



Sen. Robert Peters

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LRB103 37071 RPS 72851 a

1 AMENDMENT TO HOUSE BILL 5395

2 AMENDMENT NO. _____. Amend House Bill 5395 by replacing
3 everything after the enacting clause with the following:

4 "Article 1.

5 Section 1-1. This Act may be referred to as the Health Care
6 Protection Act.

7 Article 2.

8 Section 2-5. The Illinois Administrative Procedure Act is
9 amended by adding Section 5-45.55 as follows:

10 (5 ILCS 100/5-45.55 new)

11 Sec. 5-45.55. Emergency rulemaking; Network Adequacy and
12 Transparency Act. To provide for the expeditious and timely
13 implementation of the Network Adequacy and Transparency Act,

1 emergency rules implementing federal standards for provider
2 ratios, travel time and distance, and appointment wait times
3 if such standards apply to health insurance coverage regulated
4 by the Department of Insurance and are more stringent than the
5 State standards extant at the time the final federal standards
6 are published may be adopted in accordance with Section 5-45
7 by the Department of Insurance. The adoption of emergency
8 rules authorized by Section 5-45 and this Section is deemed to
9 be necessary for the public interest, safety, and welfare.

10 Section 2-10. The Network Adequacy and Transparency Act is
11 amended by changing Sections 3, 5, 10, 15, 20, 25, and 30 and
12 by adding Sections 35, 36, 40, 50, and 55 as follows:

13 (215 ILCS 124/3)

14 Sec. 3. Applicability of Act. This Act applies to an
15 individual or group policy of ~~accident and~~ health insurance
16 coverage with a network plan amended, delivered, issued, or
17 renewed in this State on or after January 1, 2019. This Act
18 does not apply to an individual or group policy for excepted
19 benefits or short-term, limited-duration health insurance
20 coverage dental or vision insurance or a limited health
21 service organization with a network plan amended, delivered,
22 issued, or renewed in this State on or after January 1, 2019,
23 except to the extent that federal law establishes network
24 adequacy and transparency standards for stand-alone dental

1 plans, which the Department shall enforce for plans amended,
2 delivered, issued, or renewed on or after January 1, 2025.

3 (Source: P.A. 100-502, eff. 9-15-17; 100-601, eff. 6-29-18.)

4 (215 ILCS 124/5)

5 Sec. 5. Definitions. In this Act:

6 "Authorized representative" means a person to whom a
7 beneficiary has given express written consent to represent the
8 beneficiary; a person authorized by law to provide substituted
9 consent for a beneficiary; or the beneficiary's treating
10 provider only when the beneficiary or his or her family member
11 is unable to provide consent.

12 "Beneficiary" means an individual, an enrollee, an
13 insured, a participant, or any other person entitled to
14 reimbursement for covered expenses of or the discounting of
15 provider fees for health care services under a program in
16 which the beneficiary has an incentive to utilize the services
17 of a provider that has entered into an agreement or
18 arrangement with an issuer ~~insurer~~.

19 "Department" means the Department of Insurance.

20 "Essential community provider" has the meaning ascribed to
21 that term in 45 CFR 156.235.

22 "Excepted benefits" has the meaning ascribed to that term
23 in 42 U.S.C. 300gg-91(c).

24 "Exchange" has the meaning ascribed to that term in 45 CFR
25 155.20.

1 "Director" means the Director of Insurance.

2 "Family caregiver" means a relative, partner, friend, or
3 neighbor who has a significant relationship with the patient
4 and administers or assists the patient with activities of
5 daily living, instrumental activities of daily living, or
6 other medical or nursing tasks for the quality and welfare of
7 that patient.

8 "Group health plan" has the meaning ascribed to that term
9 in Section 5 of the Illinois Health Insurance Portability and
10 Accountability Act.

11 "Health insurance coverage" has the meaning ascribed to
12 that term in Section 5 of the Illinois Health Insurance
13 Portability and Accountability Act. "Health insurance
14 coverage" does not include any coverage or benefits under
15 Medicare or under the medical assistance program established
16 under Article V of the Illinois Public Aid Code.

17 "Issuer" means a "health insurance issuer" as defined in
18 Section 5 of the Illinois Health Insurance Portability and
19 Accountability Act.

20 ~~"Insurer" means any entity that offers individual or group~~
21 ~~accident and health insurance, including, but not limited to,~~
22 ~~health maintenance organizations, preferred provider~~
23 ~~organizations, exclusive provider organizations, and other~~
24 ~~plan structures requiring network participation, excluding the~~
25 ~~medical assistance program under the Illinois Public Aid Code,~~
26 ~~the State employees group health insurance program, workers~~

1 ~~compensation insurance, and pharmacy benefit managers.~~

2 "Material change" means a significant reduction in the
3 number of providers available in a network plan, including,
4 but not limited to, a reduction of 10% or more in a specific
5 type of providers within any county, the removal of a major
6 health system that causes a network to be significantly
7 different within any county from the network when the
8 beneficiary purchased the network plan, or any change that
9 would cause the network to no longer satisfy the requirements
10 of this Act or the Department's rules for network adequacy and
11 transparency.

12 "Network" means the group or groups of preferred providers
13 providing services to a network plan.

14 "Network plan" means an individual or group policy of
15 ~~accident and~~ health insurance coverage that either requires a
16 covered person to use or creates incentives, including
17 financial incentives, for a covered person to use providers
18 managed, owned, under contract with, or employed by the issuer
19 or by a third party contracted to arrange, contract for, or
20 administer such provider-related incentives for the issuer
21 insurer.

22 "Ongoing course of treatment" means (1) treatment for a
23 life-threatening condition, which is a disease or condition
24 for which likelihood of death is probable unless the course of
25 the disease or condition is interrupted; (2) treatment for a
26 serious acute condition, defined as a disease or condition

1 requiring complex ongoing care that the covered person is
2 currently receiving, such as chemotherapy, radiation therapy,
3 ~~or~~ post-operative visits, or a serious and complex condition
4 as defined under 42 U.S.C. 300gg-113(b)(2); (3) a course of
5 treatment for a health condition that a treating provider
6 attests that discontinuing care by that provider would worsen
7 the condition or interfere with anticipated outcomes; ~~or~~ (4)
8 the third trimester of pregnancy through the post-partum
9 period; (5) undergoing a course of institutional or inpatient
10 care from the provider within the meaning of 42 U.S.C.
11 300gg-113(b)(1)(B); (6) being scheduled to undergo nonelective
12 surgery from the provider, including receipt of preoperative
13 or postoperative care from such provider with respect to such
14 a surgery; (7) being determined to be terminally ill, as
15 determined under 42 U.S.C. 1395x(dd)(3)(A), and receiving
16 treatment for such illness from such provider; or (8) any
17 other treatment of a condition or disease that requires
18 repeated health care services pursuant to a plan of treatment
19 by a provider because of the potential for changes in the
20 therapeutic regimen or because of the potential for a
21 recurrence of symptoms.

22 "Preferred provider" means any provider who has entered,
23 either directly or indirectly, into an agreement with an
24 employer or risk-bearing entity relating to health care
25 services that may be rendered to beneficiaries under a network
26 plan.

1 "Providers" means physicians licensed to practice medicine
2 in all its branches, other health care professionals,
3 hospitals, or other health care institutions or facilities
4 that provide health care services.

5 "Short-term, limited-duration insurance" means any type of
6 accident and health insurance offered or provided within this
7 State pursuant to a group or individual policy or individual
8 certificate by a company, regardless of the situs state of the
9 delivery of the policy, that has an expiration date specified
10 in the contract that is fewer than 365 days after the original
11 effective date. Regardless of the duration of coverage,
12 "short-term, limited-duration insurance" does not include
13 excepted benefits or any student health insurance coverage.

14 "Stand-alone dental plan" has the meaning ascribed to that
15 term in 45 CFR 156.400.

16 "Telehealth" has the meaning given to that term in Section
17 356z.22 of the Illinois Insurance Code.

18 "Telemedicine" has the meaning given to that term in
19 Section 49.5 of the Medical Practice Act of 1987.

20 "Tiered network" means a network that identifies and
21 groups some or all types of provider and facilities into
22 specific groups to which different provider reimbursement,
23 covered person cost-sharing or provider access requirements,
24 or any combination thereof, apply for the same services.

25 "Woman's principal health care provider" means a physician
26 licensed to practice medicine in all of its branches

1 specializing in obstetrics, gynecology, or family practice.

2 (Source: P.A. 102-92, eff. 7-9-21; 102-813, eff. 5-13-22.)

3 (215 ILCS 124/10)

4 Sec. 10. Network adequacy.

5 (a) Before issuing, delivering, or renewing a network
6 plan, an issuer ~~An insurer~~ providing a network plan shall file
7 a description of all of the following with the Director:

8 (1) The written policies and procedures for adding
9 providers to meet patient needs based on increases in the
10 number of beneficiaries, changes in the
11 patient-to-provider ratio, changes in medical and health
12 care capabilities, and increased demand for services.

13 (2) The written policies and procedures for making
14 referrals within and outside the network.

15 (3) The written policies and procedures on how the
16 network plan will provide 24-hour, 7-day per week access
17 to network-affiliated primary care, emergency services,
18 and women's principal health care providers.

19 An issuer ~~insurer~~ shall not prohibit a preferred provider
20 from discussing any specific or all treatment options with
21 beneficiaries irrespective of the insurer's position on those
22 treatment options or from advocating on behalf of
23 beneficiaries within the utilization review, grievance, or
24 appeals processes established by the issuer ~~insurer~~ in
25 accordance with any rights or remedies available under

1 applicable State or federal law.

2 (b) Before issuing, delivering, or renewing a network
3 plan, an issuer ~~Insurers~~ must file for review a description of
4 the services to be offered through a network plan. The
5 description shall include all of the following:

6 (1) A geographic map of the area proposed to be served
7 by the plan by county service area and zip code, including
8 marked locations for preferred providers.

9 (2) As deemed necessary by the Department, the names,
10 addresses, phone numbers, and specialties of the providers
11 who have entered into preferred provider agreements under
12 the network plan.

13 (3) The number of beneficiaries anticipated to be
14 covered by the network plan.

15 (4) An Internet website and toll-free telephone number
16 for beneficiaries and prospective beneficiaries to access
17 current and accurate lists of preferred providers in each
18 plan, additional information about the plan, as well as
19 any other information required by Department rule.

20 (5) A description of how health care services to be
21 rendered under the network plan are reasonably accessible
22 and available to beneficiaries. The description shall
23 address all of the following:

24 (A) the type of health care services to be
25 provided by the network plan;

26 (B) the ratio of physicians and other providers to

1 beneficiaries, by specialty and including primary care
2 physicians and facility-based physicians when
3 applicable under the contract, necessary to meet the
4 health care needs and service demands of the currently
5 enrolled population;

6 (C) the travel and distance standards for plan
7 beneficiaries in county service areas; and

8 (D) a description of how the use of telemedicine,
9 telehealth, or mobile care services may be used to
10 partially meet the network adequacy standards, if
11 applicable.

12 (6) A provision ensuring that whenever a beneficiary
13 has made a good faith effort, as evidenced by accessing
14 the provider directory, calling the network plan, and
15 calling the provider, to utilize preferred providers for a
16 covered service and it is determined the insurer does not
17 have the appropriate preferred providers due to
18 insufficient number, type, unreasonable travel distance or
19 delay, or preferred providers refusing to provide a
20 covered service because it is contrary to the conscience
21 of the preferred providers, as protected by the Health
22 Care Right of Conscience Act, the issuer ~~insurer~~ shall
23 ensure, directly or indirectly, by terms contained in the
24 payer contract, that the beneficiary will be provided the
25 covered service at no greater cost to the beneficiary than
26 if the service had been provided by a preferred provider.

1 This paragraph (6) does not apply to: (A) a beneficiary
2 who willfully chooses to access a non-preferred provider
3 for health care services available through the panel of
4 preferred providers, or (B) a beneficiary enrolled in a
5 health maintenance organization. In these circumstances,
6 the contractual requirements for non-preferred provider
7 reimbursements shall apply unless Section 356z.3a of the
8 Illinois Insurance Code requires otherwise. In no event
9 shall a beneficiary who receives care at a participating
10 health care facility be required to search for
11 participating providers under the circumstances described
12 in subsection (b) or (b-5) of Section 356z.3a of the
13 Illinois Insurance Code except under the circumstances
14 described in paragraph (2) of subsection (b-5).

15 (7) A provision that the beneficiary shall receive
16 emergency care coverage such that payment for this
17 coverage is not dependent upon whether the emergency
18 services are performed by a preferred or non-preferred
19 provider and the coverage shall be at the same benefit
20 level as if the service or treatment had been rendered by a
21 preferred provider. For purposes of this paragraph (7),
22 "the same benefit level" means that the beneficiary is
23 provided the covered service at no greater cost to the
24 beneficiary than if the service had been provided by a
25 preferred provider. This provision shall be consistent
26 with Section 356z.3a of the Illinois Insurance Code.

1 (8) A limitation that, if the plan provides that the
2 beneficiary will incur a penalty for failing to
3 pre-certify inpatient hospital treatment, the penalty may
4 not exceed \$1,000 per occurrence in addition to the plan
5 cost sharing provisions.

6 (9) For a network plan to be offered through the
7 Exchange in the individual or small group market, as well
8 as any off-Exchange mirror of such a network plan,
9 evidence that the network plan includes essential
10 community providers in accordance with rules established
11 by the Exchange that will operate in this State for the
12 applicable plan year.

13 (c) The issuer ~~network plan~~ shall demonstrate to the
14 Director a minimum ratio of providers to plan beneficiaries as
15 required by the Department for each network plan.

16 (1) The minimum ratio of physicians or other providers
17 to plan beneficiaries shall be established ~~annually~~ by the
18 Department in consultation with the Department of Public
19 Health based upon the guidance from the federal Centers
20 for Medicare and Medicaid Services. The Department shall
21 not establish ratios for vision or dental providers who
22 provide services under dental-specific or vision-specific
23 benefits, except to the extent provided under federal law
24 for stand-alone dental plans. The Department shall
25 consider establishing ratios for the following physicians
26 or other providers:

- 1 (A) Primary Care;
- 2 (B) Pediatrics;
- 3 (C) Cardiology;
- 4 (D) Gastroenterology;
- 5 (E) General Surgery;
- 6 (F) Neurology;
- 7 (G) OB/GYN;
- 8 (H) Oncology/Radiation;
- 9 (I) Ophthalmology;
- 10 (J) Urology;
- 11 (K) Behavioral Health;
- 12 (L) Allergy/Immunology;
- 13 (M) Chiropractic;
- 14 (N) Dermatology;
- 15 (O) Endocrinology;
- 16 (P) Ears, Nose, and Throat (ENT)/Otolaryngology;
- 17 (Q) Infectious Disease;
- 18 (R) Nephrology;
- 19 (S) Neurosurgery;
- 20 (T) Orthopedic Surgery;
- 21 (U) Physiatry/Rehabilitative;
- 22 (V) Plastic Surgery;
- 23 (W) Pulmonary;
- 24 (X) Rheumatology;
- 25 (Y) Anesthesiology;
- 26 (Z) Pain Medicine;

1 (AA) Pediatric Specialty Services;

2 (BB) Outpatient Dialysis; and

3 (CC) HIV.

4 (2) The Director shall establish a process for the
5 review of the adequacy of these standards, along with an
6 assessment of additional specialties to be included in the
7 list under this subsection (c).

8 (3) Notwithstanding any other law or rule, the minimum
9 ratio for each provider type shall be no less than any such
10 ratio established for qualified health plans in
11 Federally-Facilitated Exchanges by federal law or by the
12 federal Centers for Medicare and Medicaid Services, even
13 if the network plan is issued in the large group market or
14 is otherwise not issued through an exchange. Federal
15 standards for stand-alone dental plans shall only apply to
16 such network plans. In the absence of an applicable
17 Department rule, the federal standards shall apply for the
18 time period specified in the federal law, regulation, or
19 guidance. If the Centers for Medicare and Medicaid
20 Services establish standards that are more stringent than
21 the standards in effect under any Department rule, the
22 Department may amend its rules to conform to the more
23 stringent federal standards.

24 (d) The network plan shall demonstrate to the Director
25 maximum travel and distance standards and appointment wait
26 time standards for plan beneficiaries, which shall be

1 established ~~annually~~ by the Department in consultation with
2 the Department of Public Health based upon the guidance from
3 the federal Centers for Medicare and Medicaid Services. These
4 standards shall consist of the maximum minutes or miles to be
5 traveled by a plan beneficiary for each county type, such as
6 large counties, metro counties, or rural counties as defined
7 by Department rule.

8 The maximum travel time and distance standards must
9 include standards for each physician and other provider
10 category listed for which ratios have been established.

11 The Director shall establish a process for the review of
12 the adequacy of these standards along with an assessment of
13 additional specialties to be included in the list under this
14 subsection (d).

15 Notwithstanding any other law or Department rule, the
16 maximum travel time and distance standards and appointment
17 wait time standards shall be no greater than any such
18 standards established for qualified health plans in
19 Federally-Facilitated Exchanges by federal law or by the
20 federal Centers for Medicare and Medicaid Services, even if
21 the network plan is issued in the large group market or is
22 otherwise not issued through an exchange. Federal standards
23 for stand-alone dental plans shall only apply to such network
24 plans. In the absence of an applicable Department rule, the
25 federal standards shall apply for the time period specified in
26 the federal law, regulation, or guidance. If the Centers for

1 Medicare and Medicaid Services establish standards that are
2 more stringent than the standards in effect under any
3 Department rule, the Department may amend its rules to conform
4 to the more stringent federal standards.

5 If the federal area designations for the maximum time or
6 distance or appointment wait time standards required are
7 changed by the most recent Letter to Issuers in the
8 Federally-facilitated Marketplaces, the Department shall post
9 on its website notice of such changes and may amend its rules
10 to conform to those designations if the Director deems
11 appropriate.

12 (d-5) (1) Every issuer ~~insurer~~ shall ensure that
13 beneficiaries have timely and proximate access to treatment
14 for mental, emotional, nervous, or substance use disorders or
15 conditions in accordance with the provisions of paragraph (4)
16 of subsection (a) of Section 370c of the Illinois Insurance
17 Code. Issuers ~~Insurers~~ shall use a comparable process,
18 strategy, evidentiary standard, and other factors in the
19 development and application of the network adequacy standards
20 for timely and proximate access to treatment for mental,
21 emotional, nervous, or substance use disorders or conditions
22 and those for the access to treatment for medical and surgical
23 conditions. As such, the network adequacy standards for timely
24 and proximate access shall equally be applied to treatment
25 facilities and providers for mental, emotional, nervous, or
26 substance use disorders or conditions and specialists

1 providing medical or surgical benefits pursuant to the parity
2 requirements of Section 370c.1 of the Illinois Insurance Code
3 and the federal Paul Wellstone and Pete Domenici Mental Health
4 Parity and Addiction Equity Act of 2008. Notwithstanding the
5 foregoing, the network adequacy standards for timely and
6 proximate access to treatment for mental, emotional, nervous,
7 or substance use disorders or conditions shall, at a minimum,
8 satisfy the following requirements:

9 (A) For beneficiaries residing in the metropolitan
10 counties of Cook, DuPage, Kane, Lake, McHenry, and Will,
11 network adequacy standards for timely and proximate access
12 to treatment for mental, emotional, nervous, or substance
13 use disorders or conditions means a beneficiary shall not
14 have to travel longer than 30 minutes or 30 miles from the
15 beneficiary's residence to receive outpatient treatment
16 for mental, emotional, nervous, or substance use disorders
17 or conditions. Beneficiaries shall not be required to wait
18 longer than 10 business days between requesting an initial
19 appointment and being seen by the facility or provider of
20 mental, emotional, nervous, or substance use disorders or
21 conditions for outpatient treatment or to wait longer than
22 20 business days between requesting a repeat or follow-up
23 appointment and being seen by the facility or provider of
24 mental, emotional, nervous, or substance use disorders or
25 conditions for outpatient treatment; however, subject to
26 the protections of paragraph (3) of this subsection, a

1 network plan shall not be held responsible if the
2 beneficiary or provider voluntarily chooses to schedule an
3 appointment outside of these required time frames.

4 (B) For beneficiaries residing in Illinois counties
5 other than those counties listed in subparagraph (A) of
6 this paragraph, network adequacy standards for timely and
7 proximate access to treatment for mental, emotional,
8 nervous, or substance use disorders or conditions means a
9 beneficiary shall not have to travel longer than 60
10 minutes or 60 miles from the beneficiary's residence to
11 receive outpatient treatment for mental, emotional,
12 nervous, or substance use disorders or conditions.
13 Beneficiaries shall not be required to wait longer than 10
14 business days between requesting an initial appointment
15 and being seen by the facility or provider of mental,
16 emotional, nervous, or substance use disorders or
17 conditions for outpatient treatment or to wait longer than
18 20 business days between requesting a repeat or follow-up
19 appointment and being seen by the facility or provider of
20 mental, emotional, nervous, or substance use disorders or
21 conditions for outpatient treatment; however, subject to
22 the protections of paragraph (3) of this subsection, a
23 network plan shall not be held responsible if the
24 beneficiary or provider voluntarily chooses to schedule an
25 appointment outside of these required time frames.

26 (2) For beneficiaries residing in all Illinois counties,

1 network adequacy standards for timely and proximate access to
2 treatment for mental, emotional, nervous, or substance use
3 disorders or conditions means a beneficiary shall not have to
4 travel longer than 60 minutes or 60 miles from the
5 beneficiary's residence to receive inpatient or residential
6 treatment for mental, emotional, nervous, or substance use
7 disorders or conditions.

8 (3) If there is no in-network facility or provider
9 available for a beneficiary to receive timely and proximate
10 access to treatment for mental, emotional, nervous, or
11 substance use disorders or conditions in accordance with the
12 network adequacy standards outlined in this subsection, the
13 issuer ~~insurer~~ shall provide necessary exceptions to its
14 network to ensure admission and treatment with a provider or
15 at a treatment facility in accordance with the network
16 adequacy standards in this subsection.

17 (4) If the federal Centers for Medicare and Medicaid
18 Services establishes or law requires more stringent standards
19 for qualified health plans in the Federally-Facilitated
20 Exchanges, the federal standards shall control for all network
21 plans for the time period specified in the federal law,
22 regulation, or guidance, even if the network plan is issued in
23 the large group market, is issued through a different type of
24 Exchange, or is otherwise not issued through an Exchange.

