Commercially available products may be compounded
18 for dispensing to individual patients only if all of the
19 following conditions are met: (i) the commercial product is not
20 reasonably available from normal distribution channels in a
21 timely manner to meet the patient's needs and (ii) the
22 prescribing practitioner has requested that the drug be
23 compounded.

24 (p) (Blank).

25 (q) (Blank).

26 (r) "Patient counseling" means the communication between a

1 pharmacist or a student pharmacist under the supervision of a
2 pharmacist and a patient or the patient's representative about
3 the patient's medication or device for the purpose of
4 optimizing proper use of prescription medications or devices.

5 "Patient counseling" may include without limitation (1)
6 obtaining a medication history; (2) acquiring a patient's
7 allergies and health conditions; (3) facilitation of the
8 patient's understanding of the intended use of the medication;
9 (4) proper directions for use; (5) significant potential
10 adverse events; (6) potential food-drug interactions; and (7)
11 the need to be compliant with the medication therapy. A
12 pharmacy technician may only participate in the following
13 aspects of patient counseling under the supervision of a
14 pharmacist: (1) obtaining medication history; (2) providing
15 the offer for counseling by a pharmacist or student pharmacist;
16 and (3) acquiring a patient's allergies and health conditions.

17 (s) "Patient profiles" or "patient drug therapy record"
18 means the obtaining, recording, and maintenance of patient
19 prescription information, including prescriptions for
20 controlled substances, and personal information.

21 (t) (Blank).

22 (u) "Medical device" or "device" means an instrument,
23 apparatus, implement, machine, contrivance, implant, in vitro
24 reagent, or other similar or related article, including any
25 component part or accessory, required under federal law to bear
26 the label "Caution: Federal law requires dispensing by or on

1 the order of a physician". A seller of goods and services who,
2 only for the purpose of retail sales, compounds, sells, rents,
3 or leases medical devices shall not, by reasons thereof, be
4 required to be a licensed pharmacy.

5 (v) "Unique identifier" means an electronic signature,
6 handwritten signature or initials, thumb print, or other
7 acceptable biometric or electronic identification process as
8 approved by the Department.

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

11 (x) "Automated pharmacy system" means a mechanical system
12 located within the confines of the pharmacy or remote location
13 that performs operations or activities, other than compounding
14 or administration, relative to storage, packaging, dispensing,
15 or distribution of medication, and which collects, controls,
16 and maintains all transaction information.

17 (y) "Drug regimen review" means and includes the evaluation
18 of prescription drug orders and patient records for (1) known
19 allergies; (2) drug or potential therapy contraindications;
20 (3) reasonable dose, duration of use, and route of
21 administration, taking into consideration factors such as age,
22 gender, and contraindications; (4) reasonable directions for
23 use; (5) potential or actual adverse drug reactions; (6)
24 drug-drug interactions; (7) drug-food interactions; (8)
25 drug-disease contraindications; (9) therapeutic duplication;
26 (10) patient laboratory values when authorized and available;

1 (11) proper utilization (including over or under utilization)
2 and optimum therapeutic outcomes; and (12) abuse and misuse.

3 (z) "Electronically transmitted prescription" means a
4 prescription that is created, recorded, or stored by electronic
5 means; issued and validated with an electronic signature; and
6 transmitted by electronic means directly from the prescriber to
7 a pharmacy. An electronic prescription is not an image of a
8 physical prescription that is transferred by electronic means
9 from computer to computer, facsimile to facsimile, or facsimile
10 to computer.