25 (e) Except for network plans solely offered as a group
26 health plan, these ratio and time and distance standards apply

1 to the lowest cost-sharing tier of any tiered network.

2 (f) The network plan may consider use of other health care
3 service delivery options, such as telemedicine or telehealth,
4 mobile clinics, and centers of excellence, or other ways of
5 delivering care to partially meet the requirements set under
6 this Section.

7 (g) Except for the requirements set forth in subsection
8 (d-5), issuers ~~insurers~~ who are not able to comply with the
9 provider ratios and time and distance or appointment wait time
10 standards established under this Act or federal law ~~by the~~
11 ~~Department~~ may request an exception to these requirements from
12 the Department. The Department may grant an exception in the
13 following circumstances:

14 (1) if no providers or facilities meet the specific
15 time and distance standard in a specific service area and
16 the issuer ~~insurer~~ (i) discloses information on the
17 distance and travel time points that beneficiaries would
18 have to travel beyond the required criterion to reach the
19 next closest contracted provider outside of the service
20 area and (ii) provides contact information, including
21 names, addresses, and phone numbers for the next closest
22 contracted provider or facility;

23 (2) if patterns of care in the service area do not
24 support the need for the requested number of provider or
25 facility type and the issuer ~~insurer~~ provides data on
26 local patterns of care, such as claims data, referral

1 patterns, or local provider interviews, indicating where
2 the beneficiaries currently seek this type of care or
3 where the physicians currently refer beneficiaries, or
4 both; or

5 (3) other circumstances deemed appropriate by the
6 Department consistent with the requirements of this Act.

7 (h) Issuers ~~Insurers~~ are required to report to the
8 Director any material change to an approved network plan
9 within 15 business days after the change occurs and any change
10 that would result in failure to meet the requirements of this
11 Act. The issuer shall submit a revised version of the portions
12 of the network adequacy filing affected by the material
13 change, as determined by the Director by rule, and the issuer
14 shall attach versions with the changes indicated for each
15 document that was revised from the previous version of the
16 filing. Upon notice from the issuer ~~insurer~~, the Director
17 shall reevaluate the network plan's compliance with the
18 network adequacy and transparency standards of this Act. For
19 every day past 15 business days that the issuer fails to submit
20 a revised network adequacy filing to the Director, the
21 Director may order a fine of \$5,000 per day.

22 (i) If a network plan is inadequate under this Act with
23 respect to a provider type in a county, and if the network plan
24 does not have an approved exception for that provider type in
25 that county pursuant to subsection (g), an issuer shall cover
26 out-of-network claims for covered health care services

1 received from that provider type within that county at the
2 in-network benefit level and shall retroactively adjudicate
3 and reimburse beneficiaries to achieve that objective if their
4 claims were processed at the out-of-network level contrary to
5 this subsection. Nothing in this subsection shall be construed
6 to supersede Section 356z.3a of the Illinois Insurance Code.

7 (j) If the Director determines that a network is
8 inadequate in any county and no exception has been granted
9 under subsection (g) and the issuer does not have a process in
10 place to comply with subsection (d-5), the Director may
11 prohibit the network plan from being issued or renewed within
12 that county until the Director determines that the network is
13 adequate apart from processes and exceptions described in
14 subsections (d-5) and (g). Nothing in this subsection shall be
15 construed to terminate any beneficiary's health insurance
16 coverage under a network plan before the expiration of the
17 beneficiary's policy period if the Director makes a
18 determination under this subsection after the issuance or
19 renewal of the beneficiary's policy or certificate because of
20 a material change. Policies or certificates issued or renewed
21 in violation of this subsection may subject the issuer to a
22 civil penalty of \$5,000 per policy.

23 (k) For the Department to enforce any new or modified
24 federal standard before the Department adopts the standard by
25 rule, the Department must, no later than May 15 before the
26 start of the plan year, give public notice to the affected

1 health insurance issuers through a bulletin.

2 (Source: P.A. 102-144, eff. 1-1-22; 102-901, eff. 7-1-22;
3 102-1117, eff. 1-13-23.)

4 (215 ILCS 124/15)

5 Sec. 15. Notice of nonrenewal or termination.

6 (a) A network plan must give at least 60 days' notice of
7 nonrenewal or termination of a provider to the provider and to
8 the beneficiaries served by the provider. The notice shall
9 include a name and address to which a beneficiary or provider
10 may direct comments and concerns regarding the nonrenewal or
11 termination and the telephone number maintained by the
12 Department for consumer complaints. Immediate written notice
13 may be provided without 60 days' notice when a provider's
14 license has been disciplined by a State licensing board or
15 when the network plan reasonably believes direct imminent
16 physical harm to patients under the provider's ~~providers~~ care
17 may occur. The notice to the beneficiary shall provide the
18 individual with an opportunity to notify the issuer of the
19 individual's need for transitional care.

20 (b) Primary care providers must notify active affected
21 patients of nonrenewal or termination of the provider from the
22 network plan, except in the case of incapacitation.

23 (Source: P.A. 100-502, eff. 9-15-17.)

24 (215 ILCS 124/20)

1 Sec. 20. Transition of services.

2 (a) A network plan shall provide for continuity of care
3 for its beneficiaries as follows:

4 (1) If a beneficiary's ~~physician or hospital~~ provider
5 leaves the network plan's network of providers for reasons
6 other than termination of a contract in situations
7 involving imminent harm to a patient or a final
8 disciplinary action by a State licensing board and the
9 provider remains within the network plan's service area,
10 if benefits provided under such network plan with respect
11 to such provider or facility are terminated because of a
12 change in the terms of the participation of such provider
13 or facility in such plan, or if a contract between a group
14 health plan and a health insurance issuer offering a
15 network plan in connection with the group health plan is
16 terminated and results in a loss of benefits provided
17 under such plan with respect to such provider, then the
18 network plan shall permit the beneficiary to continue an
19 ongoing course of treatment with that provider during a
20 transitional period for the following duration:

21 (A) 90 days from the date of the notice to the
22 beneficiary of the provider's disaffiliation from the
23 network plan if the beneficiary has an ongoing course
24 of treatment; or

25 (B) if the beneficiary has entered the third
26 trimester of pregnancy at the time of the provider's

1 disaffiliation, a period that includes the provision
2 of post-partum care directly related to the delivery.

3 (2) Notwithstanding the provisions of paragraph (1) of
4 this subsection (a), such care shall be authorized by the
5 network plan during the transitional period in accordance
6 with the following:

7 (A) the provider receives continued reimbursement
8 from the network plan at the rates and terms and
9 conditions applicable under the terminated contract
10 prior to the start of the transitional period;

11 (B) the provider adheres to the network plan's
12 quality assurance requirements, including provision to
13 the network plan of necessary medical information
14 related to such care; and

15 (C) the provider otherwise adheres to the network
16 plan's policies and procedures, including, but not
17 limited to, procedures regarding referrals and
18 obtaining preauthorizations for treatment.

19 (3) The provisions of this Section governing health
20 care provided during the transition period do not apply if
21 the beneficiary has successfully transitioned to another
22 provider participating in the network plan, if the
23 beneficiary has already met or exceeded the benefit
24 limitations of the plan, or if the care provided is not
25 medically necessary.

26 (b) A network plan shall provide for continuity of care

1 for new beneficiaries as follows:

2 (1) If a new beneficiary whose provider is not a
3 member of the network plan's provider network, but is
4 within the network plan's service area, enrolls in the
5 network plan, the network plan shall permit the
6 beneficiary to continue an ongoing course of treatment
7 with the beneficiary's current physician during a
8 transitional period:

9 (A) of 90 days from the effective date of
10 enrollment if the beneficiary has an ongoing course of
11 treatment; or

12 (B) if the beneficiary has entered the third
13 trimester of pregnancy at the effective date of
14 enrollment, that includes the provision of post-partum
15 care directly related to the delivery.

16 (2) If a beneficiary, or a beneficiary's authorized
17 representative, elects in writing to continue to receive
18 care from such provider pursuant to paragraph (1) of this
19 subsection (b), such care shall be authorized by the
20 network plan for the transitional period in accordance
21 with the following:

22 (A) the provider receives reimbursement from the
23 network plan at rates established by the network plan;

24 (B) the provider adheres to the network plan's
25 quality assurance requirements, including provision to
26 the network plan of necessary medical information

1 related to such care; and

2 (C) the provider otherwise adheres to the network
3 plan's policies and procedures, including, but not
4 limited to, procedures regarding referrals and
5 obtaining preauthorization for treatment.

6 (3) The provisions of this Section governing health
7 care provided during the transition period do not apply if
8 the beneficiary has successfully transitioned to another
9 provider participating in the network plan, if the
10 beneficiary has already met or exceeded the benefit
11 limitations of the plan, or if the care provided is not
12 medically necessary.

13 (c) In no event shall this Section be construed to require
14 a network plan to provide coverage for benefits not otherwise
15 covered or to diminish or impair preexisting condition
16 limitations contained in the beneficiary's contract.

17 (d) A provider shall comply with the requirements of 42
18 U.S.C. 300gg-138.

19 (Source: P.A. 100-502, eff. 9-15-17.)

20 (215 ILCS 124/25)

21 Sec. 25. Network transparency.

22 (a) A network plan shall post electronically an
23 up-to-date, accurate, and complete provider directory for each
24 of its network plans, with the information and search
25 functions, as described in this Section.

1 (1) In making the directory available electronically,
2 the network plans shall ensure that the general public is
3 able to view all of the current providers for a plan
4 through a clearly identifiable link or tab and without
5 creating or accessing an account or entering a policy or
6 contract number.

7 (2) An issuer's failure to update a network plan's
8 directory shall subject the issuer to a civil penalty of
9 \$5,000 per month. The network plan shall update the online
10 provider directory at least monthly. Providers shall
11 notify the network plan electronically or in writing
12 within 10 business days of any changes to their
13 information as listed in the provider directory, including
14 the information required in subsections (b), (c), and (d)
15 subparagraph (K) of paragraph (1) of subsection (b). With
16 regard to subparagraph (I) of paragraph (1) of subsection
17 (b), the provider must give notice to the issuer within 20
18 business days of deciding to cease accepting new patients
19 covered by the plan if the new patient limitation is
20 expected to last 40 business days or longer. The network
21 plan shall update its online provider directory in a
22 manner consistent with the information provided by the
23 provider within 2 ~~10~~ business days after being notified of
24 the change by the provider. Nothing in this paragraph (2)
25 shall void any contractual relationship between the
26 provider and the plan.

1 (3) At least once every 90 days, the issuer ~~The~~
2 ~~network plan~~ shall audit each network plan's ~~periodically~~
3 ~~at least 25% of its~~ provider directories for accuracy,
4 make any corrections necessary, and retain documentation
5 of the audit. The issuer shall submit the self-audit and a
6 summary to the Department, and the Department shall make
7 the summary of each self-audit publicly available. The
8 Department shall specify the requirements of the summary,
9 which shall be statistical in nature except for a
10 high-level narrative evaluating the impact of internal and
11 external factors on the accuracy of the directory and the
12 timeliness of updates. ~~The network plan shall submit the~~
13 ~~audit to the Director upon request.~~ As part of these
14 audits, the network plan shall contact any provider in its
15 network that has not submitted a claim to the plan or
16 otherwise communicated his or her intent to continue
17 participation in the plan's network. The audits shall
18 comply with 42 U.S.C. 300gg-115(a)(2), except that
19 "provider directory information" shall include all
20 information required to be included in a provider
21 directory pursuant to this Act.

22 (4) A network plan shall provide a print copy of a
23 current provider directory or a print copy of the
24 requested directory information upon request of a
25 beneficiary or a prospective beneficiary. Except when an
26 issuer's print copies use the same provider information as

1 the electronic provider directory on each print copy's
2 date of printing, print ~~Print~~ copies must be updated at
3 least every 90 days ~~quarterly~~ and ~~an~~ errata that reflects
4 changes in the provider network must be included in each
5 update ~~updated quarterly~~.

6 (5) For each network plan, a network plan shall
7 include, in plain language in both the electronic and
8 print directory, the following general information:

9 (A) in plain language, a description of the
10 criteria the plan has used to build its provider
11 network;

12 (B) if applicable, in plain language, a
13 description of the criteria the issuer ~~insurer~~ or
14 network plan has used to create tiered networks;

15 (C) if applicable, in plain language, how the
16 network plan designates the different provider tiers
17 or levels in the network and identifies for each
18 specific provider, hospital, or other type of facility
19 in the network which tier each is placed, for example,
20 by name, symbols, or grouping, in order for a
21 beneficiary-covered person or a prospective
22 beneficiary-covered person to be able to identify the
23 provider tier; ~~and~~

24 (D) if applicable, a notation that authorization
25 or referral may be required to access some providers; ~~i-~~

26 (E) a telephone number and email address for a

1 customer service representative to whom directory
2 inaccuracies may be reported; and

3 (F) a detailed description of the process to
4 dispute charges for out-of-network providers,
5 hospitals, or facilities that were incorrectly listed
6 as in-network prior to the provision of care and a
7 telephone number and email address to dispute such
8 charges.

9 (6) A network plan shall make it clear for both its
10 electronic and print directories what provider directory
11 applies to which network plan, such as including the
12 specific name of the network plan as marketed and issued
13 in this State. The network plan shall include in both its
14 electronic and print directories a customer service email
15 address and telephone number or electronic link that
16 beneficiaries or the general public may use to notify the
17 network plan of inaccurate provider directory information
18 and contact information for the Department's Office of
19 Consumer Health Insurance.

20 (7) A provider directory, whether in electronic or
21 print format, shall accommodate the communication needs of
22 individuals with disabilities, and include a link to or
23 information regarding available assistance for persons
24 with limited English proficiency.

25 (b) For each network plan, a network plan shall make
26 available through an electronic provider directory the

1 following information in a searchable format:

2 (1) for health care professionals:

3 (A) name;

4 (B) gender;

5 (C) participating office locations;

6 (D) patient population served (such as pediatric,
7 adult, elderly, or women) and specialty or
8 subspecialty, if applicable;

9 (E) medical group affiliations, if applicable;

10 (F) facility affiliations, if applicable;

11 (G) participating facility affiliations, if
12 applicable;

13 (H) languages spoken other than English, if
14 applicable;

15 (I) whether accepting new patients;

16 (J) board certifications, if applicable; ~~and~~

17 (K) use of telehealth or telemedicine, including,
18 but not limited to:

19 (i) whether the provider offers the use of
20 telehealth or telemedicine to deliver services to
21 patients for whom it would be clinically
22 appropriate;

23 (ii) what modalities are used and what types
24 of services may be provided via telehealth or
25 telemedicine; and

26 (iii) whether the provider has the ability and

1 willingness to include in a telehealth or
2 telemedicine encounter a family caregiver who is
3 in a separate location than the patient if the
4 patient wishes and provides his or her consent;

5 (L) whether the health care professional accepts
6 appointment requests from patients; and

7 (M) the anticipated date the provider will leave
8 the network, if applicable, which shall be included no
9 more than 10 days after the issuer confirms that the
10 provider is scheduled to leave the network;

11 (2) for hospitals:

12 (A) hospital name;

13 (B) hospital type (such as acute, rehabilitation,
14 children's, or cancer);

15 (C) participating hospital location; ~~and~~

16 (D) hospital accreditation status; and

17 (E) the anticipated date the hospital will leave
18 the network, if applicable, which shall be included no
19 more than 10 days after the issuer confirms the
20 hospital is scheduled to leave the network; and

21 (3) for facilities, other than hospitals, by type:

22 (A) facility name;

23 (B) facility type;

24 (C) types of services performed; ~~and~~

25 (D) participating facility location or locations;

26 and-

1 (E) the anticipated date the facility will leave
2 the network, if applicable, which shall be included no
3 more than 10 days after the issuer confirms the
4 facility is scheduled to leave the network.

5 (c) For the electronic provider directories, for each
6 network plan, a network plan shall make available all of the
7 following information in addition to the searchable
8 information required in this Section:

9 (1) for health care professionals:

10 (A) contact information, including both a
11 telephone number and digital contact information if
12 the provider has supplied digital contact information;

13 and

14 (B) languages spoken other than English by
15 clinical staff, if applicable;

16 (2) for hospitals, telephone number and digital
17 contact information; and

18 (3) for facilities other than hospitals, telephone
19 number.

20 (d) The issuer ~~insurer~~ or network plan shall make
21 available in print, upon request, the following provider
22 directory information for the applicable network plan:

23 (1) for health care professionals:

24 (A) name;

25 (B) contact information, including a telephone
26 number and digital contact information if the provider

1 has supplied digital contact information;

2 (C) participating office location or locations;

3 (D) patient population (such as pediatric, adult,
4 elderly, or women) and specialty or subspecialty, if
5 applicable;

6 (E) languages spoken other than English, if
7 applicable;

8 (F) whether accepting new patients; ~~and~~

9 (G) use of telehealth or telemedicine, including,
10 but not limited to:

11 (i) whether the provider offers the use of
12 telehealth or telemedicine to deliver services to
13 patients for whom it would be clinically
14 appropriate;

15 (ii) what modalities are used and what types
16 of services may be provided via telehealth or
17 telemedicine; and

18 (iii) whether the provider has the ability and
19 willingness to include in a telehealth or
20 telemedicine encounter a family caregiver who is
21 in a separate location than the patient if the
22 patient wishes and provides his or her consent;
23 and

24 (H) whether the health care professional accepts
25 appointment requests from patients.

26 (2) for hospitals:

- 1 (A) hospital name;
- 2 (B) hospital type (such as acute, rehabilitation,
3 children's, or cancer); and
- 4 (C) participating hospital location, ~~and~~ telephone
5 number, and digital contact information; and
- 6 (3) for facilities, other than hospitals, by type:
- 7 (A) facility name;
- 8 (B) facility type;
- 9 (C) patient population (such as pediatric, adult,
10 elderly, or women) served, if applicable, and types of
11 services performed; and
- 12 (D) participating facility location or locations, ~~and~~
13 telephone numbers, and digital contact information
14 for each location.
- 15 (e) The network plan shall include a disclosure in the
16 print format provider directory that the information included
17 in the directory is accurate as of the date of printing and
18 that beneficiaries or prospective beneficiaries should consult
19 the issuer's ~~insurer's~~ electronic provider directory on its
20 website and contact the provider. The network plan shall also
21 include a telephone number and email address in the print
22 format provider directory for a customer service
23 representative where the beneficiary can obtain current
24 provider directory information or report provider directory
25 inaccuracies. The printed provider directory shall include a
26 detailed description of the process to dispute charges for

1 out-of-network providers, hospitals, or facilities that were
2 incorrectly listed as in-network prior to the provision of
3 care and a telephone number and email address to dispute those
4 charges.

5 (f) The Director may conduct periodic audits of the
6 accuracy of provider directories. A network plan shall not be
7 subject to any fines or penalties for information required in
8 this Section that a provider submits that is inaccurate or
9 incomplete.

10 (g) To the extent not otherwise provided in this Act, an
11 issuer shall comply with the requirements of 42 U.S.C.
12 300gg-115, except that "provider directory information" shall
13 include all information required to be included in a provider
14 directory pursuant to this Section.

15 (h) If the issuer or the Department identifies a provider
16 incorrectly listed in the provider directory, the issuer shall
17 do all of the following:

18 (1) Check each of the issuer's network plan provider
19 directories for the provider within 2 business days to
20 ascertain whether the provider is a preferred provider in
21 that network plan and, if the provider is incorrectly
22 listed in the directory, remove the provider without
23 delay.

24 (2) Identify the dates across each of the issuer's
25 network plan provider directories that the provider was
26 listed when the provider was not a preferred provider.

1 (3) For covered services furnished by the provider
2 during the period the provider was incorrectly listed in
3 the network directory, identify all claims that have been
4 paid, are pending, or, for a network plan that does not
5 require a referral for in-network covered services
6 rendered by that type of provider, have been denied as
7 out-of-network. For claims that a beneficiary submits to
8 the issuer for reimbursement, the issuer shall reimburse
9 or supplement a prior reimbursement to the beneficiary in
10 the amount necessary to ensure the beneficiary is held
11 harmless for all billed amounts for covered services that
12 exceed the in-network cost-sharing amount for the covered
13 services. For claims that the issuer pays directly to the
14 provider, the issuer shall notify the provider and the
15 beneficiary in writing of the beneficiary's right to
16 reimbursement from the provider for any payments in excess
17 of the in-network cost-sharing amount pursuant to 42
18 U.S.C. 300gg-139(b), and the issuer's notice shall specify
19 the in-network cost-sharing amount for the covered
20 services. All out-of-pocket costs incurred by the
21 beneficiary within the in-network cost-sharing amount
22 shall apply toward the in-network deductible and
23 out-of-pocket maximum.

24 (4) For each beneficiary who had an in-network claim
25 for services from the incorrectly included provider during
26 the year prior to the date that the provider ceased to

1 participate in the network plan, send a written
2 communication to the beneficiary of the inaccurate
3 provider listing, including the dates thereof, and the
4 beneficiary's right to reimbursement by the issuer or, if
5 the issuer paid the claim to the provider directly,
6 reimbursement by the provider, for any costs incurred
7 incorrectly in excess of the in-network cost-sharing on
8 the dates that the provider was incorrectly listed as
9 in-network in the provider directory.

10 (i) Issuers must maintain a copy of each network plan's
11 provider directory for a minimum of 5 years from the date of
12 publication and make it available to beneficiaries and the
13 Department upon request and at no cost.

14 (j) If an issuer fails to provide notice to beneficiaries
15 of a nonrenewal or termination of a provider in accordance
16 with Section 15 and that nonrenewal or termination occurs,
17 services delivered by the provider shall be reimbursed to the
18 beneficiary as if the provider were in-network until the
19 requirements, including the notice period of Section 15, have
20 been met. For claims that a beneficiary submits to the issuer
21 for reimbursement, the issuer shall reimburse or supplement a
22 prior reimbursement to the beneficiary in the amount necessary
23 to ensure the beneficiary is held harmless for all billed
24 amounts for covered services that exceed the in-network
25 cost-sharing amount for the covered services. For claims that
26 the issuer pays directly to the provider, the issuer shall

1 notify the provider and the beneficiary of the in-network
2 cost-sharing amount for the covered services, and the provider
3 shall hold harmless and reimburse the beneficiary for all
4 payments in excess of that amount. The amounts paid by the
5 beneficiary shall apply towards the in-network deductible and
6 out-of-pocket maximum, if any.

7 (k) If the Director determines that an issuer violated
8 this Section, the Director may assess a fine up to \$5,000 per
9 violation, except for inaccurate information given by a
10 provider to the issuer. If an issuer, or any entity or person
11 acting on the issuer's behalf, knew or reasonably should have
12 known that a provider was incorrectly included in a provider
13 directory, the Director may assess a fine of up to \$25,000 per
14 violation against the issuer.

15 (l) This Section applies to network plans not otherwise
16 exempt under Section 3, including stand-alone dental plans.

17 (Source: P.A. 102-92, eff. 7-9-21; revised 9-26-23.)

18 (215 ILCS 124/30)

19 Sec. 30. Administration and enforcement.

20 (a) Issuers ~~Insurers~~, as defined in this Act, have a
21 continuing obligation to comply with the requirements of this
22 Act. Other than the duties specifically created in this Act,
23 nothing in this Act is intended to preclude, prevent, or
24 require the adoption, modification, or termination of any
25 utilization management, quality management, or claims

1 processing methodologies of an issuer ~~insurer~~.

2 (b) Nothing in this Act precludes, prevents, or requires
3 the adoption, modification, or termination of any network plan
4 term, benefit, coverage or eligibility provision, or payment
5 methodology.

6 (c) The Director shall enforce the provisions of this Act
7 pursuant to the enforcement powers granted to it by law.

8 (d) The Department shall adopt rules to enforce compliance
9 with this Act to the extent necessary.

10 (e) In accordance with Section 5-45 of the Illinois
11 Administrative Procedure Act, the Department may adopt
12 emergency rules to implement federal standards for provider
13 ratios, travel time and distance, and appointment wait times
14 if such standards apply to health insurance coverage regulated
15 by the Department and are more stringent than the State
16 standards extant at the time the final federal standards are
17 published.

18 (Source: P.A. 100-502, eff. 9-15-17.)

19 (215 ILCS 124/35 new)

20 Sec. 35. Provider requirements. Providers shall comply
21 with 42 U.S.C. 300gg-138 and 300gg-139 and the regulations
22 promulgated thereunder, as well as Section 20, paragraph (2)
23 of subsection (a) of Section 25, subsections (h) and (j) of
24 Section 25, and Section 36 of this Act, except that "provider
25 directory information" includes all information required to be

1 included in a provider directory pursuant to Section 25 of
2 this Act.

3 (215 ILCS 124/36 new)

4 Sec. 36. Complaint of incorrect charges.

5 (a) A beneficiary who, taking into account the
6 reimbursement, if any, by the issuer, incurs a cost in excess
7 of the in-network cost-sharing for a covered service from a
8 provider, facility, or hospital that was listed as in-network
9 in the plan's provider directory prior to or at the time of the
10 provision of services may file a complaint with the
11 Department. The Department shall investigate the complaint and
12 determine if the provider was incorrectly included in the
13 plan's provider directory when the beneficiary made the
14 appointment or received the service.

15 (b) Upon the Department's confirmation of the allegations
16 in the complaint that the beneficiary incurred a cost in
17 excess of the in-network cost-sharing for covered services
18 provided by an incorrectly included provider when the
19 appointment was made or service was provided, the issuer shall
20 reimburse the beneficiary for all costs incurred in excess of
21 the in-network cost-sharing. However, if the issuer has paid
22 the claim to the provider directly, the issuer shall notify
23 the beneficiary and the provider of the beneficiary's right to
24 reimbursement from the provider for any payments in excess of
25 the in-network cost-sharing amount pursuant to 42 U.S.C.

1 300gg-139(b), and the issuer's notice shall specify the
2 in-network cost-sharing amount for the covered services. The
3 amounts paid by the beneficiary within the in-network
4 cost-sharing amount shall apply towards the in-network
5 deductible and out-of-pocket maximum, if any.

6 (215 ILCS 124/40 new)

7 Sec. 40. Confidentiality.

8 (a) All records in the custody or possession of the
9 Department are presumed to be open to public inspection or
10 copying unless exempt from disclosure by Section 7 or 7.5 of
11 the Freedom of Information Act. Except as otherwise provided
12 in this Section or other applicable law, the filings required
13 under this Act shall be open to public inspection or copying.

14 (b) The following information shall not be deemed
15 confidential:

16 (1) actual or projected ratios of providers to
17 beneficiaries;

18 (2) actual or projected time and distance between
19 network providers and beneficiaries or actual or projected
20 waiting times for a beneficiary to see a network provider;

21 (3) geographic maps of network providers;

22 (4) requests for exceptions under subsection (g) of
23 Section 10, except with respect to any discussion of
24 ongoing or planned contractual negotiations with providers
25 that the issuer requests to be treated as confidential;

1 (5) provider directories and provider lists;

2 (6) self-audit summaries required under paragraph (3)
3 of subsection (a) of Section 25 of this Act; and

4 (7) issuer or Department statements of determination
5 as to whether a network plan has satisfied this Act's
6 requirements regarding the information described in this
7 subsection.

8 (c) An issuer's work papers and reports on the results of a
9 self-audit of its provider directories, including any
10 communications between the issuer and the Department, shall
11 remain confidential unless expressly waived by the issuer or
12 unless deemed public information under federal law.

13 (d) The filings required under Section 10 of this Act
14 shall be confidential while they remain under the Department's
15 review but shall become open to public inspection and copying
16 upon completion of the review, except as provided in this
17 Section or under other applicable law.

18 (e) Nothing in this Section shall supersede the statutory
19 requirement that work papers obtained during a market conduct
20 examination be deemed confidential.

21 (215 ILCS 124/50 new)

22 Sec. 50. Funds for enforcement. Moneys from fines and
23 penalties collected from issuers for violations of this Act
24 shall be deposited into the Insurance Producer Administration
25 Fund for appropriation by the General Assembly to the

1 Department to be used for providing financial support of the
2 Department's enforcement of this Act.

3 (215 ILCS 124/55 new)

4 Sec. 55. Uniform electronic provider directory information
5 notification forms.

6 (a) On or before January 1, 2026, the Department shall
7 develop and publish a uniform electronic provider directory
8 information form that issuers shall make available to
9 onboarding, current, and former preferred providers to notify
10 the issuer of the provider's currently accurate provider
11 directory information under Section 25 of this Act and 42
12 U.S.C. 300gg-139. The form shall address information needed
13 from newly onboarding preferred providers, updates to
14 previously supplied provider directory information, reporting
15 an inaccurate directory entry of previously supplied
16 information, contract terminations, and differences in
17 information for specific network plans offered by an issuer,
18 such as whether the provider is a preferred provider for the
19 network plan or is accepting new patients under that plan. The
20 Department shall allow issuers to implement this form through
21 either a PDF or a web portal that requests the same
22 information.

23 (b) Notwithstanding any other provision of law to the
24 contrary, beginning 6 months after the Department publishes
25 the uniform electronic provider directory information form and

1 no later than July 1, 2026, every provider must use the uniform
2 electronic provider directory information form to notify
3 issuers of their provider directory information as required
4 under Section 25 of this Act and 42 U.S.C. 300gg-139. Issuers
5 shall accept this form as sufficient to update their provider
6 directories. Issuers shall not accept paper or fax submissions
7 of provider directory information from providers.

8 (c) The Uniform Electronic Provider Directory Information
9 Form Task Force is created. The purpose of this task force is
10 to provide input and advice to the Department of Insurance in
11 the development of a uniform electronic provider directory
12 information form. The task force shall include at least the
13 following individuals:

14 (1) the Director of Insurance or a designee, as chair;

15 (2) the Marketplace Director or a designee;

16 (3) the Director of the Division of Professional
17 Regulation or a designee;

18 (4) the Director of Public Health or a designee;

19 (5) the Secretary of Innovation and Technology or a
20 designee;

21 (6) the Director of Healthcare and Family Services or
22 a designee;

23 (7) the following individuals appointed by the
24 Director:

25 (A) one representative of a statewide association
26 representing physicians;

1 (B) one representative of a statewide association
2 representing nurses;

3 (C) one representative of a statewide organization
4 representing a majority of Illinois hospitals;

5 (D) one representative of a statewide organization
6 representing Illinois pharmacies;

7 (E) one representative of a statewide organization
8 representing mental health care providers;

9 (F) one representative of a statewide organization
10 representing substance use disorder health care
11 providers;

12 (G) 2 representatives of health insurance issuers
13 doing business in this State or issuer trade
14 associations, at least one of which represents a
15 State-domiciled mutual health insurance company, with
16 a demonstrated expertise in the business of health
17 insurance or health benefits administration; and

18 (H) 2 representatives of a health insurance
19 consumer advocacy group.

20 (d) The Department shall convene the task force described
21 in this Section no later than April 1, 2025.

22 (e) The Department, in development of the uniform
23 electronic provider directory information form, and the task
24 force, in offering input, shall take into consideration the
25 following:

26 (1) readability and user experience;

1 (2) interoperability;

2 (3) existing regulations established by the federal
3 Centers for Medicare and Medicaid Services, the Department
4 of Insurance, the Department of Healthcare and Family
5 Service, the Department of Financial and Professional
6 Regulation, and the Department of Public Health;

7 (4) potential opportunities to avoid duplication of
8 data collection efforts, including, but not limited to,
9 opportunities related to:

10 (A) integrating any provider reporting required
11 under Section 25 of this Act and 42 U.S.C. 300gg-139
12 with the provider reporting required under the Health
13 Care Professional Credentials Data Collection Act;

14 (B) furnishing information to any national
15 provider directory established by the federal Centers
16 for Medicare and Medicaid Services or another federal
17 agency with jurisdiction over health care providers;
18 and

19 (C) furnishing information in compliance with the
20 Patients' Right to Know Act;

21 (5) compatibility with the Illinois Health Benefits
22 Exchange;

23 (6) provider licensing requirements and forms; and

24 (7) information needed to classify a provider under
25 any specialty type for which a network adequacy standard
26 may be established under this Act when a specialty board

1 certification or State license does not currently exist.

2 Section 2-15. The Managed Care Reform and Patient Rights
3 Act is amended by changing Sections 20 and 25 as follows:

4 (215 ILCS 134/20)

5 Sec. 20. Notice of nonrenewal or termination. A health
6 care plan must give at least 60 days notice of nonrenewal or
7 termination of a health care provider to the health care
8 provider and to the enrollees served by the health care
9 provider. The notice shall include a name and address to which
10 an enrollee or health care provider may direct comments and
11 concerns regarding the nonrenewal or termination. Immediate
12 written notice may be provided without 60 days notice when a
13 health care provider's license has been disciplined by a State
14 licensing board. The notice to the enrollee shall provide the
15 individual with an opportunity to notify the health care plan
16 of the individual's need for transitional care.

17 (Source: P.A. 91-617, eff. 1-1-00.)

18 (215 ILCS 134/25)

19 Sec. 25. Transition of services.

20 (a) A health care plan shall provide for continuity of
21 care for its enrollees as follows:

22 (1) If an enrollee's health care provider ~~physician~~
23 leaves the health care plan's network of health care

1 providers for reasons other than termination of a contract
2 in situations involving imminent harm to a patient or a
3 final disciplinary action by a State licensing board and
4 the provider ~~physician~~ remains within the health care
5 plan's service area, or if benefits provided under such
6 health care plan with respect to such provider are
7 terminated because of a change in the terms of the
8 participation of such provider in such plan, or if a
9 contract between a group health plan, as defined in
10 Section 5 of the Illinois Health Insurance Portability and
11 Accountability Act, and a health care plan offered in
12 connection with the group health plan is terminated and
13 results in a loss of benefits provided under such plan
14 with respect to such provider, the health care plan shall
15 permit the enrollee to continue an ongoing course of
16 treatment with that provider ~~physician~~ during a
17 transitional period:

18 (A) of 90 days from the date of the notice of
19 provider's ~~physician's~~ termination from the health
20 care plan to the enrollee of the provider's
21 ~~physician's~~ disaffiliation from the health care plan
22 if the enrollee has an ongoing course of treatment; or

23 (B) if the enrollee has entered the third
24 trimester of pregnancy at the time of the provider's
25 ~~physician's~~ disaffiliation, that includes the
26 provision of post-partum care directly related to the

1 delivery.

2 (2) Notwithstanding the provisions in item (1) of this
3 subsection, such care shall be authorized by the health
4 care plan during the transitional period only if the
5 provider ~~physician~~ agrees:

6 (A) to continue to accept reimbursement from the
7 health care plan at the rates applicable prior to the
8 start of the transitional period;

9 (B) to adhere to the health care plan's quality
10 assurance requirements and to provide to the health
11 care plan necessary medical information related to
12 such care; and

13 (C) to otherwise adhere to the health care plan's
14 policies and procedures, including but not limited to
15 procedures regarding referrals and obtaining
16 preauthorizations for treatment.

17 (3) During an enrollee's plan year, a health care plan
18 shall not remove a drug from its formulary or negatively
19 change its preferred or cost-tier sharing unless, at least
20 60 days before making the formulary change, the health
21 care plan:

22 (A) provides general notification of the change in
23 its formulary to current and prospective enrollees;

24 (B) directly notifies enrollees currently
25 receiving coverage for the drug, including information
26 on the specific drugs involved and the steps they may

1 take to request coverage determinations and
2 exceptions, including a statement that a certification
3 of medical necessity by the enrollee's prescribing
4 provider will result in continuation of coverage at
5 the existing level; and

6 (C) directly notifies in writing ~~by first class~~
7 ~~mail and~~ through an electronic transmission, ~~if~~
8 ~~available,~~ the prescribing provider of all health care
9 plan enrollees currently prescribed the drug affected
10 by the proposed change; the notice shall include a
11 one-page form by which the prescribing provider can
12 notify the health care plan in writing or
13 electronically ~~by first class mail~~ that coverage of
14 the drug for the enrollee is medically necessary.

15 The notification in paragraph (C) may direct the
16 prescribing provider to an electronic portal through which
17 the prescribing provider may electronically file a
18 certification to the health care plan that coverage of the
19 drug for the enrollee is medically necessary. The
20 prescribing provider may make a secure electronic
21 signature beside the words "certification of medical
22 necessity", and this certification shall authorize
23 continuation of coverage for the drug.

24 If the prescribing provider certifies to the health
25 care plan either in writing or electronically that the
26 drug is medically necessary for the enrollee as provided

1 in paragraph (C), a health care plan shall authorize
2 coverage for the drug prescribed based solely on the
3 prescribing provider's assertion that coverage is
4 medically necessary, and the health care plan is
5 prohibited from making modifications to the coverage
6 related to the covered drug, including, but not limited
7 to:

8 (i) increasing the out-of-pocket costs for the
9 covered drug;

10 (ii) moving the covered drug to a more restrictive
11 tier; or

12 (iii) denying an enrollee coverage of the drug for
13 which the enrollee has been previously approved for
14 coverage by the health care plan.

15 Nothing in this item (3) prevents a health care plan
16 from removing a drug from its formulary or denying an
17 enrollee coverage if the United States Food and Drug
18 Administration has issued a statement about the drug that
19 calls into question the clinical safety of the drug, the
20 drug manufacturer has notified the United States Food and
21 Drug Administration of a manufacturing discontinuance or
22 potential discontinuance of the drug as required by
23 Section 506C of the Federal Food, Drug, and Cosmetic Act,
24 as codified in 21 U.S.C. 356c, or the drug manufacturer
25 has removed the drug from the market.

26 Nothing in this item (3) prohibits a health care plan,

1 by contract, written policy or procedure, or any other
2 agreement or course of conduct, from requiring a
3 pharmacist to effect substitutions of prescription drugs
4 consistent with Section 19.5 of the Pharmacy Practice Act,
5 under which a pharmacist may substitute an interchangeable
6 biologic for a prescribed biologic product, and Section 25
7 of the Pharmacy Practice Act, under which a pharmacist may
8 select a generic drug determined to be therapeutically
9 equivalent by the United States Food and Drug
10 Administration and in accordance with the Illinois Food,
11 Drug and Cosmetic Act.

12 This item (3) applies to a policy or contract that is
13 amended, delivered, issued, or renewed on or after January
14 1, 2019. This item (3) does not apply to a health plan as
15 defined in the State Employees Group Insurance Act of 1971
16 or medical assistance under Article V of the Illinois
17 Public Aid Code.

18 (b) A health care plan shall provide for continuity of
19 care for new enrollees as follows:

20 (1) If a new enrollee whose physician is not a member
21 of the health care plan's provider network, but is within
22 the health care plan's service area, enrolls in the health
23 care plan, the health care plan shall permit the enrollee
24 to continue an ongoing course of treatment with the
25 enrollee's current physician during a transitional period:

26 (A) of 90 days from the effective date of

1 enrollment if the enrollee has an ongoing course of
2 treatment; or

3 (B) if the enrollee has entered the third
4 trimester of pregnancy at the effective date of
5 enrollment, that includes the provision of post-partum
6 care directly related to the delivery.

7 (2) If an enrollee elects to continue to receive care
8 from such physician pursuant to item (1) of this
9 subsection, such care shall be authorized by the health
10 care plan for the transitional period only if the
11 physician agrees:

12 (A) to accept reimbursement from the health care
13 plan at rates established by the health care plan;
14 such rates shall be the level of reimbursement
15 applicable to similar physicians within the health
16 care plan for such services;

17 (B) to adhere to the health care plan's quality
18 assurance requirements and to provide to the health
19 care plan necessary medical information related to
20 such care; and

21 (C) to otherwise adhere to the health care plan's
22 policies and procedures including, but not limited to
23 procedures regarding referrals and obtaining
24 preauthorization for treatment.

25 (c) In no event shall this Section be construed to require
26 a health care plan to provide coverage for benefits not

1 otherwise covered or to diminish or impair preexisting
2 condition limitations contained in the enrollee's contract. In
3 no event shall this Section be construed to prohibit the
4 addition of prescription drugs to a health care plan's list of
5 covered drugs during the coverage year.

6 (d) In this Section, "ongoing course of treatment" has the
7 meaning ascribed to that term in Section 5 of the Network
8 Adequacy and Transparency Act.

9 (Source: P.A. 100-1052, eff. 8-24-18.)

10 Article 3.

11 Section 3-5. The Illinois Insurance Code is amended by
12 changing Section 355 as follows:

13 (215 ILCS 5/355) (from Ch. 73, par. 967)

14 Sec. 355. Accident and health policies; provisions.

15 (a) As used in this Section:

16 "Inadequate rate" means a rate:

17 (1) that is insufficient to sustain projected losses
18 and expenses to which the rate applies; and

19 (2) the continued use of which endangers the solvency
20 of an insurer using that rate.

21 "Large employer" has the meaning provided in the Illinois
22 Health Insurance Portability and Accountability Act.

23 "Plain language" has the meaning provided in the federal

1 Plain Writing Act of 2010 and subsequent guidance documents,
2 including the Federal Plain Language Guidelines.

3 "Unreasonable rate increase" means a rate increase that
4 the Director determines to be excessive, unjustified, or
5 unfairly discriminatory in accordance with 45 CFR 154.205.

6 (b) No policy of insurance against loss or damage from the
7 sickness, or from the bodily injury or death of the insured by
8 accident shall be issued or delivered to any person in this
9 State until a copy of the form thereof and of the
10 classification of risks and the premium rates pertaining
11 thereto have been filed with the Director; nor shall it be so
12 issued or delivered until the Director shall have approved
13 such policy pursuant to the provisions of Section 143. If the
14 Director disapproves the policy form, he or she shall make a
15 written decision stating the respects in which such form does
16 not comply with the requirements of law and shall deliver a
17 copy thereof to the company and it shall be unlawful
18 thereafter for any such company to issue any policy in such
19 form. On and after January 1, 2025, any form filing submitted
20 for large employer group accident and health insurance shall
21 be automatically deemed approved within 90 days of the
22 submission date unless the Director extends by not more than
23 an additional 30 days the period within which the form shall be
24 approved or disapproved by giving written notice to the
25 insurer of such extension before the expiration of the 90
26 days. Any form in receipt of such an extension shall be

1 automatically deemed approved within 120 days of the
2 submission date. The Director may toll the filing due to a
3 conflict in legal interpretation of federal or State law as
4 long as the tolling is applied uniformly to all applicable
5 forms, written notification is provided to the insurer prior
6 to the tolling, the duration of the tolling is provided within
7 the notice to the insurer, and justification for the tolling
8 is posted to the Department's website. The Director may
9 disapprove the filing if the insurer fails to respond to an
10 objection or request for additional information within the
11 timeframe identified for response. As used in this subsection,
12 "large employer" has the meaning given in Section 5 of the
13 federal Health Insurance Portability and Accountability Act.

14 (c) For plan year 2026 and thereafter, premium rates for
15 all individual and small group accident and health insurance
16 policies must be filed with the Department for approval.
17 Unreasonable rate increases or inadequate rates shall be
18 modified or disapproved. For any plan year during which the
19 Illinois Health Benefits Exchange operates as a full
20 State-based exchange, the Department shall provide insurers at
21 least 30 days' notice of the deadline to submit rate filings.

22 (c-5) Unless prohibited under federal law, for plan year
23 2026 and thereafter, each insurer proposing to offer a
24 qualified health plan issued in the individual market through
25 the Illinois Health Benefits Exchange must incorporate the
26 following approach in its rate filing under this Section:

1 (1) The rate filing must apply a cost-sharing
2 reduction defunding adjustment factor within a range that:

3 (A) is uniform across all insurers;

4 (B) is consistent with the total adjustment
5 expected to be needed to cover actual cost-sharing
6 reduction costs across all silver plans on the
7 Illinois Health Benefits Exchange statewide, provided
8 that such costs are calculated assuming utilization by
9 the State's full individual-market risk pool; and

10 (C) assumes that the only on-Exchange silver plans
11 that will be purchased are the 87% and 94%
12 cost-sharing reduction variations.

13 (2) The rate filing must apply an induced demand
14 factor based on the following formula: (Plan Actuarial
15 Value)² - (Plan Actuarial Value) + 1.24.

16 In the annual notice to insurers described in subsection
17 (c), the Department must include the specific numerical range
18 calculated for the applicable plan year under paragraph (1) of
19 this subsection (c-5) and the formula in paragraph (2) of this
20 subsection (c-5).

21 (d) For plan year 2025 and thereafter, the Department
22 shall post all insurers' rate filings and summaries on the
23 Department's website 5 business days after the rate filing
24 deadline set by the Department in annual guidance. The rate
25 filings and summaries posted to the Department's website shall
26 exclude information that is proprietary or trade secret

1 information protected under paragraph (g) of subsection (1) of
2 Section 7 of the Freedom of Information Act or confidential or
3 privileged under any applicable insurance law or rule. All
4 summaries shall include a brief justification of any rate
5 increase or decrease requested, including the number of
6 individual members, the medical loss ratio, medical trend,
7 administrative costs, and any other information required by
8 rule. The plain writing summary shall include notification of
9 the public comment period established in subsection (e).