11 (aa) "Medication therapy management services" means a
12 distinct service or group of services offered by licensed
13 pharmacists, physicians licensed to practice medicine in all
14 its branches, advanced practice registered nurses authorized
15 in a written agreement with a physician licensed to practice
16 medicine in all its branches, or physician assistants
17 authorized in guidelines by a supervising physician that
18 optimize therapeutic outcomes for individual patients through
19 improved medication use. In a retail or other non-hospital
20 pharmacy, medication therapy management services shall consist
21 of the evaluation of prescription drug orders and patient
22 medication records to resolve conflicts with the following:

23 (1) known allergies;

24 (2) drug or potential therapy contraindications;

25 (3) reasonable dose, duration of use, and route of
26 administration, taking into consideration factors such as

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

13 "Medication therapy management services" includes the
14 following:

- 15 (1) documenting the services delivered and
- 16 communicating the information provided to patients'
- 17 prescribers within an appropriate time frame, not to exceed
- 18 48 hours;
- 19 (2) providing patient counseling designed to enhance a
- 20 patient's understanding and the appropriate use of his or
- 21 her medications; and
- 22 (3) providing information, support services, and
- 23 resources designed to enhance a patient's adherence with
- 24 his or her prescribed therapeutic regimens.

25 "Medication therapy management services" may also include
26 patient care functions authorized by a physician licensed to

1 practice medicine in all its branches for his or her identified
2 patient or groups of patients under specified conditions or
3 limitations in a standing order from the physician.

4 "Medication therapy management services" in a licensed
5 hospital may also include the following:

6 (1) reviewing assessments of the patient's health
7 status; and

8 (2) following protocols of a hospital pharmacy and
9 therapeutics committee with respect to the fulfillment of
10 medication orders.

11 (bb) "Pharmacist care" means the provision by a pharmacist
12 of medication therapy management services, with or without the
13 dispensing of drugs or devices, intended to achieve outcomes
14 that improve patient health, quality of life, and comfort and
15 enhance patient safety.

16 (cc) "Protected health information" means individually
17 identifiable health information that, except as otherwise
18 provided, is:

19 (1) transmitted by electronic media;

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

23 (3) transmitted or maintained in any other form or
24 medium.

25 "Protected health information" does not include
26 individually identifiable health information found in:

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

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

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

8 (ee) "Address of record" means the designated address
9 recorded by the Department in the applicant's application file
10 or licensee's license file maintained by the Department's
11 licensure maintenance unit.

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

14 (gg) "Email address of record" means the designated email
15 address recorded by the Department in the applicant's
16 application file or the licensee's license file, as maintained
17 by the Department's licensure maintenance unit.

18 (Source: P.A. 99-180, eff. 7-29-15; 100-208, eff. 1-1-18;
19 100-497, eff. 9-8-17; 100-513, eff. 1-1-18; revised 9-29-17.)

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

22 (305 ILCS 5/5-5.12b new)

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

1 (a) Subject to approval by the federal Centers for Medicare
2 and Medicaid Services, the medical assistance program,
3 including both the fee-for-service and managed care medical
4 assistance programs established under this Article, shall
5 cover patient care services provided by a pharmacist for
6 hormonal contraceptives assessment and consultation.

7 (b) The Department shall establish a fee schedule for
8 patient care services provided by a pharmacist for hormonal
9 contraceptives assessment and consultation.

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

14 (d) A pharmacist must be enrolled in the medical assistance
15 program as an ordering and referring provider prior to
16 providing hormonal contraceptives assessment and consultation
17 that is submitted by a pharmacy or pharmacist provider for
18 reimbursement pursuant to this Section.

19 (e) The Director shall seek any necessary federal waivers
20 or approvals to implement this Section. This Section shall not
21 be implemented until the receipt of all necessary federal
22 wavers or approvals or until January 1, 2021, whichever comes
23 first. If federal approval is not obtained by January 1, 2021,
24 the provisions of this Section shall be implemented using State
25 funds.

26 (f) This Section does not restrict or prohibit any services

1 currently provided by pharmacists as authorized by law,
2 including, but not limited to, pharmacist services provided
3 under this Code or authorized under the Illinois Title XIX
4 State Plan.

5 (g) The Department shall adopt administrative rules for
6 this Section as soon as practicable but no later than May 1,
7 2019.

8 Section 99. Effective date. This Act takes effect January
9 1, 2019.