10 (e) The Department shall open a 30-day public comment
11 period on the rate filings beginning on the date that all of
12 the rate filings are posted on the Department's website. The
13 Department shall post all of the comments received to the
14 Department's website within 5 business days after the comment
15 period ends.

16 (f) After the close of the public comment period described
17 in subsection (e), the Department, beginning for plan year
18 2026, shall issue a decision to approve, disapprove, or modify
19 a rate filing within 60 days. Any rate filing or any rates
20 within a filing on which the Director does not issue a decision
21 within 60 days shall automatically be deemed approved. The
22 Director's decision shall take into account the actuarial
23 justifications and public comments. The Department shall
24 notify the insurer of the decision, make the decision
25 available to the public by posting it on the Department's
26 website, and include an explanation of the findings, actuarial

1 justifications, and rationale that are the basis for the
2 decision. Any company whose rate has been modified or
3 disapproved shall be allowed to request a hearing within 10
4 days after the action taken. The action of the Director in
5 disapproving a rate shall be subject to judicial review under
6 the Administrative Review Law.

7 (g) If, following the issuance of a decision but before
8 the effective date of the premium rates approved by the
9 decision, an event occurs that materially affects the
10 Director's decision to approve, deny, or modify the rates, the
11 Director may consider supplemental facts or data reasonably
12 related to the event.

13 (h) The Department shall adopt rules implementing the
14 procedures described in subsections (d) through (g) by March
15 31, 2024.

16 (i) Subsection (a) and subsections (c) through (h) of this
17 Section do not apply to grandfathered health plans as defined
18 in 45 CFR 147.140; excepted benefits as defined in 42 U.S.C.
19 300gg-91; student health insurance coverage as defined in 45
20 CFR 147.145; the large group market as defined in Section 5 of
21 the Illinois Health Insurance Portability and Accountability
22 Act; or short-term, limited-duration health insurance coverage
23 as defined in Section 5 of the Short-Term, Limited-Duration
24 Health Insurance Coverage Act. For a filing of premium rates
25 or classifications of risk for any of these types of coverage,
26 the Director's initial review period shall not exceed 60 days

1 to issue informal objections to the company that request
2 additional clarification, explanation, substantiating
3 documentation, or correction of concerns identified in the
4 filing before the company implements the premium rates,
5 classifications, or related rate-setting methodologies
6 described in the filing, except that the Director may extend
7 by not more than an additional 30 days the period of initial
8 review by giving written notice to the company of such
9 extension before the expiration of the initial 60-day period.
10 Nothing in this subsection shall confer authority upon the
11 Director to approve, modify, or disapprove rates where that
12 authority is not provided by other law. Nothing in this
13 subsection shall prohibit the Director from conducting any
14 investigation, examination, hearing, or other formal
15 administrative or enforcement proceeding with respect to a
16 company's rate filing or implementation thereof under
17 applicable law at any time, including after the period of
18 initial review.

19 (Source: P.A. 103-106, eff. 1-1-24.)

20 Section 3-10. The Illinois Health Benefits Exchange Law is
21 amended by changing Section 5-5 as follows:

22 (215 ILCS 122/5-5)

23 Sec. 5-5. State health benefits exchange. It is declared
24 that this State, beginning October 1, 2013, in accordance with

1 Section 1311 of the federal Patient Protection and Affordable
2 Care Act, shall establish a State health benefits exchange to
3 be known as the Illinois Health Benefits Exchange in order to
4 help individuals and small employers with no more than 50
5 employees shop for, select, and enroll in qualified,
6 affordable private health plans that fit their needs at
7 competitive prices. The Exchange shall separate coverage pools
8 for individuals and small employers and shall supplement and
9 not supplant any existing private health insurance market for
10 individuals and small employers. The Department of Insurance
11 shall operate the Illinois Health Benefits Exchange as a
12 State-based exchange using the federal platform by plan year
13 2025 and as a State-based exchange by plan year 2026. The
14 Director of Insurance may require that all plans in the
15 individual and small group markets, other than grandfathered
16 health plans, be made available for comparison on the Illinois
17 Health Benefits Exchange, but may not require that all plans
18 in the individual and small group markets be purchased
19 exclusively on the Illinois Health Benefits Exchange. Through
20 the adoption of rules, the Director of Insurance may require
21 that plans offered on the exchange conform with standardized
22 plan designs that provide for standardized cost sharing for
23 covered health services. Except when it is inconsistent with
24 State law, the Department of Insurance shall enforce the
25 coverage requirements under the federal Patient Protection and
26 Affordable Care Act, including the coverage of all United

1 States Preventive Services Task Force Grade A and B preventive
2 services without cost sharing notwithstanding any federal
3 overturning or repeal of 42 U.S.C. 300gg-13(a)(1), that apply
4 to the individual and small group markets. Beginning for plan
5 year 2026, if a health insurance issuer offers a product as
6 defined under 45 CFR 144.103 at the gold or silver level
7 through the Illinois Health Benefits Exchange, the issuer must
8 offer that product at both the gold and silver levels. The
9 Director of Insurance may elect to add a small business health
10 options program to the Illinois Health Benefits Exchange to
11 help small employers enroll their employees in qualified
12 health plans in the small group market. The General Assembly
13 shall appropriate funds to establish the Illinois Health
14 Benefits Exchange.

15 (Source: P.A. 103-103, eff. 6-27-23.)

16 Article 4.

17 Section 4-5. The Illinois Insurance Code is amended by
18 changing Section 355 as follows:

19 (215 ILCS 5/355) (from Ch. 73, par. 967)

20 Sec. 355. Accident and health policies; provisions.

21 (a) As used in this Section:

22 "Inadequate rate" means a rate:

23 (1) that is insufficient to sustain projected losses

1 and expenses to which the rate applies; and

2 (2) the continued use of which endangers the solvency
3 of an insurer using that rate.

4 "Large employer" has the meaning provided in the Illinois
5 Health Insurance Portability and Accountability Act.

6 "Plain language" has the meaning provided in the federal
7 Plain Writing Act of 2010 and subsequent guidance documents,
8 including the Federal Plain Language Guidelines.

9 "Unreasonable rate increase" means a rate increase that
10 the Director determines to be excessive, unjustified, or
11 unfairly discriminatory in accordance with 45 CFR 154.205.

12 (b) No policy of insurance against loss or damage from the
13 sickness, or from the bodily injury or death of the insured by
14 accident shall be issued or delivered to any person in this
15 State until a copy of the form thereof and of the
16 classification of risks and the premium rates pertaining
17 thereto have been filed with the Director; nor shall it be so
18 issued or delivered until the Director shall have approved
19 such policy pursuant to the provisions of Section 143. If the
20 Director disapproves the policy form, he or she shall make a
21 written decision stating the respects in which such form does
22 not comply with the requirements of law and shall deliver a
23 copy thereof to the company and it shall be unlawful
24 thereafter for any such company to issue any policy in such
25 form. On and after January 1, 2025, any form filing submitted
26 for large employer group accident and health insurance shall

1 be automatically deemed approved within 90 days of the
2 submission date unless the Director extends by not more than
3 an additional 30 days the period within which the form shall be
4 approved or disapproved by giving written notice to the
5 insurer of such extension before the expiration of the 90
6 days. Any form in receipt of such an extension shall be
7 automatically deemed approved within 120 days of the
8 submission date. The Director may toll the filing due to a
9 conflict in legal interpretation of federal or State law as
10 long as the tolling is applied uniformly to all applicable
11 forms, written notification is provided to the insurer prior
12 to the tolling, the duration of the tolling is provided within
13 the notice to the insurer, and justification for the tolling
14 is posted to the Department's website. The Director may
15 disapprove the filing if the insurer fails to respond to an
16 objection or request for additional information within the
17 timeframe identified for response. As used in this subsection,
18 "large employer" has the meaning given in Section 5 of the
19 federal Health Insurance Portability and Accountability Act.

20 (c) For plan year 2026 and thereafter, premium rates for
21 all individual and small group accident and health insurance
22 policies must be filed with the Department for approval.
23 Unreasonable rate increases or inadequate rates shall be
24 modified or disapproved. For any plan year during which the
25 Illinois Health Benefits Exchange operates as a full
26 State-based exchange, the Department shall provide insurers at

1 least 30 days' notice of the deadline to submit rate filings.

2 (d) For plan year 2025 and thereafter, the Department
3 shall post all insurers' rate filings and summaries on the
4 Department's website 5 business days after the rate filing
5 deadline set by the Department in annual guidance. The rate
6 filings and summaries posted to the Department's website shall
7 exclude information that is proprietary or trade secret
8 information protected under paragraph (g) of subsection (1) of
9 Section 7 of the Freedom of Information Act or confidential or
10 privileged under any applicable insurance law or rule. All
11 summaries shall include a brief justification of any rate
12 increase or decrease requested, including the number of
13 individual members, the medical loss ratio, medical trend,
14 administrative costs, and any other information required by
15 rule. The plain writing summary shall include notification of
16 the public comment period established in subsection (e).

17 (e) The Department shall open a 30-day public comment
18 period on the rate filings beginning on the date that all of
19 the rate filings are posted on the Department's website. The
20 Department shall post all of the comments received to the
21 Department's website within 5 business days after the comment
22 period ends.

23 (f) After the close of the public comment period described
24 in subsection (e), the Department, beginning for plan year
25 2026, shall issue a decision to approve, disapprove, or modify
26 a rate filing within 60 days. Any rate filing or any rates

1 within a filing on which the Director does not issue a decision
2 within 60 days shall automatically be deemed approved. The
3 Director's decision shall take into account the actuarial
4 justifications and public comments. The Department shall
5 notify the insurer of the decision, make the decision
6 available to the public by posting it on the Department's
7 website, and include an explanation of the findings, actuarial
8 justifications, and rationale that are the basis for the
9 decision. Any company whose rate has been modified or
10 disapproved shall be allowed to request a hearing within 10
11 days after the action taken. The action of the Director in
12 disapproving a rate shall be subject to judicial review under
13 the Administrative Review Law.

14 (g) If, following the issuance of a decision but before
15 the effective date of the premium rates approved by the
16 decision, an event occurs that materially affects the
17 Director's decision to approve, deny, or modify the rates, the
18 Director may consider supplemental facts or data reasonably
19 related to the event.

20 (h) The Department shall adopt rules implementing the
21 procedures described in subsections (d) through (g) by March
22 31, 2024.

23 (i) Subsection (a), ~~and~~ subsections (c) through (h), and
24 subsection (j) of this Section do not apply to grandfathered
25 health plans as defined in 45 CFR 147.140; excepted benefits
26 as defined in 42 U.S.C. 300gg-91; or student health insurance

1 coverage as defined in 45 CFR 147.145, ~~the large group market~~
2 ~~as defined in Section 5 of the Illinois Health Insurance~~
3 ~~Portability and Accountability Act; or short term,~~
4 ~~limited duration health insurance coverage as defined in~~
5 ~~Section 5 of the Short Term, Limited Duration Health Insurance~~
6 ~~Coverage Act.~~ For a filing of premium rates or classifications
7 of risk for any of these types of coverage, the Director's
8 initial review period shall not exceed 60 days to issue
9 informal objections to the company that request additional
10 clarification, explanation, substantiating documentation, or
11 correction of concerns identified in the filing before the
12 company implements the premium rates, classifications, or
13 related rate-setting methodologies described in the filing,
14 except that the Director may extend by not more than an
15 additional 30 days the period of initial review by giving
16 written notice to the company of such extension before the
17 expiration of the initial 60-day period. Nothing in this
18 subsection shall confer authority upon the Director to
19 approve, modify, or disapprove rates where that authority is
20 not provided by other law. Nothing in this subsection shall
21 prohibit the Director from conducting any investigation,
22 examination, hearing, or other formal administrative or
23 enforcement proceeding with respect to a company's rate filing
24 or implementation thereof under applicable law at any time,
25 including after the period of initial review.

26 (j) Subsections (c) through (h) do not apply to group

1 policies issued to large employers. For large employer group
2 policies issued, delivered, amended, or renewed on or after
3 January 1, 2026 that are not described in subsection (i), the
4 premium rates and risk classifications, including any rate
5 manuals and rules used to arrive at the rates, must be filed
6 with the Department annually for approval at least 120 days
7 before the rates are intended to take effect.

8 (1) A rate filing shall be modified or disapproved if
9 rates will be unreasonable in relation to the benefits,
10 unjustified, or unfairly discriminatory, or otherwise in
11 violation of applicable State or federal law.

12 (2) Within 60 days of receipt of the rate filing, the
13 Director shall issue a decision to approve, disapprove, or
14 modify the filing along with the reasons and actuarial
15 justification for the decision. Any rate filing or rates
16 within a filing on which the Director does not issue a
17 decision within 60 days shall be automatically deemed
18 approved.

19 (3) Any company whose rate or rate filing has been
20 modified or disapproved shall be allowed to request a
21 hearing within 10 days after the action taken. The action
22 of the Director in disapproving a rate or rate filing
23 shall be subject to judicial review under the
24 Administrative Review Law.

25 (4) Nothing in this subsection requires a company to
26 file a large employer group policy's final premium rates

1 for prior approval if the company negotiates the final
2 rates or rate adjustments with the large employer in
3 accordance with the rate manual and rules of the currently
4 approved rate filing for the policy.

5 (Source: P.A. 103-106, eff. 1-1-24.)

6 Section 4-10. The Health Maintenance Organization Act is
7 amended by changing Section 4-12 as follows:

8 (215 ILCS 125/4-12) (from Ch. 111 1/2, par. 1409.5)

9 Sec. 4-12. Changes in rate methodology and benefits,
10 material modifications. A health maintenance organization
11 shall file with the Director, prior to use, a notice of any
12 change in rate methodology, or benefits and of any material
13 modification of any matter or document furnished pursuant to
14 Section 2-1, together with such supporting documents as are
15 necessary to fully explain the change or modification.

16 (a) Contract modifications described in subsections
17 (c) (5), (c) (6) and (c) (7) of Section 2-1 shall include all
18 form agreements between the organization and enrollees,
19 providers, administrators of services and insurers of health
20 maintenance organizations.

21 (b) Material transactions or series of transactions other
22 than those described in subsection (a) of this Section, the
23 total annual value of which exceeds the greater of \$100,000 or
24 5% of net earned subscription revenue for the most current

1 12-month period as determined from filed financial statements.

2 (c) Any agreement between the organization and an insurer
3 shall be subject to the provisions of the laws of this State
4 regarding reinsurance as provided in Article XI of the
5 Illinois Insurance Code. All reinsurance agreements must be
6 filed. Approval of the Director is required for all agreements
7 except the following: individual stop loss, aggregate excess,
8 hospitalization benefits or out-of-area of the participating
9 providers unless 20% or more of the organization's total risk
10 is reinsured, in which case all reinsurance agreements require
11 approval.

12 (d) In addition to any applicable provisions of this Act,
13 premium rate filings shall be subject to subsections (a) and
14 (c) through (j) ~~(i)~~ of Section 355 of the Illinois Insurance
15 Code.

16 (Source: P.A. 103-106, eff. 1-1-24.)

17 Section 4-15. The Limited Health Service Organization Act
18 is amended by changing Section 3006 as follows:

19 (215 ILCS 130/3006) (from Ch. 73, par. 1503-6)

20 Sec. 3006. Changes in rate methodology and benefits;
21 material modifications; addition of limited health services.

22 (a) A limited health service organization shall file with
23 the Director prior to use, a notice of any change in rate
24 methodology, charges, or benefits and of any material

1 modification of any matter or document furnished pursuant to
2 Section 2001, together with such supporting documents as are
3 necessary to fully explain the change or modification.

4 (1) Contract modifications described in paragraphs (5)
5 and (6) of subsection (c) of Section 2001 shall include
6 all agreements between the organization and enrollees,
7 providers, administrators of services, and insurers of
8 limited health services; also other material transactions
9 or series of transactions, the total annual value of which
10 exceeds the greater of \$100,000 or 5% of net earned
11 subscription revenue for the most current 12-month ~~12~~
12 ~~month~~ period as determined from filed financial
13 statements.

14 (2) Contract modification for reinsurance. Any
15 agreement between the organization and an insurer shall be
16 subject to the provisions of Article XI of the Illinois
17 Insurance Code, as now or hereafter amended. All
18 reinsurance agreements must be filed with the Director.
19 Approval of the Director in required agreements must be
20 filed. Approval of the director is required for all
21 agreements except individual stop loss, aggregate excess,
22 hospitalization benefits, or out-of-area of the
23 participating providers, unless 20% or more of the
24 organization's total risk is reinsured, in which case all
25 reinsurance agreements shall require approval.

26 (b) If a limited health service organization desires to

1 add one or more additional limited health services, it shall
2 file a notice with the Director and, at the same time, submit
3 the information required by Section 2001 if different from
4 that filed with the prepaid limited health service
5 organization's application. Issuance of such an amended
6 certificate of authority shall be subject to the conditions of
7 Section 2002 of this Act.

8 (c) In addition to any applicable provisions of this Act,
9 premium rate filings shall be subject to subsection (i) and,
10 for pharmaceutical policies, subsection (j) of Section 355 of
11 the Illinois Insurance Code.

12 (Source: P.A. 103-106, eff. 1-1-24; revised 1-2-24.)

13 Article 5.

14 Section 5-5. The Illinois Insurance Code is amended by
15 changing Sections 121-2.05, 356z.18, 367.3, 367a, and 368f and
16 by adding Section 352c as follows:

17 (215 ILCS 5/121-2.05) (from Ch. 73, par. 733-2.05)

18 Sec. 121-2.05. Group insurance policies issued and
19 delivered in other State-Transactions in this State. With the
20 exception of insurance transactions authorized under Sections
21 230.2 or 367.3 of this Code or transactions described under
22 Section 352c, transactions in this State involving group
23 legal, group life and group accident and health or blanket

1 accident and health insurance or group annuities where the
2 master policy of such groups was lawfully issued and delivered
3 in, and under the laws of, a State in which the insurer was
4 authorized to do an insurance business, to a group properly
5 established pursuant to law or regulation, and where the
6 policyholder is domiciled or otherwise has a bona fide situs.

7 (Source: P.A. 86-753.)

8 (215 ILCS 5/352c new)

9 Sec. 352c. Short-term, limited-duration insurance
10 prohibited.

11 (a) In this Section:

12 "Excepted benefits" has the meaning given to that term in
13 42 U.S.C. 300gg-91 and implementing regulations. "Excepted
14 benefits" includes individual, group, or blanket coverage.

15 "Short-term, limited-duration insurance" means any type of
16 accident and health insurance offered or provided within this
17 State pursuant to a group or individual policy or individual
18 certificate by a company, regardless of the situs state of the
19 delivery of the policy, that has an expiration date specified
20 in the contract that is fewer than 365 days after the original
21 effective date. Regardless of the duration of coverage,
22 "short-term, limited-duration insurance" does not include
23 excepted benefits or any student health insurance coverage.

24 (b) On and after January 1, 2025, no company shall issue,
25 deliver, amend, or renew short-term, limited-duration

1 insurance to any natural or legal person that is a resident or
2 domiciled in this State.

3 (215 ILCS 5/356z.18)

4 (Text of Section before amendment by P.A. 103-512)

5 Sec. 356z.18. Prosthetic and customized orthotic devices.

6 (a) For the purposes of this Section:

7 "Customized orthotic device" means a supportive device for
8 the body or a part of the body, the head, neck, or extremities,
9 and includes the replacement or repair of the device based on
10 the patient's physical condition as medically necessary,
11 excluding foot orthotics defined as an in-shoe device designed
12 to support the structural components of the foot during
13 weight-bearing activities.

14 "Licensed provider" means a prosthetist, orthotist, or
15 pedorthist licensed to practice in this State.

16 "Prosthetic device" means an artificial device to replace,
17 in whole or in part, an arm or leg and includes accessories
18 essential to the effective use of the device and the
19 replacement or repair of the device based on the patient's
20 physical condition as medically necessary.

21 (b) This amendatory Act of the 96th General Assembly shall
22 provide benefits to any person covered thereunder for expenses
23 incurred in obtaining a prosthetic or custom orthotic device
24 from any Illinois licensed prosthetist, licensed orthotist, or
25 licensed pedorthist as required under the Orthotics,

1 Prosthetics, and Pedorthics Practice Act.

2 (c) A group or individual major medical policy of accident
3 or health insurance or managed care plan or medical, health,
4 or hospital service corporation contract that provides
5 coverage for prosthetic or custom orthotic care and is
6 amended, delivered, issued, or renewed 6 months after the
7 effective date of this amendatory Act of the 96th General
8 Assembly must provide coverage for prosthetic and orthotic
9 devices in accordance with this subsection (c). The coverage
10 required under this Section shall be subject to the other
11 general exclusions, limitations, and financial requirements of
12 the policy, including coordination of benefits, participating
13 provider requirements, utilization review of health care
14 services, including review of medical necessity, case
15 management, and experimental and investigational treatments,
16 and other managed care provisions under terms and conditions
17 that are no less favorable than the terms and conditions that
18 apply to substantially all medical and surgical benefits
19 provided under the plan or coverage.

20 (d) The policy or plan or contract may require prior
21 authorization for the prosthetic or orthotic devices in the
22 same manner that prior authorization is required for any other
23 covered benefit.

24 (e) Repairs and replacements of prosthetic and orthotic
25 devices are also covered, subject to the co-payments and
26 deductibles, unless necessitated by misuse or loss.

1 (f) A policy or plan or contract may require that, if
2 coverage is provided through a managed care plan, the benefits
3 mandated pursuant to this Section shall be covered benefits
4 only if the prosthetic or orthotic devices are provided by a
5 licensed provider employed by a provider service who contracts
6 with or is designated by the carrier, to the extent that the
7 carrier provides in-network and out-of-network service, the
8 coverage for the prosthetic or orthotic device shall be
9 offered no less extensively.

10 (g) The policy or plan or contract shall also meet
11 adequacy requirements as established by the Health Care
12 Reimbursement Reform Act of 1985 of the Illinois Insurance
13 Code.

14 (h) This Section shall not apply to accident only,
15 specified disease, short-term travel ~~hospital or medical~~,
16 hospital confinement indemnity or other fixed indemnity,
17 credit, dental, vision, Medicare supplement, long-term care,
18 basic hospital and medical-surgical expense coverage,
19 disability income insurance coverage, coverage issued as a
20 supplement to liability insurance, workers' compensation
21 insurance, or automobile medical payment insurance.

22 (Source: P.A. 96-833, eff. 6-1-10.)

23 (Text of Section after amendment by P.A. 103-512)

24 Sec. 356z.18. Prosthetic and customized orthotic devices.

25 (a) For the purposes of this Section:

1 "Customized orthotic device" means a supportive device for
2 the body or a part of the body, the head, neck, or extremities,
3 and includes the replacement or repair of the device based on
4 the patient's physical condition as medically necessary,
5 excluding foot orthotics defined as an in-shoe device designed
6 to support the structural components of the foot during
7 weight-bearing activities.

8 "Licensed provider" means a prosthetist, orthotist, or
9 pedorthist licensed to practice in this State.

10 "Prosthetic device" means an artificial device to replace,
11 in whole or in part, an arm or leg and includes accessories
12 essential to the effective use of the device and the
13 replacement or repair of the device based on the patient's
14 physical condition as medically necessary.

15 (b) This amendatory Act of the 96th General Assembly shall
16 provide benefits to any person covered thereunder for expenses
17 incurred in obtaining a prosthetic or custom orthotic device
18 from any Illinois licensed prosthetist, licensed orthotist, or
19 licensed pedorthist as required under the Orthotics,
20 Prosthetics, and Pedorthics Practice Act.

21 (c) A group or individual major medical policy of accident
22 or health insurance or managed care plan or medical, health,
23 or hospital service corporation contract that provides
24 coverage for prosthetic or custom orthotic care and is
25 amended, delivered, issued, or renewed 6 months after the
26 effective date of this amendatory Act of the 96th General

1 Assembly must provide coverage for prosthetic and orthotic
2 devices in accordance with this subsection (c). The coverage
3 required under this Section shall be subject to the other
4 general exclusions, limitations, and financial requirements of
5 the policy, including coordination of benefits, participating
6 provider requirements, utilization review of health care
7 services, including review of medical necessity, case
8 management, and experimental and investigational treatments,
9 and other managed care provisions under terms and conditions
10 that are no less favorable than the terms and conditions that
11 apply to substantially all medical and surgical benefits
12 provided under the plan or coverage.

13 (d) With respect to an enrollee at any age, in addition to
14 coverage of a prosthetic or custom orthotic device required by
15 this Section, benefits shall be provided for a prosthetic or
16 custom orthotic device determined by the enrollee's provider
17 to be the most appropriate model that is medically necessary
18 for the enrollee to perform physical activities, as
19 applicable, such as running, biking, swimming, and lifting
20 weights, and to maximize the enrollee's whole body health and
21 strengthen the lower and upper limb function.

22 (e) The requirements of this Section do not constitute an
23 addition to this State's essential health benefits that
24 requires defrayal of costs by this State pursuant to 42 U.S.C.
25 18031(d)(3)(B).

26 (f) The policy or plan or contract may require prior

1 authorization for the prosthetic or orthotic devices in the
2 same manner that prior authorization is required for any other
3 covered benefit.

4 (g) Repairs and replacements of prosthetic and orthotic
5 devices are also covered, subject to the co-payments and
6 deductibles, unless necessitated by misuse or loss.

7 (h) A policy or plan or contract may require that, if
8 coverage is provided through a managed care plan, the benefits
9 mandated pursuant to this Section shall be covered benefits
10 only if the prosthetic or orthotic devices are provided by a
11 licensed provider employed by a provider service who contracts
12 with or is designated by the carrier, to the extent that the
13 carrier provides in-network and out-of-network service, the
14 coverage for the prosthetic or orthotic device shall be
15 offered no less extensively.

16 (i) The policy or plan or contract shall also meet
17 adequacy requirements as established by the Health Care
18 Reimbursement Reform Act of 1985 of the Illinois Insurance
19 Code.

20 (j) This Section shall not apply to accident only,
21 specified disease, short-term travel ~~hospital or medical~~,
22 hospital confinement indemnity or other fixed indemnity,
23 credit, dental, vision, Medicare supplement, long-term care,
24 basic hospital and medical-surgical expense coverage,
25 disability income insurance coverage, coverage issued as a
26 supplement to liability insurance, workers' compensation

1 insurance, or automobile medical payment insurance.

2 (Source: P.A. 103-512, eff. 1-1-25.)

3 (215 ILCS 5/367.3) (from Ch. 73, par. 979.3)

4 Sec. 367.3. Group accident and health insurance;
5 discretionary groups.

6 (a) No group health insurance offered to a resident of
7 this State under a policy issued to a group, other than one
8 specifically described in Section 367(1), shall be delivered
9 or issued for delivery in this State unless the Director
10 determines that:

11 (1) the issuance of the policy is not contrary to the
12 public interest;

13 (2) the issuance of the policy will result in
14 economies of acquisition and administration; and

15 (3) the benefits under the policy are reasonable in
16 relation to the premium charged.

17 (b) No such group health insurance may be offered in this
18 State under a policy issued in another state unless this State
19 or the state in which the group policy is issued has made a
20 determination that the requirements of subsection (a) have
21 been met.

22 Where insurance is to be offered in this State under a
23 policy described in this subsection, the insurer shall file
24 for informational review purposes:

25 (1) a copy of the group master contract;

1 (2) a copy of the statute authorizing the issuance of
2 the group policy in the state of situs, which statute has
3 the same or similar requirements as this State, or in the
4 absence of such statute, a certification by an officer of
5 the company that the policy meets the Illinois minimum
6 standards required for individual accident and health
7 policies under authority of Section 401 of this Code, as
8 now or hereafter amended, as promulgated by rule at 50
9 Illinois Administrative Code, Ch. I, Sec. 2007, et seq.,
10 as now or hereafter amended, or by a successor rule;

11 (3) evidence of approval by the state of situs of the
12 group master policy; and

13 (4) copies of all supportive material furnished to the
14 state of situs to satisfy the criteria for approval.

15 (c) The Director may, at any time after receipt of the
16 information required under subsection (b) and after finding
17 that the standards of subsection (a) have not been met, order
18 the insurer to cease the issuance or marketing of that
19 coverage in this State.

20 (d) Notwithstanding subsections (a) and (b), group ~~Group~~
21 accident and health insurance subject to the provisions of
22 this Section is also subject to the provisions of Sections
23 352c and Section 367i of this Code and rules thereunder.

24 (Source: P.A. 90-655, eff. 7-30-98.)

25 (215 ILCS 5/367a) (from Ch. 73, par. 979a)

1 Sec. 367a. Blanket accident and health insurance.

2 (1) Blanket accident and health insurance is the ~~that~~ form
3 of accident and health insurance providing excepted benefits,
4 as defined in Section 352c, that covers ~~covering~~ special
5 groups of persons as enumerated in one of the following
6 paragraphs (a) to (g), inclusive:

7 (a) Under a policy or contract issued to any carrier for
8 hire, which shall be deemed the policyholder, covering a group
9 defined as all persons who may become passengers on such
10 carrier.

11 (b) Under a policy or contract issued to an employer, who
12 shall be deemed the policyholder, covering all employees or
13 any group of employees defined by reference to exceptional
14 hazards incident to such employment.

15 (c) Under a policy or contract issued to a college,
16 school, or other institution of learning or to the head or
17 principal thereof, who or which shall be deemed the
18 policyholder, covering students or teachers. However, student
19 health insurance coverage, as defined in 45 CFR 147.145, shall
20 remain subject to the standards and requirements for
21 individual health insurance coverage except where inconsistent
22 with that regulation. An issuer providing student health
23 insurance coverage or a policy or contract covering students
24 for limited-scope dental or vision under 45 CFR 148.220 shall
25 require an individual application or enrollment form and shall
26 furnish each insured individual a certificate, which shall

1 have been approved by the Director under Section 355.

2 (d) Under a policy or contract issued in the name of any
3 volunteer fire department, first aid, or other such volunteer
4 group, which shall be deemed the policyholder, covering all of
5 the members of such department or group.

6 (e) Under a policy or contract issued to a creditor, who
7 shall be deemed the policyholder, to insure debtors of the
8 creditors; Provided, however, that in the case of a loan which
9 is subject to the Small Loans Act, no insurance premium or
10 other cost shall be directly or indirectly charged or assessed
11 against, or collected or received from the borrower.

12 (f) Under a policy or contract issued to a sports team or
13 to a camp, which team or camp sponsor shall be deemed the
14 policyholder, covering members or campers.

15 (g) Under a policy or contract issued to any other
16 substantially similar group which, in the discretion of the
17 Director, may be subject to the issuance of a blanket accident
18 and health policy or contract.

19 (2) Any insurance company authorized to write accident and
20 health insurance in this state shall have the power to issue
21 blanket accident and health insurance. No such blanket policy
22 may be issued or delivered in this State unless a copy of the
23 form thereof shall have been filed in accordance with Section
24 355, and it contains in substance such of those provisions
25 contained in Sections 357.1 through 357.30 as may be
26 applicable to blanket accident and health insurance and the

1 following provisions:

2 (a) A provision that the policy and the application shall
3 constitute the entire contract between the parties, and that
4 all statements made by the policyholder shall, in absence of
5 fraud, be deemed representations and not warranties, and that
6 no such statements shall be used in defense to a claim under
7 the policy, unless it is contained in a written application.

8 (b) A provision that to the group or class thereof
9 originally insured shall be added from time to time all new
10 persons or individuals eligible for coverage.

11 (3) An individual application shall not be required from a
12 person covered under a blanket accident or health policy or
13 contract, nor shall it be necessary for the insurer to furnish
14 each person a certificate.

15 (4) All benefits under any blanket accident and health
16 policy shall be payable to the person insured, or to his
17 designated beneficiary or beneficiaries, or to his or her
18 estate, except that if the person insured be a minor or person
19 under legal disability, such benefits may be made payable to
20 his or her parent, guardian, or other person actually
21 supporting him or her. Provided further, however, that the
22 policy may provide that all or any portion of any indemnities
23 provided by any such policy on account of hospital, nursing,
24 medical or surgical services may, at the insurer's option, be
25 paid directly to the hospital or person rendering such
26 services; but the policy may not require that the service be

1 rendered by a particular hospital or person. Payment so made
2 shall discharge the insurer's obligation with respect to the
3 amount of insurance so paid.

4 (5) Nothing contained in this section shall be deemed to
5 affect the legal liability of policyholders for the death of
6 or injury to, any such member of such group.

7 (Source: P.A. 83-1362.)

8 (215 ILCS 5/368f)

9 Sec. 368f. Military service member insurance
10 reinstatement.

11 (a) No Illinois resident activated for military service
12 and no spouse or dependent of the resident who becomes
13 eligible for a federal government-sponsored health insurance
14 program, including the TriCare program providing coverage for
15 civilian dependents of military personnel, as a result of the
16 activation shall be denied reinstatement into the same
17 individual health insurance coverage with the health insurer
18 that the resident lapsed as a result of activation or becoming
19 covered by the federal government-sponsored health insurance
20 program. The resident shall have the right to reinstatement in
21 the same individual health insurance coverage without medical
22 underwriting, subject to payment of the current premium
23 charged to other persons of the same age and gender that are
24 covered under the same individual health coverage. Except in
25 the case of birth or adoption that occurs during the period of

1 activation, reinstatement must be into the same coverage type
2 as the resident held prior to lapsing the individual health
3 insurance coverage and at the same or, at the option of the
4 resident, higher deductible level. The reinstatement rights
5 provided under this subsection (a) are not available to a
6 resident or dependents if the activated person is discharged
7 from the military under other than honorable conditions.

8 (b) The health insurer with which the reinstatement is
9 being requested must receive a request for reinstatement no
10 later than 63 days following the later of (i) deactivation or
11 (ii) loss of coverage under the federal government-sponsored
12 health insurance program. The health insurer may request proof
13 of loss of coverage and the timing of the loss of coverage of
14 the government-sponsored coverage in order to determine
15 eligibility for reinstatement into the individual coverage.
16 The effective date of the reinstatement of individual health
17 coverage shall be the first of the month following receipt of
18 the notice requesting reinstatement.

19 (c) All insurers must provide written notice to the
20 policyholder of individual health coverage of the rights
21 described in subsection (a) of this Section. In lieu of the
22 inclusion of the notice in the individual health insurance
23 policy, an insurance company may satisfy the notification
24 requirement by providing a single written notice:

25 (1) in conjunction with the enrollment process for a
26 policyholder initially enrolling in the individual

1 coverage on or after the effective date of this amendatory
2 Act of the 94th General Assembly; or

3 (2) by mailing written notice to policyholders whose
4 coverage was effective prior to the effective date of this
5 amendatory Act of the 94th General Assembly no later than
6 90 days following the effective date of this amendatory
7 Act of the 94th General Assembly.

8 (d) The provisions of subsection (a) of this Section do
9 not apply to any policy or certificate providing coverage for
10 any specified disease, specified accident or accident-only
11 coverage, credit, dental, disability income, hospital
12 indemnity or other fixed indemnity, long-term care, Medicare
13 supplement, vision care, or short-term travel nonrenewable
14 ~~health policy~~ or other limited-benefit supplemental insurance,
15 or any coverage issued as a supplement to any liability
16 insurance, workers' compensation or similar insurance, or any
17 insurance under which benefits are payable with or without
18 regard to fault, whether written on a group, blanket, or
19 individual basis.

20 (e) Nothing in this Section shall require an insurer to
21 reinstate the resident if the insurer requires residency in an
22 enrollment area and those residency requirements are not met
23 after deactivation or loss of coverage under the
24 government-sponsored health insurance program.

25 (f) All terms, conditions, and limitations of the
26 individual coverage into which reinstatement is made apply

1 equally to all insureds enrolled in the coverage.

2 (g) The Secretary may adopt rules as may be necessary to
3 carry out the provisions of this Section.

4 (Source: P.A. 94-1037, eff. 7-20-06.)

5 Section 5-10. The Health Maintenance Organization Act is
6 amended by changing Section 5-3 as follows:

7 (215 ILCS 125/5-3) (from Ch. 111 1/2, par. 1411.2)

8 Sec. 5-3. Insurance Code provisions.

9 (a) Health Maintenance Organizations shall be subject to
10 the provisions of Sections 133, 134, 136, 137, 139, 140,
11 141.1, 141.2, 141.3, 143, 143c, 147, 148, 149, 151, 152, 153,
12 154, 154.5, 154.6, 154.7, 154.8, 155.04, 155.22a, 155.49,
13 352c, 355.2, 355.3, 355b, 355c, 356f, 356g.5-1, 356m, 356q,
14 356v, 356w, 356x, 356z.2, 356z.3a, 356z.4, 356z.4a, 356z.5,
15 356z.6, 356z.8, 356z.9, 356z.10, 356z.11, 356z.12, 356z.13,
16 356z.14, 356z.15, 356z.17, 356z.18, 356z.19, 356z.20, 356z.21,
17 356z.22, 356z.23, 356z.24, 356z.25, 356z.26, 356z.28, 356z.29,
18 356z.30, 356z.30a, 356z.31, 356z.32, 356z.33, 356z.34,
19 356z.35, 356z.36, 356z.37, 356z.38, 356z.39, 356z.40, 356z.41,
20 356z.44, 356z.45, 356z.46, 356z.47, 356z.48, 356z.49, 356z.50,
21 356z.51, 356z.53, 356z.54, 356z.55, 356z.56, 356z.57, 356z.58,
22 356z.59, 356z.60, 356z.61, 356z.62, 356z.64, 356z.65, 356z.67,
23 356z.68, 364, 364.01, 364.3, 367.2, 367.2-5, 367i, 368a, 368b,
24 368c, 368d, 368e, 370c, 370c.1, 401, 401.1, 402, 403, 403A,

1 408, 408.2, 409, 412, 444, and 444.1, paragraph (c) of
2 subsection (2) of Section 367, and Articles IIA, VIII 1/2,
3 XII, XII 1/2, XIII, XIII 1/2, XXV, XXVI, and XXXIIB of the
4 Illinois Insurance Code.

5 (b) For purposes of the Illinois Insurance Code, except
6 for Sections 444 and 444.1 and Articles XIII and XIII 1/2,
7 Health Maintenance Organizations in the following categories
8 are deemed to be "domestic companies":

9 (1) a corporation authorized under the Dental Service
10 Plan Act or the Voluntary Health Services Plans Act;

11 (2) a corporation organized under the laws of this
12 State; or

13 (3) a corporation organized under the laws of another
14 state, 30% or more of the enrollees of which are residents
15 of this State, except a corporation subject to
16 substantially the same requirements in its state of
17 organization as is a "domestic company" under Article VIII
18 1/2 of the Illinois Insurance Code.

19 (c) In considering the merger, consolidation, or other
20 acquisition of control of a Health Maintenance Organization
21 pursuant to Article VIII 1/2 of the Illinois Insurance Code,

22 (1) the Director shall give primary consideration to
23 the continuation of benefits to enrollees and the
24 financial conditions of the acquired Health Maintenance
25 Organization after the merger, consolidation, or other
26 acquisition of control takes effect;

1 (2) (i) the criteria specified in subsection (1) (b) of
2 Section 131.8 of the Illinois Insurance Code shall not
3 apply and (ii) the Director, in making his determination
4 with respect to the merger, consolidation, or other
5 acquisition of control, need not take into account the
6 effect on competition of the merger, consolidation, or
7 other acquisition of control;

8 (3) the Director shall have the power to require the
9 following information:

10 (A) certification by an independent actuary of the
11 adequacy of the reserves of the Health Maintenance
12 Organization sought to be acquired;

13 (B) pro forma financial statements reflecting the
14 combined balance sheets of the acquiring company and
15 the Health Maintenance Organization sought to be
16 acquired as of the end of the preceding year and as of
17 a date 90 days prior to the acquisition, as well as pro
18 forma financial statements reflecting projected
19 combined operation for a period of 2 years;

20 (C) a pro forma business plan detailing an
21 acquiring party's plans with respect to the operation
22 of the Health Maintenance Organization sought to be
23 acquired for a period of not less than 3 years; and

24 (D) such other information as the Director shall
25 require.

26 (d) The provisions of Article VIII 1/2 of the Illinois

1 Insurance Code and this Section 5-3 shall apply to the sale by
2 any health maintenance organization of greater than 10% of its
3 enrollee population (including, without limitation, the health
4 maintenance organization's right, title, and interest in and
5 to its health care certificates).

6 (e) In considering any management contract or service
7 agreement subject to Section 141.1 of the Illinois Insurance
8 Code, the Director (i) shall, in addition to the criteria
9 specified in Section 141.2 of the Illinois Insurance Code,
10 take into account the effect of the management contract or
11 service agreement on the continuation of benefits to enrollees
12 and the financial condition of the health maintenance
13 organization to be managed or serviced, and (ii) need not take
14 into account the effect of the management contract or service
15 agreement on competition.

16 (f) Except for small employer groups as defined in the
17 Small Employer Rating, Renewability and Portability Health
18 Insurance Act and except for medicare supplement policies as
19 defined in Section 363 of the Illinois Insurance Code, a
20 Health Maintenance Organization may by contract agree with a
21 group or other enrollment unit to effect refunds or charge
22 additional premiums under the following terms and conditions:

23 (i) the amount of, and other terms and conditions with
24 respect to, the refund or additional premium are set forth
25 in the group or enrollment unit contract agreed in advance
26 of the period for which a refund is to be paid or

1 additional premium is to be charged (which period shall
2 not be less than one year); and

3 (ii) the amount of the refund or additional premium
4 shall not exceed 20% of the Health Maintenance
5 Organization's profitable or unprofitable experience with
6 respect to the group or other enrollment unit for the
7 period (and, for purposes of a refund or additional
8 premium, the profitable or unprofitable experience shall
9 be calculated taking into account a pro rata share of the
10 Health Maintenance Organization's administrative and
11 marketing expenses, but shall not include any refund to be
12 made or additional premium to be paid pursuant to this
13 subsection (f)). The Health Maintenance Organization and
14 the group or enrollment unit may agree that the profitable
15 or unprofitable experience may be calculated taking into
16 account the refund period and the immediately preceding 2
17 plan years.

18 The Health Maintenance Organization shall include a
19 statement in the evidence of coverage issued to each enrollee
20 describing the possibility of a refund or additional premium,
21 and upon request of any group or enrollment unit, provide to
22 the group or enrollment unit a description of the method used
23 to calculate (1) the Health Maintenance Organization's
24 profitable experience with respect to the group or enrollment
25 unit and the resulting refund to the group or enrollment unit
26 or (2) the Health Maintenance Organization's unprofitable

1 experience with respect to the group or enrollment unit and
2 the resulting additional premium to be paid by the group or
3 enrollment unit.

4 In no event shall the Illinois Health Maintenance
5 Organization Guaranty Association be liable to pay any
6 contractual obligation of an insolvent organization to pay any
7 refund authorized under this Section.

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

14 (Source: P.A. 102-30, eff. 1-1-22; 102-34, eff. 6-25-21;
15 102-203, eff. 1-1-22; 102-306, eff. 1-1-22; 102-443, eff.
16 1-1-22; 102-589, eff. 1-1-22; 102-642, eff. 1-1-22; 102-665,
17 eff. 10-8-21; 102-731, eff. 1-1-23; 102-775, eff. 5-13-22;
18 102-804, eff. 1-1-23; 102-813, eff. 5-13-22; 102-816, eff.
19 1-1-23; 102-860, eff. 1-1-23; 102-901, eff. 7-1-22; 102-1093,
20 eff. 1-1-23; 102-1117, eff. 1-13-23; 103-84, eff. 1-1-24;
21 103-91, eff. 1-1-24; 103-123, eff. 1-1-24; 103-154, eff.
22 6-30-23; 103-420, eff. 1-1-24; 103-426, eff. 8-4-23; 103-445,
23 eff. 1-1-24; 103-551, eff. 8-11-23; revised 8-29-23.)

24 Section 5-15. The Limited Health Service Organization Act
25 is amended by changing Section 4003 as follows:

1 (215 ILCS 130/4003) (from Ch. 73, par. 1504-3)

2 Sec. 4003. Illinois Insurance Code provisions. Limited
3 health service organizations shall be subject to the
4 provisions of Sections 133, 134, 136, 137, 139, 140, 141.1,
5 141.2, 141.3, 143, 143c, 147, 148, 149, 151, 152, 153, 154,
6 154.5, 154.6, 154.7, 154.8, 155.04, 155.37, 155.49, 352c,
7 355.2, 355.3, 355b, 356q, 356v, 356z.4, 356z.4a, 356z.10,
8 356z.21, 356z.22, 356z.25, 356z.26, 356z.29, 356z.30a,
9 356z.32, 356z.33, 356z.41, 356z.46, 356z.47, 356z.51, 356z.53,
10 356z.54, 356z.57, 356z.59, 356z.61, 356z.64, 356z.67, 356z.68,
11 364.3, 368a, 401, 401.1, 402, 403, 403A, 408, 408.2, 409, 412,
12 444, and 444.1 and Articles IIA, VIII 1/2, XII, XII 1/2, XIII,
13 XIII 1/2, XXV, and XXVI of the Illinois Insurance Code.
14 Nothing in this Section shall require a limited health care
15 plan to cover any service that is not a limited health service.
16 For purposes of the Illinois Insurance Code, except for
17 Sections 444 and 444.1 and Articles XIII and XIII 1/2, limited
18 health service organizations in the following categories are
19 deemed to be domestic companies:

20 (1) a corporation under the laws of this State; or

21 (2) a corporation organized under the laws of another
22 state, 30% or more of the enrollees of which are residents
23 of this State, except a corporation subject to
24 substantially the same requirements in its state of
25 organization as is a domestic company under Article VIII

1 condition" and any other term in Section 10 of the Managed Care
2 Reform and Patient Rights Act that is used in the other
3 Sections listed in this Section.

4 (Source: P.A. 102-409, eff. 1-1-22; 103-426, eff. 8-4-23.)

5 (215 ILCS 5/155.37)

6 Sec. 155.37. Drug formulary; notice.

7 (a) Insurance companies that transact the kinds of
8 insurance authorized under Class 1(b) or Class 2(a) of Section
9 4 of this Code and provide coverage for prescription drugs
10 through the use of a drug formulary must notify insureds of any
11 change in the formulary. A company may comply with this
12 Section by posting changes in the formulary on its website.

13 (b) No later than October 1, 2025, insurance companies
14 that use a drug formulary shall post the formulary on their
15 websites in a manner that is searchable and accessible to the
16 general public without requiring an individual to create any
17 account. This formulary shall adhere to a template developed
18 by the Department by March 31, 2025, which shall take into
19 consideration existing requirements for reporting of
20 information established by the federal Centers for Medicare
21 and Medicaid Services as well as display of cost-sharing
22 information. This template and all formularies also shall do
23 all the following:

24 (1) include information on cost-sharing tiers and
25 utilization controls, such as prior authorization, for

1 each covered drug;

2 (2) indicate any drugs on the formulary that are
3 preferred over other drugs on the formulary;

4 (3) include information to educate insureds about the
5 differences between drugs administered or provided under a
6 policy's medical benefit and drugs covered under a drug
7 benefit and how to obtain coverage information about drugs
8 that are not covered under the drug benefit;

9 (4) include information to educate insureds that
10 policies that provide drug benefits are required to have a
11 method for enrollees to obtain drugs not listed in the
12 formulary if they are deemed medically necessary by a
13 clinician under Section 45.1 of the Managed Care Reform
14 and Patient Rights Act;

15 (5) include information on which medications are
16 covered, including both generic and brand name; and

17 (6) include information on what tier of the plan's
18 drug formulary each medication is in.

19 (c) No formulary may establish a step therapy requirement
20 for any formulary drug or any drug covered as a result of a
21 medical exceptions procedure.

22 (Source: P.A. 92-440, eff. 8-17-01; 92-651, eff. 7-11-02.)

23 (215 ILCS 5/356z.40)

24 Sec. 356z.40. Pregnancy and postpartum coverage.

25 (a) An individual or group policy of accident and health

1 insurance or managed care plan amended, delivered, issued, or
2 renewed on or after the effective date of this amendatory Act
3 of the 102nd General Assembly shall provide coverage for
4 pregnancy and newborn care in accordance with 42 U.S.C.
5 18022(b) regarding essential health benefits.

6 (b) Benefits under this Section shall be as follows:

7 (1) An individual who has been identified as
8 experiencing a high-risk pregnancy by the individual's
9 treating provider shall have access to clinically
10 appropriate case management programs. As used in this
11 subsection, "case management" means a mechanism to
12 coordinate and assure continuity of services, including,
13 but not limited to, health services, social services, and
14 educational services necessary for the individual. "Case
15 management" involves individualized assessment of needs,
16 planning of services, referral, monitoring, and advocacy
17 to assist an individual in gaining access to appropriate
18 services and closure when services are no longer required.
19 "Case management" is an active and collaborative process
20 involving a single qualified case manager, the individual,
21 the individual's family, the providers, and the community.
22 This includes close coordination and involvement with all
23 service providers in the management plan for that
24 individual or family, including assuring that the
25 individual receives the services. As used in this
26 subsection, "high-risk pregnancy" means a pregnancy in

1 which the pregnant or postpartum individual or baby is at
2 an increased risk for poor health or complications during
3 pregnancy or childbirth, including, but not limited to,
4 hypertension disorders, gestational diabetes, and
5 hemorrhage.

6 (2) An individual shall have access to medically
7 necessary treatment of a mental, emotional, nervous, or
8 substance use disorder or condition consistent with the
9 requirements set forth in this Section and in Sections
10 370c and 370c.1 of this Code.

11 (3) The benefits provided for inpatient and outpatient
12 services for the treatment of a mental, emotional,
13 nervous, or substance use disorder or condition related to
14 pregnancy or postpartum complications shall be provided if
15 determined to be medically necessary, consistent with the
16 requirements of Sections 370c and 370c.1 of this Code. The
17 facility or provider shall notify the insurer of both the
18 admission and the initial treatment plan within 48 hours
19 after admission or initiation of treatment. Subject to the
20 requirements of Sections 370c and 370c.1 of this Code,
21 nothing ~~Nothing~~ in this paragraph shall prevent an insurer
22 from applying concurrent and post-service utilization
23 review of health care services, including review of
24 medical necessity, case management, experimental and
25 investigational treatments, managed care provisions, and
26 other terms and conditions of the insurance policy.

1 (4) The benefits for the first 48 hours of initiation
2 of services for an inpatient admission, detoxification or
3 withdrawal management program, or partial hospitalization
4 admission for the treatment of a mental, emotional,
5 nervous, or substance use disorder or condition related to
6 pregnancy or postpartum complications shall be provided
7 without post-service or concurrent review of medical
8 necessity, as the medical necessity for the first 48 hours
9 of such services shall be determined solely by the covered
10 pregnant or postpartum individual's provider. Subject to
11 Section 370c and 370c.1 of this Code, nothing ~~Nothing~~ in
12 this paragraph shall prevent an insurer from applying
13 concurrent and post-service utilization review, including
14 the review of medical necessity, case management,
15 experimental and investigational treatments, managed care
16 provisions, and other terms and conditions of the
17 insurance policy, of any inpatient admission,
18 detoxification or withdrawal management program admission,
19 or partial hospitalization admission services for the
20 treatment of a mental, emotional, nervous, or substance
21 use disorder or condition related to pregnancy or
22 postpartum complications received 48 hours after the
23 initiation of such services. If an insurer determines that
24 the services are no longer medically necessary, then the
25 covered person shall have the right to external review
26 pursuant to the requirements of the Health Carrier

1 External Review Act.

2 (5) If an insurer determines that continued inpatient
3 care, detoxification or withdrawal management, partial
4 hospitalization, intensive outpatient treatment, or
5 outpatient treatment in a facility is no longer medically
6 necessary, the insurer shall, within 24 hours, provide
7 written notice to the covered pregnant or postpartum
8 individual and the covered pregnant or postpartum
9 individual's provider of its decision and the right to
10 file an expedited internal appeal of the determination.
11 The insurer shall review and make a determination with
12 respect to the internal appeal within 24 hours and
13 communicate such determination to the covered pregnant or
14 postpartum individual and the covered pregnant or
15 postpartum individual's provider. If the determination is
16 to uphold the denial, the covered pregnant or postpartum
17 individual and the covered pregnant or postpartum
18 individual's provider have the right to file an expedited
19 external appeal. An independent ~~utilization~~ review
20 organization shall make a determination within 72 hours.
21 If the insurer's determination is upheld and it is
22 determined that continued inpatient care, detoxification
23 or withdrawal management, partial hospitalization,
24 intensive outpatient treatment, or outpatient treatment is
25 not medically necessary, the insurer shall remain
26 responsible for providing benefits for the inpatient care,

1 detoxification or withdrawal management, partial
2 hospitalization, intensive outpatient treatment, or
3 outpatient treatment through the day following the date
4 the determination is made, and the covered pregnant or
5 postpartum individual shall only be responsible for any
6 applicable copayment, deductible, and coinsurance for the
7 stay through that date as applicable under the policy. The
8 covered pregnant or postpartum individual shall not be
9 discharged or released from the inpatient facility,
10 detoxification or withdrawal management, partial
11 hospitalization, intensive outpatient treatment, or
12 outpatient treatment until all internal appeals and
13 independent utilization review organization appeals are
14 exhausted. A decision to reverse an adverse determination
15 shall comply with the Health Carrier External Review Act.

16 (6) Except as otherwise stated in this subsection (b),
17 the benefits and cost-sharing shall be provided to the
18 same extent as for any other medical condition covered
19 under the policy.

20 (7) The benefits required by paragraphs (2) and (6) of
21 this subsection (b) are to be provided to all covered
22 pregnant or postpartum individuals with a diagnosis of a
23 mental, emotional, nervous, or substance use disorder or
24 condition. The presence of additional related or unrelated
25 diagnoses shall not be a basis to reduce or deny the
26 benefits required by this subsection (b).

1 (Source: P.A. 102-665, eff. 10-8-21.)

2 (215 ILCS 5/370c) (from Ch. 73, par. 982c)

3 Sec. 370c. Mental and emotional disorders.

4 (a)(1) On and after January 1, 2022 (the effective date of
5 Public Act 102-579), every insurer that amends, delivers,
6 issues, or renews group accident and health policies providing
7 coverage for hospital or medical treatment or services for
8 illness on an expense-incurred basis shall provide coverage
9 for the medically necessary treatment of mental, emotional,
10 nervous, or substance use disorders or conditions consistent
11 with the parity requirements of Section 370c.1 of this Code.

12 (2) Each insured that is covered for mental, emotional,
13 nervous, or substance use disorders or conditions shall be
14 free to select the physician licensed to practice medicine in
15 all its branches, licensed clinical psychologist, licensed
16 clinical social worker, licensed clinical professional
17 counselor, licensed marriage and family therapist, licensed
18 speech-language pathologist, or other licensed or certified
19 professional at a program licensed pursuant to the Substance
20 Use Disorder Act of his or her choice to treat such disorders,
21 and the insurer shall pay the covered charges of such
22 physician licensed to practice medicine in all its branches,
23 licensed clinical psychologist, licensed clinical social
24 worker, licensed clinical professional counselor, licensed
25 marriage and family therapist, licensed speech-language

1 pathologist, or other licensed or certified professional at a
2 program licensed pursuant to the Substance Use Disorder Act up
3 to the limits of coverage, provided (i) the disorder or
4 condition treated is covered by the policy, and (ii) the
5 physician, licensed psychologist, licensed clinical social
6 worker, licensed clinical professional counselor, licensed
7 marriage and family therapist, licensed speech-language
8 pathologist, or other licensed or certified professional at a
9 program licensed pursuant to the Substance Use Disorder Act is
10 authorized to provide said services under the statutes of this
11 State and in accordance with accepted principles of his or her
12 profession.

13 (3) Insofar as this Section applies solely to licensed
14 clinical social workers, licensed clinical professional
15 counselors, licensed marriage and family therapists, licensed
16 speech-language pathologists, and other licensed or certified
17 professionals at programs licensed pursuant to the Substance
18 Use Disorder Act, those persons who may provide services to
19 individuals shall do so after the licensed clinical social
20 worker, licensed clinical professional counselor, licensed
21 marriage and family therapist, licensed speech-language
22 pathologist, or other licensed or certified professional at a
23 program licensed pursuant to the Substance Use Disorder Act
24 has informed the patient of the desirability of the patient
25 conferring with the patient's primary care physician.

26 (4) "Mental, emotional, nervous, or substance use disorder

1 or condition" means a condition or disorder that involves a
2 mental health condition or substance use disorder that falls
3 under any of the diagnostic categories listed in the mental
4 and behavioral disorders chapter of the current edition of the
5 World Health Organization's International Classification of
6 Disease or that is listed in the most recent version of the
7 American Psychiatric Association's Diagnostic and Statistical
8 Manual of Mental Disorders. "Mental, emotional, nervous, or
9 substance use disorder or condition" includes any mental
10 health condition that occurs during pregnancy or during the
11 postpartum period and includes, but is not limited to,
12 postpartum depression.

13 (5) Medically necessary treatment and medical necessity
14 determinations shall be interpreted and made in a manner that
15 is consistent with and pursuant to subsections (h) through
16 (t).

17 (b) (1) (Blank).

18 (2) (Blank).

19 (2.5) (Blank).

20 (3) Unless otherwise prohibited by federal law and
21 consistent with the parity requirements of Section 370c.1 of
22 this Code, the reimbursing insurer that amends, delivers,
23 issues, or renews a group or individual policy of accident and
24 health insurance, a qualified health plan offered through the
25 health insurance marketplace, or a provider of treatment of
26 mental, emotional, nervous, or substance use disorders or

1 conditions shall furnish medical records or other necessary
2 data that substantiate that initial or continued treatment is
3 at all times medically necessary. An insurer shall provide a
4 mechanism for the timely review by a provider holding the same
5 license and practicing in the same specialty as the patient's
6 provider, who is unaffiliated with the insurer, jointly
7 selected by the patient (or the patient's next of kin or legal
8 representative if the patient is unable to act for himself or
9 herself), the patient's provider, and the insurer in the event
10 of a dispute between the insurer and patient's provider
11 regarding the medical necessity of a treatment proposed by a
12 patient's provider. If the reviewing provider determines the
13 treatment to be medically necessary, the insurer shall provide
14 reimbursement for the treatment. Future contractual or
15 employment actions by the insurer regarding the patient's
16 provider may not be based on the provider's participation in
17 this procedure. Nothing prevents the insured from agreeing in
18 writing to continue treatment at his or her expense. When
19 making a determination of the medical necessity for a
20 treatment modality for mental, emotional, nervous, or
21 substance use disorders or conditions, an insurer must make
22 the determination in a manner that is consistent with the
23 manner used to make that determination with respect to other
24 diseases or illnesses covered under the policy, including an
25 appeals process. Medical necessity determinations for
26 substance use disorders shall be made in accordance with

1 appropriate patient placement criteria established by the
2 American Society of Addiction Medicine. No additional criteria
3 may be used to make medical necessity determinations for
4 substance use disorders.

5 (4) A group health benefit plan amended, delivered,
6 issued, or renewed on or after January 1, 2019 (the effective
7 date of Public Act 100-1024) or an individual policy of
8 accident and health insurance or a qualified health plan
9 offered through the health insurance marketplace amended,
10 delivered, issued, or renewed on or after January 1, 2019 (the
11 effective date of Public Act 100-1024):

12 (A) shall provide coverage based upon medical
13 necessity for the treatment of a mental, emotional,
14 nervous, or substance use disorder or condition consistent
15 with the parity requirements of Section 370c.1 of this
16 Code; provided, however, that in each calendar year
17 coverage shall not be less than the following:

18 (i) 45 days of inpatient treatment; and

19 (ii) beginning on June 26, 2006 (the effective
20 date of Public Act 94-921), 60 visits for outpatient
21 treatment including group and individual outpatient
22 treatment; and

23 (iii) for plans or policies delivered, issued for
24 delivery, renewed, or modified after January 1, 2007
25 (the effective date of Public Act 94-906), 20
26 additional outpatient visits for speech therapy for

1 treatment of pervasive developmental disorders that
2 will be in addition to speech therapy provided
3 pursuant to item (ii) of this subparagraph (A); and

4 (B) may not include a lifetime limit on the number of
5 days of inpatient treatment or the number of outpatient
6 visits covered under the plan.

7 (C) (Blank).

8 (5) An issuer of a group health benefit plan or an
9 individual policy of accident and health insurance or a
10 qualified health plan offered through the health insurance
11 marketplace may not count toward the number of outpatient
12 visits required to be covered under this Section an outpatient
13 visit for the purpose of medication management and shall cover
14 the outpatient visits under the same terms and conditions as
15 it covers outpatient visits for the treatment of physical
16 illness.

17 (5.5) An individual or group health benefit plan amended,
18 delivered, issued, or renewed on or after September 9, 2015
19 (the effective date of Public Act 99-480) shall offer coverage
20 for medically necessary acute treatment services and medically
21 necessary clinical stabilization services. The treating
22 provider shall base all treatment recommendations and the
23 health benefit plan shall base all medical necessity
24 determinations for substance use disorders in accordance with
25 the most current edition of the Treatment Criteria for
26 Addictive, Substance-Related, and Co-Occurring Conditions

1 established by the American Society of Addiction Medicine. The
2 treating provider shall base all treatment recommendations and
3 the health benefit plan shall base all medical necessity
4 determinations for medication-assisted treatment in accordance
5 with the most current Treatment Criteria for Addictive,
6 Substance-Related, and Co-Occurring Conditions established by
7 the American Society of Addiction Medicine.

8 As used in this subsection:

9 "Acute treatment services" means 24-hour medically
10 supervised addiction treatment that provides evaluation and
11 withdrawal management and may include biopsychosocial
12 assessment, individual and group counseling, psychoeducational
13 groups, and discharge planning.

14 "Clinical stabilization services" means 24-hour treatment,
15 usually following acute treatment services for substance
16 abuse, which may include intensive education and counseling
17 regarding the nature of addiction and its consequences,
18 relapse prevention, outreach to families and significant
19 others, and aftercare planning for individuals beginning to
20 engage in recovery from addiction.

21 (6) An issuer of a group health benefit plan may provide or
22 offer coverage required under this Section through a managed
23 care plan.

24 (6.5) An individual or group health benefit plan amended,
25 delivered, issued, or renewed on or after January 1, 2019 (the
26 effective date of Public Act 100-1024):

1 (A) shall not impose prior authorization requirements,
2 other than those established under the Treatment Criteria
3 for Addictive, Substance-Related, and Co-Occurring
4 Conditions established by the American Society of
5 Addiction Medicine, on a prescription medication approved
6 by the United States Food and Drug Administration that is
7 prescribed or administered for the treatment of substance
8 use disorders;

9 (B) shall not impose any step therapy requirements,
10 ~~other than those established under the Treatment Criteria~~
11 ~~for Addictive, Substance-Related, and Co-Occurring~~
12 ~~Conditions established by the American Society of~~
13 ~~Addiction Medicine, before authorizing coverage for a~~
14 ~~prescription medication approved by the United States Food~~
15 ~~and Drug Administration that is prescribed or administered~~
16 ~~for the treatment of substance use disorders;~~

17 (C) shall place all prescription medications approved
18 by the United States Food and Drug Administration
19 prescribed or administered for the treatment of substance
20 use disorders on, for brand medications, the lowest tier
21 of the drug formulary developed and maintained by the
22 individual or group health benefit plan that covers brand
23 medications and, for generic medications, the lowest tier
24 of the drug formulary developed and maintained by the
25 individual or group health benefit plan that covers
26 generic medications; and

1 (D) shall not exclude coverage for a prescription
2 medication approved by the United States Food and Drug
3 Administration for the treatment of substance use
4 disorders and any associated counseling or wraparound
5 services on the grounds that such medications and services
6 were court ordered.

7 (7) (Blank).

8 (8) (Blank).

9 (9) With respect to all mental, emotional, nervous, or
10 substance use disorders or conditions, coverage for inpatient
11 treatment shall include coverage for treatment in a
12 residential treatment center certified or licensed by the
13 Department of Public Health or the Department of Human
14 Services.

15 (c) This Section shall not be interpreted to require
16 coverage for speech therapy or other rehabilitative services for
17 those individuals covered under Section 356z.15 of this Code.

18 (d) With respect to a group or individual policy of
19 accident and health insurance or a qualified health plan
20 offered through the health insurance marketplace, the
21 Department and, with respect to medical assistance, the
22 Department of Healthcare and Family Services shall each
23 enforce the requirements of this Section and Sections 356z.23
24 and 370c.1 of this Code, the Paul Wellstone and Pete Domenici
25 Mental Health Parity and Addiction Equity Act of 2008, 42
26 U.S.C. 18031(j), and any amendments to, and federal guidance

1 or regulations issued under, those Acts, including, but not
2 limited to, final regulations issued under the Paul Wellstone
3 and Pete Domenici Mental Health Parity and Addiction Equity
4 Act of 2008 and final regulations applying the Paul Wellstone
5 and Pete Domenici Mental Health Parity and Addiction Equity
6 Act of 2008 to Medicaid managed care organizations, the
7 Children's Health Insurance Program, and alternative benefit
8 plans. Specifically, the Department and the Department of
9 Healthcare and Family Services shall take action:

10 (1) proactively ensuring compliance by individual and
11 group policies, including by requiring that insurers
12 submit comparative analyses, as set forth in paragraph (6)
13 of subsection (k) of Section 370c.1, demonstrating how
14 they design and apply nonquantitative treatment
15 limitations, both as written and in operation, for mental,
16 emotional, nervous, or substance use disorder or condition
17 benefits as compared to how they design and apply
18 nonquantitative treatment limitations, as written and in
19 operation, for medical and surgical benefits;

20 (2) evaluating all consumer or provider complaints
21 regarding mental, emotional, nervous, or substance use
22 disorder or condition coverage for possible parity
23 violations;

24 (3) performing parity compliance market conduct
25 examinations or, in the case of the Department of
26 Healthcare and Family Services, parity compliance audits

1 of individual and group plans and policies, including, but
2 not limited to, reviews of:

3 (A) nonquantitative treatment limitations,
4 including, but not limited to, prior authorization
5 requirements, concurrent review, retrospective review,
6 step therapy, network admission standards,
7 reimbursement rates, and geographic restrictions;

8 (B) denials of authorization, payment, and
9 coverage; and

10 (C) other specific criteria as may be determined
11 by the Department.

12 The findings and the conclusions of the parity compliance
13 market conduct examinations and audits shall be made public.

14 The Director may adopt rules to effectuate any provisions
15 of the Paul Wellstone and Pete Domenici Mental Health Parity
16 and Addiction Equity Act of 2008 that relate to the business of
17 insurance.

18 (e) Availability of plan information.

19 (1) The criteria for medical necessity determinations
20 made under a group health plan, an individual policy of
21 accident and health insurance, or a qualified health plan
22 offered through the health insurance marketplace with
23 respect to mental health or substance use disorder
24 benefits (or health insurance coverage offered in
25 connection with the plan with respect to such benefits)
26 must be made available by the plan administrator (or the

1 health insurance issuer offering such coverage) to any
2 current or potential participant, beneficiary, or
3 contracting provider upon request.

4 (2) The reason for any denial under a group health
5 benefit plan, an individual policy of accident and health
6 insurance, or a qualified health plan offered through the
7 health insurance marketplace (or health insurance coverage
8 offered in connection with such plan or policy) of
9 reimbursement or payment for services with respect to
10 mental, emotional, nervous, or substance use disorders or
11 conditions benefits in the case of any participant or
12 beneficiary must be made available within a reasonable
13 time and in a reasonable manner and in readily
14 understandable language by the plan administrator (or the
15 health insurance issuer offering such coverage) to the
16 participant or beneficiary upon request.

17 (f) As used in this Section, "group policy of accident and
18 health insurance" and "group health benefit plan" includes (1)
19 State-regulated employer-sponsored group health insurance
20 plans written in Illinois or which purport to provide coverage
21 for a resident of this State; and (2) State employee health
22 plans.

23 (g) (1) As used in this subsection:

24 "Benefits", with respect to insurers, means the benefits
25 provided for treatment services for inpatient and outpatient
26 treatment of substance use disorders or conditions at American

1 Society of Addiction Medicine levels of treatment 2.1
2 (Intensive Outpatient), 2.5 (Partial Hospitalization), 3.1
3 (Clinically Managed Low-Intensity Residential), 3.3
4 (Clinically Managed Population-Specific High-Intensity
5 Residential), 3.5 (Clinically Managed High-Intensity
6 Residential), and 3.7 (Medically Monitored Intensive
7 Inpatient) and OMT (Opioid Maintenance Therapy) services.

8 "Benefits", with respect to managed care organizations,
9 means the benefits provided for treatment services for
10 inpatient and outpatient treatment of substance use disorders
11 or conditions at American Society of Addiction Medicine levels
12 of treatment 2.1 (Intensive Outpatient), 2.5 (Partial
13 Hospitalization), 3.5 (Clinically Managed High-Intensity
14 Residential), and 3.7 (Medically Monitored Intensive
15 Inpatient) and OMT (Opioid Maintenance Therapy) services.

16 "Substance use disorder treatment provider or facility"
17 means a licensed physician, licensed psychologist, licensed
18 psychiatrist, licensed advanced practice registered nurse, or
19 licensed, certified, or otherwise State-approved facility or
20 provider of substance use disorder treatment.

21 (2) A group health insurance policy, an individual health
22 benefit plan, or qualified health plan that is offered through
23 the health insurance marketplace, small employer group health
24 plan, and large employer group health plan that is amended,
25 delivered, issued, executed, or renewed in this State, or
26 approved for issuance or renewal in this State, on or after

1 January 1, 2019 (the effective date of Public Act 100-1023)
2 shall comply with the requirements of this Section and Section
3 370c.1. The services for the treatment and the ongoing
4 assessment of the patient's progress in treatment shall follow
5 the requirements of 77 Ill. Adm. Code 2060.

6 (3) Prior authorization shall not be utilized for the
7 benefits under this subsection. The substance use disorder
8 treatment provider or facility shall notify the insurer of the
9 initiation of treatment. For an insurer that is not a managed
10 care organization, the substance use disorder treatment
11 provider or facility notification shall occur for the
12 initiation of treatment of the covered person within 2
13 business days. For managed care organizations, the substance
14 use disorder treatment provider or facility notification shall
15 occur in accordance with the protocol set forth in the
16 provider agreement for initiation of treatment within 24
17 hours. If the managed care organization is not capable of
18 accepting the notification in accordance with the contractual
19 protocol during the 24-hour period following admission, the
20 substance use disorder treatment provider or facility shall
21 have one additional business day to provide the notification
22 to the appropriate managed care organization. Treatment plans
23 shall be developed in accordance with the requirements and
24 timeframes established in 77 Ill. Adm. Code 2060. If the
25 substance use disorder treatment provider or facility fails to
26 notify the insurer of the initiation of treatment in

1 accordance with these provisions, the insurer may follow its
2 normal prior authorization processes.

3 (4) For an insurer that is not a managed care
4 organization, if an insurer determines that benefits are no
5 longer medically necessary, the insurer shall notify the
6 covered person, the covered person's authorized
7 representative, if any, and the covered person's health care
8 provider in writing of the covered person's right to request
9 an external review pursuant to the Health Carrier External
10 Review Act. The notification shall occur within 24 hours
11 following the adverse determination.

12 Pursuant to the requirements of the Health Carrier
13 External Review Act, the covered person or the covered
14 person's authorized representative may request an expedited
15 external review. An expedited external review may not occur if
16 the substance use disorder treatment provider or facility
17 determines that continued treatment is no longer medically
18 necessary.

19 If an expedited external review request meets the criteria
20 of the Health Carrier External Review Act, an independent
21 review organization shall make a final determination of
22 medical necessity within 72 hours. If an independent review
23 organization upholds an adverse determination, an insurer
24 shall remain responsible to provide coverage of benefits
25 through the day following the determination of the independent
26 review organization. A decision to reverse an adverse

1 determination shall comply with the Health Carrier External
2 Review Act.

3 (5) The substance use disorder treatment provider or
4 facility shall provide the insurer with 7 business days'
5 advance notice of the planned discharge of the patient from
6 the substance use disorder treatment provider or facility and
7 notice on the day that the patient is discharged from the
8 substance use disorder treatment provider or facility.

9 (6) The benefits required by this subsection shall be
10 provided to all covered persons with a diagnosis of substance
11 use disorder or conditions. The presence of additional related
12 or unrelated diagnoses shall not be a basis to reduce or deny
13 the benefits required by this subsection.

14 (7) Nothing in this subsection shall be construed to
15 require an insurer to provide coverage for any of the benefits
16 in this subsection.

17 (h) As used in this Section:

18 "Generally accepted standards of mental, emotional,
19 nervous, or substance use disorder or condition care" means
20 standards of care and clinical practice that are generally
21 recognized by health care providers practicing in relevant
22 clinical specialties such as psychiatry, psychology, clinical
23 sociology, social work, addiction medicine and counseling, and
24 behavioral health treatment. Valid, evidence-based sources
25 reflecting generally accepted standards of mental, emotional,
26 nervous, or substance use disorder or condition care include

1 peer-reviewed scientific studies and medical literature,
2 recommendations of nonprofit health care provider professional
3 associations and specialty societies, including, but not
4 limited to, patient placement criteria and clinical practice
5 guidelines, recommendations of federal government agencies,
6 and drug labeling approved by the United States Food and Drug
7 Administration.

8 "Medically necessary treatment of mental, emotional,
9 nervous, or substance use disorders or conditions" means a
10 service or product addressing the specific needs of that
11 patient, for the purpose of screening, preventing, diagnosing,
12 managing, or treating an illness, injury, or condition or its
13 symptoms and comorbidities, including minimizing the
14 progression of an illness, injury, or condition or its
15 symptoms and comorbidities in a manner that is all of the
16 following:

17 (1) in accordance with the generally accepted
18 standards of mental, emotional, nervous, or substance use
19 disorder or condition care;

20 (2) clinically appropriate in terms of type,
21 frequency, extent, site, and duration; and

22 (3) not primarily for the economic benefit of the
23 insurer, purchaser, or for the convenience of the patient,
24 treating physician, or other health care provider.

25 "Utilization review" means either of the following:

26 (1) prospectively, retrospectively, or concurrently

1 reviewing and approving, modifying, delaying, or denying,
2 based in whole or in part on medical necessity, requests
3 by health care providers, insureds, or their authorized
4 representatives for coverage of health care services
5 before, retrospectively, or concurrently with the
6 provision of health care services to insureds.

7 (2) evaluating the medical necessity, appropriateness,
8 level of care, service intensity, efficacy, or efficiency
9 of health care services, benefits, procedures, or
10 settings, under any circumstances, to determine whether a
11 health care service or benefit subject to a medical
12 necessity coverage requirement in an insurance policy is
13 covered as medically necessary for an insured.

14 "Utilization review criteria" means patient placement
15 criteria or any criteria, standards, protocols, or guidelines
16 used by an insurer to conduct utilization review.

17 (i)(1) Every insurer that amends, delivers, issues, or
18 renews a group or individual policy of accident and health
19 insurance or a qualified health plan offered through the
20 health insurance marketplace in this State and Medicaid
21 managed care organizations providing coverage for hospital or
22 medical treatment on or after January 1, 2023 shall, pursuant
23 to subsections (h) through (s), provide coverage for medically
24 necessary treatment of mental, emotional, nervous, or
25 substance use disorders or conditions.

26 (2) An insurer shall not set a specific limit on the

1 duration of benefits or coverage of medically necessary
2 treatment of mental, emotional, nervous, or substance use
3 disorders or conditions or limit coverage only to alleviation
4 of the insured's current symptoms.

5 (3) All utilization review conducted ~~medical necessity~~
6 ~~determinations made~~ by the insurer concerning diagnosis,
7 prevention, and treatment ~~service intensity, level of care~~
8 ~~placement, continued stay, and transfer or discharge~~ of
9 insureds diagnosed with mental, emotional, nervous, or
10 substance use disorders or conditions shall be conducted in
11 accordance with the requirements of subsections (k) through
12 (w) ~~(u)~~.

13 (4) An insurer that authorizes a specific type of
14 treatment by a provider pursuant to this Section shall not
15 rescind or modify the authorization after that provider
16 renders the health care service in good faith and pursuant to
17 this authorization for any reason, including, but not limited
18 to, the insurer's subsequent cancellation or modification of
19 the insured's or policyholder's contract, or the insured's or
20 policyholder's eligibility. Nothing in this Section shall
21 require the insurer to cover a treatment when the
22 authorization was granted based on a material
23 misrepresentation by the insured, the policyholder, or the
24 provider. Nothing in this Section shall require Medicaid
25 managed care organizations to pay for services if the
26 individual was not eligible for Medicaid at the time the

1 service was rendered. Nothing in this Section shall require an
2 insurer to pay for services if the individual was not the
3 insurer's enrollee at the time services were rendered. As used
4 in this paragraph, "material" means a fact or situation that
5 is not merely technical in nature and results in or could
6 result in a substantial change in the situation.

7 (j) An insurer shall not limit benefits or coverage for
8 medically necessary services on the basis that those services
9 should be or could be covered by a public entitlement program,
10 including, but not limited to, special education or an
11 individualized education program, Medicaid, Medicare,
12 Supplemental Security Income, or Social Security Disability
13 Insurance, and shall not include or enforce a contract term
14 that excludes otherwise covered benefits on the basis that
15 those services should be or could be covered by a public
16 entitlement program. Nothing in this subsection shall be
17 construed to require an insurer to cover benefits that have
18 been authorized and provided for a covered person by a public
19 entitlement program. Medicaid managed care organizations are
20 not subject to this subsection.

21 (k) An insurer shall base any medical necessity
22 determination or the utilization review criteria that the
23 insurer, and any entity acting on the insurer's behalf,
24 applies to determine the medical necessity of health care
25 services and benefits for the diagnosis, prevention, and
26 treatment of mental, emotional, nervous, or substance use

1 disorders or conditions on current generally accepted
2 standards of mental, emotional, nervous, or substance use
3 disorder or condition care. All denials and appeals shall be
4 reviewed by a professional with experience or expertise
5 comparable to the provider requesting the authorization.

6 (l) In conducting utilization review of all covered health
7 care services for the diagnosis, prevention, and treatment of
8 ~~For medical necessity determinations relating to level of care~~
9 ~~placement, continued stay, and transfer or discharge of~~
10 ~~insureds diagnosed with~~ mental, emotional, and nervous
11 disorders or conditions, an insurer shall apply the ~~patient~~
12 ~~placement~~ criteria and guidelines set forth in the most recent
13 version of the treatment criteria developed by an unaffiliated
14 nonprofit professional association for the relevant clinical
15 specialty or, for Medicaid managed care organizations, ~~patient~~
16 ~~placement~~ criteria and guidelines determined by the Department
17 of Healthcare and Family Services that are consistent with
18 generally accepted standards of mental, emotional, nervous or
19 substance use disorder or condition care. Pursuant to
20 subsection (b), in conducting utilization review of all
21 covered services and benefits for the diagnosis, prevention,
22 and treatment of substance use disorders an insurer shall use
23 the most recent edition of the patient placement criteria
24 established by the American Society of Addiction Medicine.

25 (m) In conducting utilization review ~~For medical necessity~~
26 ~~determinations~~ relating to level of care placement, continued

1 stay, ~~and transfer, or discharge,~~ or any other patient care
2 decisions that are within the scope of the sources specified
3 in subsection (l), an insurer shall not apply different,
4 additional, conflicting, or more restrictive utilization
5 review criteria than the criteria set forth in those sources.
6 For all level of care placement decisions, the insurer shall
7 authorize placement at the level of care consistent with the
8 assessment of the insured using the relevant patient placement
9 criteria as specified in subsection (l). If that level of
10 placement is not available, the insurer shall authorize the
11 next higher level of care. In the event of disagreement, the
12 insurer shall provide full detail of its assessment using the
13 relevant criteria as specified in subsection (l) to the
14 provider of the service and the patient.

15 ~~Nothing in this subsection or subsection (l) prohibits an~~
16 ~~insurer from applying utilization review criteria that were~~
17 ~~developed in accordance with subsection (k) to health care~~
18 ~~services and benefits for mental, emotional, and nervous~~
19 ~~disorders or conditions that are not related to medical~~
20 ~~necessity determinations for level of care placement,~~
21 ~~continued stay, and transfer or discharge.~~ If an insurer
22 purchases or licenses utilization review criteria pursuant to
23 this subsection, the insurer shall verify and document before
24 use that the criteria were developed in accordance with
25 subsection (k).

26 (n) In conducting utilization review that is outside the

1 scope of the criteria as specified in subsection (l) or
2 relates to the advancements in technology or in the types or
3 levels of care that are not addressed in the most recent
4 versions of the sources specified in subsection (l), an
5 insurer shall conduct utilization review in accordance with
6 subsection (k).

7 (o) This Section does not in any way limit the rights of a
8 patient under the Medical Patient Rights Act.

9 (p) This Section does not in any way limit early and
10 periodic screening, diagnostic, and treatment benefits as
11 defined under 42 U.S.C. 1396d(r).

12 (q) To ensure the proper use of the criteria described in
13 subsection (l), every insurer shall do all of the following:

14 (1) Educate the insurer's staff, including any third
15 parties contracted with the insurer to review claims,
16 conduct utilization reviews, or make medical necessity
17 determinations about the utilization review criteria.

18 (2) Make the educational program available to other
19 stakeholders, including the insurer's participating or
20 contracted providers and potential participants,
21 beneficiaries, or covered lives. The education program
22 must be provided at least once a year, in-person or
23 digitally, or recordings of the education program must be
24 made available to the aforementioned stakeholders.

25 (3) Provide, at no cost, the utilization review
26 criteria and any training material or resources to

1 providers and insured patients upon request. For
2 utilization review criteria not concerning level of care
3 placement, continued stay, ~~and transfer,~~ ~~or discharge,~~ or
4 other patient care decisions used by the insurer pursuant
5 to subsection (m), the insurer may place the criteria on a
6 secure, password-protected website so long as the access
7 requirements of the website do not unreasonably restrict
8 access to insureds or their providers. No restrictions
9 shall be placed upon the insured's or treating provider's
10 access right to utilization review criteria obtained under
11 this paragraph at any point in time, including before an
12 initial request for authorization.

13 (4) Track, identify, and analyze how the utilization
14 review criteria are used to certify care, deny care, and
15 support the appeals process.

16 (5) Conduct interrater reliability testing to ensure
17 consistency in utilization review decision making that
18 covers how medical necessity decisions are made; this
19 assessment shall cover all aspects of utilization review
20 as defined in subsection (h).

21 (6) Run interrater reliability reports about how the
22 clinical guidelines are used in conjunction with the
23 utilization review process and parity compliance
24 activities.

25 (7) Achieve interrater reliability pass rates of at
26 least 90% and, if this threshold is not met, immediately

1 provide for the remediation of poor interrater reliability
2 and interrater reliability testing for all new staff
3 before they can conduct utilization review without
4 supervision.

5 (8) Maintain documentation of interrater reliability
6 testing and the remediation actions taken for those with
7 pass rates lower than 90% and submit to the Department of
8 Insurance or, in the case of Medicaid managed care
9 organizations, the Department of Healthcare and Family
10 Services the testing results and a summary of remedial
11 actions as part of parity compliance reporting set forth
12 in subsection (k) of Section 370c.1.

13 (r) This Section applies to all health care services and
14 benefits for the diagnosis, prevention, and treatment of
15 mental, emotional, nervous, or substance use disorders or
16 conditions covered by an insurance policy, including
17 prescription drugs.

18 (s) This Section applies to an insurer that amends,
19 delivers, issues, or renews a group or individual policy of
20 accident and health insurance or a qualified health plan
21 offered through the health insurance marketplace in this State
22 providing coverage for hospital or medical treatment and
23 conducts utilization review as defined in this Section,
24 including Medicaid managed care organizations, and any entity
25 or contracting provider that performs utilization review or
26 utilization management functions on an insurer's behalf.

1 (t) If the Director determines that an insurer has
2 violated this Section, the Director may, after appropriate
3 notice and opportunity for hearing, by order, assess a civil
4 penalty between \$1,000 and \$5,000 for each violation. Moneys
5 collected from penalties shall be deposited into the Parity
6 Advancement Fund established in subsection (i) of Section
7 370c.1.

8 (u) An insurer shall not adopt, impose, or enforce terms
9 in its policies or provider agreements, in writing or in
10 operation, that undermine, alter, or conflict with the
11 requirements of this Section.

12 (v) The provisions of this Section are severable. If any
13 provision of this Section or its application is held invalid,
14 that invalidity shall not affect other provisions or
15 applications that can be given effect without the invalid
16 provision or application.

17 (w) Beginning January 1, 2026, coverage for inpatient
18 mental health treatment at participating hospitals shall
19 comply with the following requirements:

20 (1) Subject to paragraphs (2) and (3) of this
21 subsection, no policy shall require prior authorization
22 for admission for such treatment at any participating
23 hospital.

24 (2) Coverage provided under this subsection also shall
25 not be subject to concurrent review for the first 72
26 hours, provided that the hospital must notify the insurer

1 of both the admission and the initial treatment plan
2 within 48 hours of admission. A discharge plan must be
3 fully developed and continuity services prepared to meet
4 the patient's needs and the patient's community preference
5 upon release. Nothing in this paragraph supersedes a
6 health maintenance organization's referral requirement for
7 services from nonparticipating providers upon a patient's
8 discharge from a hospital.

9 (3) Treatment provided under this subsection may be
10 reviewed retrospectively. If coverage is denied
11 retrospectively, neither the insurer nor the participating
12 hospital shall bill, and the insured shall not be liable,
13 for any treatment under this subsection through the date
14 the adverse determination is issued, other than any
15 copayment, coinsurance, or deductible for the stay through
16 that date as applicable under the policy. Coverage shall
17 not be retrospectively denied for the first 72 hours of
18 treatment at a participating hospital except:

19 (A) upon reasonable determination that the
20 inpatient mental health treatment was not provided;

21 (B) upon determination that the patient receiving
22 the treatment was not an insured, enrollee, or
23 beneficiary under the policy;

24 (C) upon material misrepresentation by the patient
25 or health care provider. In this item (C), "material"
26 means a fact or situation that is not merely technical

1 in nature and results or could result in a substantial
2 change in the situation; or

3 (D) upon determination that a service was excluded
4 under the terms of coverage. In that case, the
5 limitation to billing for a copayment, coinsurance, or
6 deductible shall not apply.

7 (4) Nothing in this subsection shall be construed to
8 require a policy to cover any health care service excluded
9 under the terms of coverage.

10 (x) Notwithstanding any provision of this Section, nothing
11 shall require the medical assistance program under Article V
12 of the Illinois Public Aid Code to violate any applicable
13 federal laws, regulations, or grant requirements or any State
14 or federal consent decrees. Nothing in subsection (w) shall
15 prevent the Department of Healthcare and Family Services from
16 requiring a health care provider to use specified level of
17 care, admission, continued stay, or discharge criteria,
18 including, but not limited to, those under Section 5-5.23 of
19 the Illinois Public Aid Code, as long as the Department of
20 Healthcare and Family Services does not require a health care
21 provider to seek prior authorization or concurrent review from
22 the Department of Healthcare and Family Services, a Medicaid
23 managed care organization, or a utilization review
24 organization under the circumstances expressly prohibited by
25 subsection (w).

26 (y) Children's Mental Health. Nothing in this Section

1 shall suspend the screening and assessment requirements for
2 mental health services for children participating in the
3 State's medical assistance program as required in Section
4 5-5.23 of the Illinois Public Aid Code.

5 (Source: P.A. 102-558, eff. 8-20-21; 102-579, eff. 1-1-22;
6 102-813, eff. 5-13-22; 103-426, eff. 8-4-23.)

7 Section 6-10. The Managed Care Reform and Patient Rights
8 Act is amended by changing Sections 10, 45.1, and 85 and by
9 adding Section 87 as follows:

10 (215 ILCS 134/10)

11 Sec. 10. Definitions. In this Act:

12 "Adverse determination" means a determination by a health
13 care plan under Section 45 or by a utilization review program
14 under Section 85 that a health care service is not medically
15 necessary.

16 "Clinical peer" means a health care professional who is in
17 the same profession and the same or similar specialty as the
18 health care provider who typically manages the medical
19 condition, procedures, or treatment under review.

20 "Department" means the Department of Insurance.

21 "Emergency medical condition" means a medical condition
22 manifesting itself by acute symptoms of sufficient severity,
23 regardless of the final diagnosis given, such that a prudent
24 layperson, who possesses an average knowledge of health and

1 medicine, could reasonably expect the absence of immediate
2 medical attention to result in:

3 (1) placing the health of the individual (or, with
4 respect to a pregnant woman, the health of the woman or her
5 unborn child) in serious jeopardy;

6 (2) serious impairment to bodily functions;

7 (3) serious dysfunction of any bodily organ or part;

8 (4) inadequately controlled pain; or

9 (5) with respect to a pregnant woman who is having
10 contractions:

11 (A) inadequate time to complete a safe transfer to
12 another hospital before delivery; or

13 (B) a transfer to another hospital may pose a
14 threat to the health or safety of the woman or unborn
15 child.

16 "Emergency medical screening examination" means a medical
17 screening examination and evaluation by a physician licensed
18 to practice medicine in all its branches, or to the extent
19 permitted by applicable laws, by other appropriately licensed
20 personnel under the supervision of or in collaboration with a
21 physician licensed to practice medicine in all its branches to
22 determine whether the need for emergency services exists.

23 "Emergency services" means, with respect to an enrollee of
24 a health care plan, transportation services, including but not
25 limited to ambulance services, and covered inpatient and
26 outpatient hospital services furnished by a provider qualified

1 to furnish those services that are needed to evaluate or
2 stabilize an emergency medical condition. "Emergency services"
3 does not refer to post-stabilization medical services.

4 "Enrollee" means any person and his or her dependents
5 enrolled in or covered by a health care plan.

6 "Generally accepted standards of care" means standards of
7 care and clinical practice that are generally recognized by
8 health care providers practicing in relevant clinical
9 specialties for the illness, injury, or condition or its
10 symptoms and comorbidities. Valid, evidence-based sources
11 reflecting generally accepted standards of care include
12 peer-reviewed scientific studies and medical literature,
13 recommendations of nonprofit health care provider professional
14 associations and specialty societies, including, but not
15 limited to, patient placement criteria and clinical practice
16 guidelines, recommendations of federal government agencies,
17 and drug labeling approved by the United States Food and Drug
18 Administration.

19 "Health care plan" means a plan, including, but not
20 limited to, a health maintenance organization, a managed care
21 community network as defined in the Illinois Public Aid Code,
22 or an accountable care entity as defined in the Illinois
23 Public Aid Code that receives capitated payments to cover
24 medical services from the Department of Healthcare and Family
25 Services, that establishes, operates, or maintains a network
26 of health care providers that has entered into an agreement

1 with the plan to provide health care services to enrollees to
2 whom the plan has the ultimate obligation to arrange for the
3 provision of or payment for services through organizational
4 arrangements for ongoing quality assurance, utilization review
5 programs, or dispute resolution. Nothing in this definition
6 shall be construed to mean that an independent practice
7 association or a physician hospital organization that
8 subcontracts with a health care plan is, for purposes of that
9 subcontract, a health care plan.

10 For purposes of this definition, "health care plan" shall
11 not include the following:

12 (1) indemnity health insurance policies including
13 those using a contracted provider network;

14 (2) health care plans that offer only dental or only
15 vision coverage;

16 (3) preferred provider administrators, as defined in
17 Section 370g(g) of the Illinois Insurance Code;

18 (4) employee or employer self-insured health benefit
19 plans under the federal Employee Retirement Income
20 Security Act of 1974;

21 (5) health care provided pursuant to the Workers'
22 Compensation Act or the Workers' Occupational Diseases
23 Act; and

24 (6) except with respect to subsections (a) and (b) of
25 Section 65 and subsection (a-5) of Section 70,
26 not-for-profit voluntary health services plans with health

1 maintenance organization authority in existence as of
2 January 1, 1999 that are affiliated with a union and that
3 only extend coverage to union members and their
4 dependents.

5 "Health care professional" means a physician, a registered
6 professional nurse, or other individual appropriately licensed
7 or registered to provide health care services.

8 "Health care provider" means any physician, hospital
9 facility, facility licensed under the Nursing Home Care Act,
10 long-term care facility as defined in Section 1-113 of the
11 Nursing Home Care Act, or other person that is licensed or
12 otherwise authorized to deliver health care services. Nothing
13 in this Act shall be construed to define Independent Practice
14 Associations or Physician-Hospital Organizations as health
15 care providers.

16 "Health care services" means any services included in the
17 furnishing to any individual of medical care, or the
18 hospitalization incident to the furnishing of such care, as
19 well as the furnishing to any person of any and all other
20 services for the purpose of preventing, alleviating, curing,
21 or healing human illness or injury including behavioral
22 health, mental health, home health, and pharmaceutical
23 services and products.

24 "Medical director" means a physician licensed in any state
25 to practice medicine in all its branches appointed by a health
26 care plan.

1 "Medically necessary" means that a service or product
2 addresses the specific needs of a patient for the purpose of
3 screening, preventing, diagnosing, managing, or treating an
4 illness, injury, or condition or its symptoms and
5 comorbidities, including minimizing the progression of an
6 illness, injury, or condition or its symptoms and
7 comorbidities, in a manner that is all of the following:

8 (1) in accordance with generally accepted standards of
9 care;

10 (2) clinically appropriate in terms of type,
11 frequency, extent, site, and duration; and

12 (3) not primarily for the economic benefit of the
13 health care plan, purchaser, or utilization review
14 organization, or for the convenience of the patient,
15 treating physician, or other health care provider.

16 "Person" means a corporation, association, partnership,
17 limited liability company, sole proprietorship, or any other
18 legal entity.

19 "Physician" means a person licensed under the Medical
20 Practice Act of 1987.

21 "Post-stabilization medical services" means health care
22 services provided to an enrollee that are furnished in a
23 licensed hospital by a provider that is qualified to furnish
24 such services, and determined to be medically necessary and
25 directly related to the emergency medical condition following
26 stabilization.

1 "Stabilization" means, with respect to an emergency
2 medical condition, to provide such medical treatment of the
3 condition as may be necessary to assure, within reasonable
4 medical probability, that no material deterioration of the
5 condition is likely to result.

6 "Step therapy requirement" means a utilization review or
7 formulary requirement that specifies, as a condition of
8 coverage under a health care plan, the order in which certain
9 health care services must be used to treat or manage an
10 enrollee's health condition.

11 "Step therapy requirement" does not include:

12 (i) the use of utilization review to identify when a
13 treatment is contraindicated or to limit quantity or
14 dosage for an enrollee based on utilization review
15 criteria consistent with generally accepted standards of
16 care;

17 (ii) the removal of a drug from a formulary or
18 negatively changing a formulary drug's preferred or
19 cost-sharing tier;

20 (iii) the fact that an enrollee or the enrollee's
21 authorized representative must use the medical exceptions
22 process under Section 45.1 of this Act to obtain coverage
23 for a drug that is not concurrently listed on the
24 formulary for the enrollee's health care plan. However, if
25 a medical exceptions procedure requires an enrollee to try
26 a formulary drug before an off-formulary drug, that is a

1 step therapy requirement unless the enrollee or
2 prescribing provider demonstrates that: (1) the formulary
3 drug is not likely to be as effective for the enrollee or
4 has less likelihood of patient compliance with the
5 formulary drug than with the off-formulary drug; (2) the
6 enrollee is already stable on an off-formulary drug; or
7 (3) the formulary drug is contraindicated for the
8 enrollee. Any off-formulary coverage decision during a
9 medical exceptions procedure based on cost is step therapy
10 and prohibited;

11 (iv) a requirement that an enrollee or the enrollee's
12 authorized representative obtain prior authorization for
13 the requested treatment;

14 (v) for health care plans operated or overseen by the
15 Department of Healthcare and Family Services, including
16 Medicaid managed care plans, any utilization controls
17 mandated by 42 CFR 456.703;

18 (vi) the creation and maintenance by the Department of
19 Healthcare and Family Services of a Preferred Drug List,
20 and any requirement that Medicaid managed care
21 organizations comply with the Preferred Drug List
22 utilization control process, as described in Section
23 5-30.14 of the Illinois Public Aid Code; or

24 (vii) the use of utilization review criteria allowed
25 under subsections (c) through (e) of Section 87 of this
26 Act for any health care service other than prescription

1 drugs.

2 "Utilization review" means the evaluation of the medical
3 necessity, appropriateness, and efficiency of the use of
4 health care services, procedures, and facilities.

5 "Utilization review" includes either of the following:

6 (1) prospectively, retrospectively, or concurrently
7 reviewing and approving, modifying, delaying, or denying,
8 based, in whole or in part, on medical necessity, requests
9 by health care providers, enrollees, or their authorized
10 representatives for coverage of health care services
11 before, retrospectively, or concurrently with the
12 provision of health care services to enrollees; or

13 (2) evaluating the medical necessity, appropriateness,
14 level of care, service intensity, efficacy, or efficiency
15 of health care services, benefits, procedures, or
16 settings, under any circumstances, to determine whether a
17 health care service or benefit subject to a medical
18 necessity coverage requirement in a health care plan is
19 covered as medically necessary for an enrollee.

20 "Utilization review criteria" means criteria, standards,
21 protocols, or guidelines used by a utilization review program
22 to conduct utilization review to ensure that a patient's care
23 is aligned with generally accepted standards of care and
24 consistent with State law.

25 "Utilization review program" means a program established
26 by a person to perform utilization review.

1 (Source: P.A. 102-409, eff. 1-1-22; 103-426, eff. 8-4-23.)

2 (215 ILCS 134/45.1)

3 Sec. 45.1. Medical exceptions procedures required.

4 (a) Notwithstanding any other provision of law, on or
5 after January 1, 2018 (the effective date of Public Act
6 99-761), every insurer licensed in this State to sell a policy
7 of group or individual accident and health insurance or a
8 health benefits plan shall establish and maintain a medical
9 exceptions process that allows covered persons or their
10 authorized representatives to request any clinically
11 appropriate prescription drug when (1) the drug is not covered
12 based on the health benefit plan's formulary; (2) the health
13 benefit plan is discontinuing coverage of the drug on the
14 plan's formulary for reasons other than safety or other than
15 because the prescription drug has been withdrawn from the
16 market by the drug's manufacturer; (3) (blank) ~~the~~
17 ~~prescription drug alternatives required to be used in~~
18 ~~accordance with a step therapy requirement (A) has been~~
19 ~~ineffective in the treatment of the enrollee's disease or~~
20 ~~medical condition or, based on both sound clinical evidence~~
21 ~~and medical and scientific evidence, the known relevant~~
22 ~~physical or mental characteristics of the enrollee, and the~~
23 ~~known characteristics of the drug regimen, is likely to be~~
24 ~~ineffective or adversely affect the drug's effectiveness or~~
25 ~~patient compliance or (B) has caused or, based on sound~~

1 ~~medical evidence, is likely to cause an adverse reaction or~~
2 ~~harm to the enrollee;~~ or (4) the number of doses available
3 under a dose restriction for the prescription drug (A) has
4 been ineffective in the treatment of the enrollee's disease or
5 medical condition or (B) based on both sound clinical evidence
6 and medical and scientific evidence, the known relevant
7 physical and mental characteristics of the enrollee, and known
8 characteristics of the drug regimen, is likely to be
9 ineffective or adversely affect the drug's effective or
10 patient compliance.

11 (b) The health carrier's established medical exceptions
12 procedures must require, at a minimum, the following:

13 (1) Any request for approval of coverage made verbally
14 or in writing (regardless of whether made using a paper or
15 electronic form or some other writing) at any time shall
16 be reviewed by appropriate health care professionals.

17 (2) The health carrier must, within 72 hours after
18 receipt of a request made under subsection (a) of this
19 Section, either approve or deny the request. In the case
20 of a denial, the health carrier shall provide the covered
21 person or the covered person's authorized representative
22 and the covered person's prescribing provider with the
23 reason for the denial, an alternative covered medication,
24 if applicable, and information regarding the procedure for
25 submitting an appeal to the denial. A health carrier shall
26 not use the authorization of alternative covered

1 medications under this Section in a manner that
2 effectively creates a step therapy requirement.

3 (3) In the case of an expedited coverage
4 determination, the health carrier must either approve or
5 deny the request within 24 hours after receipt of the
6 request. In the case of a denial, the health carrier shall
7 provide the covered person or the covered person's
8 authorized representative and the covered person's
9 prescribing provider with the reason for the denial, an
10 alternative covered medication, if applicable, and
11 information regarding the procedure for submitting an
12 appeal to the denial.

13 (c) (Blank). ~~A step therapy requirement exception request~~
14 ~~shall be approved if:~~

15 ~~(1) the required prescription drug is contraindicated;~~

16 ~~(2) the patient has tried the required prescription~~
17 ~~drug while under the patient's current or previous health~~
18 ~~insurance or health benefit plan and the prescribing~~
19 ~~provider submits evidence of failure or intolerance; or~~

20 ~~(3) the patient is stable on a prescription drug~~
21 ~~selected by his or her health care provider for the~~
22 ~~medical condition under consideration while on a current~~
23 ~~or previous health insurance or health benefit plan.~~

24 (d) Upon the granting of an exception request, the
25 insurer, health plan, utilization review organization, or
26 other entity shall authorize the coverage for the drug

1 prescribed by the enrollee's treating health care provider, to
2 the extent the prescribed drug is a covered drug under the
3 policy or contract up to the quantity covered.

4 (e) Any approval of a medical exception request made
5 pursuant to this Section shall be honored for 12 months
6 following the date of the approval or until renewal of the
7 plan.

8 (f) Notwithstanding any other provision of this Section,
9 nothing in this Section shall be interpreted or implemented in
10 a manner not consistent with the federal Patient Protection
11 and Affordable Care Act (Public Law 111-148), as amended by
12 the federal Health Care and Education Reconciliation Act of
13 2010 (Public Law 111-152), and any amendments thereto, or
14 regulations or guidance issued under those Acts.

15 (g) Nothing in this Section shall require or authorize the
16 State agency responsible for the administration of the medical
17 assistance program established under the Illinois Public Aid
18 Code to approve, supply, or cover prescription drugs pursuant
19 to the procedure established in this Section.

20 (Source: P.A. 103-154, eff. 6-30-23.)

21 (215 ILCS 134/85)

22 Sec. 85. Utilization review program registration.

23 (a) No person may conduct a utilization review program in
24 this State unless once every 2 years the person registers the
25 utilization review program with the Department and certifies

1 compliance with the Health Utilization Management Standards of
2 the American Accreditation Healthcare Commission (URAC)
3 sufficient to achieve American Accreditation Healthcare
4 Commission (URAC) accreditation or submits evidence of
5 accreditation by the American Accreditation Healthcare
6 Commission (URAC) for its Health Utilization Management
7 Standards. Nothing in this Act shall be construed to require a
8 health care plan or its subcontractors to become American
9 Accreditation Healthcare Commission (URAC) accredited.

10 (b) In addition, the Director of the Department, in
11 consultation with the Director of the Department of Public
12 Health, may certify alternative utilization review standards
13 of national accreditation organizations or entities in order
14 for plans to comply with this Section. Any alternative
15 utilization review standards shall meet or exceed those
16 standards required under subsection (a).

17 (b-5) The Department shall recognize the Accreditation
18 Association for Ambulatory Health Care among the list of
19 accreditors from which utilization organizations may receive
20 accreditation and qualify for reduced registration and renewal
21 fees.

22 (c) The provisions of this Section do not apply to:

23 (1) persons providing utilization review program
24 services only to the federal government;

25 (2) self-insured health plans under the federal
26 Employee Retirement Income Security Act of 1974, however,

1 this Section does apply to persons conducting a
2 utilization review program on behalf of these health
3 plans;

4 (3) hospitals and medical groups performing
5 utilization review activities for internal purposes unless
6 the utilization review program is conducted for another
7 person.

8 Nothing in this Act prohibits a health care plan or other
9 entity from contractually requiring an entity designated in
10 item (3) of this subsection to adhere to the utilization
11 review program requirements of this Act.

12 (d) This registration shall include submission of all of
13 the following information regarding utilization review program
14 activities:

15 (1) The name, address, and telephone number of the
16 utilization review programs.

17 (2) The organization and governing structure of the
18 utilization review programs.

19 (3) The number of lives for which utilization review
20 is conducted by each utilization review program.

21 (4) Hours of operation of each utilization review
22 program.

23 (5) Description of the grievance process for each
24 utilization review program.

25 (6) Number of covered lives for which utilization
26 review was conducted for the previous calendar year for

1 each utilization review program.

2 (7) Written policies and procedures for protecting
3 confidential information according to applicable State and
4 federal laws for each utilization review program.

5 (e) (1) A utilization review program shall have written
6 procedures for assuring that patient-specific information
7 obtained during the process of utilization review will be:

8 (A) kept confidential in accordance with applicable
9 State and federal laws; and

10 (B) shared only with the enrollee, the enrollee's
11 designee, the enrollee's health care provider, and those
12 who are authorized by law to receive the information.

13 Summary data shall not be considered confidential if it
14 does not provide information to allow identification of
15 individual patients or health care providers.

16 (2) Only a clinical peer ~~health care professional~~ may
17 make adverse determinations regarding the medical
18 necessity of health care services during the course of
19 utilization review. Either a health care professional or
20 an accredited algorithmic automated process, or both in
21 combination, may certify the medical necessity of a health
22 care service in accordance with accreditation standards.
23 Nothing in this subsection prohibits an accredited
24 algorithmic automated process from being used to refer a
25 case to a clinical peer for a potential adverse
26 determination.

1 (3) When making retrospective reviews, utilization
2 review programs shall base reviews solely on the medical
3 information available to the attending physician or
4 ordering provider at the time the health care services
5 were provided.

6 (4) When making prospective, concurrent, and
7 retrospective determinations, utilization review programs
8 shall collect only information that is necessary to make
9 the determination and shall not routinely require health
10 care providers to numerically code diagnoses or procedures
11 to be considered for certification, unless required under
12 State or federal Medicare or Medicaid rules or
13 regulations, but may request such code if available, or
14 routinely request copies of medical records of all
15 enrollees reviewed. During prospective or concurrent
16 review, copies of medical records shall only be required
17 when necessary to verify that the health care services
18 subject to review are medically necessary. In these cases,
19 only the necessary or relevant sections of the medical
20 record shall be required.

21 (f) If the Department finds that a utilization review
22 program is not in compliance with this Section, the Department
23 shall issue a corrective action plan and allow a reasonable
24 amount of time for compliance with the plan. If the
25 utilization review program does not come into compliance, the
26 Department may issue a cease and desist order. Before issuing

1 a cease and desist order under this Section, the Department
2 shall provide the utilization review program with a written
3 notice of the reasons for the order and allow a reasonable
4 amount of time to supply additional information demonstrating
5 compliance with requirements of this Section and to request a
6 hearing. The hearing notice shall be sent by certified mail,
7 return receipt requested, and the hearing shall be conducted
8 in accordance with the Illinois Administrative Procedure Act.

9 (g) A utilization review program subject to a corrective
10 action may continue to conduct business until a final decision
11 has been issued by the Department.

12 (h) Any adverse determination made by a health care plan
13 or its subcontractors may be appealed in accordance with
14 subsection (f) of Section 45.

15 (i) The Director may by rule establish a registration fee
16 for each person conducting a utilization review program. All
17 fees paid to and collected by the Director under this Section
18 shall be deposited into the Insurance Producer Administration
19 Fund.

20 (Source: P.A. 99-111, eff. 1-1-16.)

21 (215 ILCS 134/87 new)

22 Sec. 87. General standards for use of utilization review
23 criteria.

24 (a) Except as provided in subsection (h), beginning
25 January 1, 2026, all medical necessity determinations made by

1 a utilization review program shall be conducted in accordance
2 with the requirements of this Section. No policy, contract,
3 certificate, or evidence of coverage issued to any enrollee,
4 nor any formulary, may contain terms or conditions to the
5 contrary.

6 (b) A utilization review program shall base any medical
7 necessity determination or the utilization review criteria
8 that the program applies to determine the medical necessity of
9 health care services and benefits on current generally
10 accepted standards of care.

11 (c) Subject to subsection (i), a utilization review
12 program shall apply the most recent version of:

13 (1) the treatment criteria, at the time the service or
14 treatment was delivered, developed by an unaffiliated
15 nonprofit professional association for the relevant
16 clinical specialty;

17 (2) nationally recognized, evidence-based treatment
18 criteria reflecting current generally accepted standards
19 of care when:

20 (A) such national criteria are developed and
21 updated annually by a third-party entity that does not
22 receive direct payments based on the outcome of the
23 clinical care decisions; and

24 (B) for utilization review programs with respect
25 to health care plans subject to this Act, neither the
26 developing entity nor the utilization review program

1 customizes or adapts such national criteria, and the
2 developing entity does not offer the utilization
3 review program a choice the among more than one
4 distinct set of criteria for the same health care
5 service, except to the extent necessary for all
6 utilization review programs subject to this Section to
7 comply with State or federal requirements applicable
8 to each health care plan that they offer or administer
9 as provided in subsection (i); or

10 (3) for health care plans operated or overseen by the
11 Department of Healthcare and Family Services, including
12 Medicaid managed care plans, when neither of the preceding
13 types of sources offers treatment criteria for a covered
14 item or service, treatment criteria determined by the
15 Department of Healthcare and Family Services that are not
16 inconsistent with generally accepted standards of care.

17 (d) For medical necessity determinations that are within
18 the scope of the sources specified in subsection (c), a
19 utilization review program shall not apply different,
20 additional, conflicting, or more restrictive utilization
21 review criteria than the criteria set forth in those sources.
22 For all level of care placement decisions, the utilization
23 review program or health care plan shall authorize placement
24 at the level of care consistent with the assessment of the
25 enrollee using the relevant patient placement criteria as
26 specified in subsection (c). If that level of placement is not

1 available, the utilization review program or health care plan
2 shall authorize the next highest level of care. In the event of
3 disagreement, the utilization review program shall provide
4 full detail of its assessment using the relevant criteria as
5 specified in subsection (c) to the provider of the service and
6 the patient.

7 (e) If a utilization review program conducts utilization
8 review that is outside the scope of the criteria specified in
9 subsection (c) or that relates to the advancements in
10 technology or in the types or levels of care that are not
11 addressed in the most recent versions of the sources specified
12 in subsection (c), then the utilization review program shall
13 conduct utilization review in accordance with subsection (b).
14 If a utilization review program purchases or licenses
15 utilization review criteria pursuant to this subsection, then
16 the utilization review program shall verify and document
17 before use that the criteria were developed in accordance with
18 subsection (b).

19 (f) To ensure the proper use of utilization review
20 criteria that were not developed under or that diverge from
21 those developed under subsection (c), every health care plan
22 shall do all of the following:

23 (1) Make an educational program available to the
24 health care plan's staff, as well as the staff of any other
25 utilization review program contracted to review claims,
26 conduct utilization reviews, or make medical necessity

1 determinations about the utilization review criteria.

2 (2) Make the educational program available, at no
3 cost, to other stakeholders, including the health care
4 plan's participating or contracted providers and potential
5 enrollees. The education program must be provided at least
6 once a year, in person or digitally, or recordings of the
7 education program must be made available to those
8 stakeholders.

9 (3) Provide, at no cost, the utilization review
10 criteria and any training material or resources to
11 providers and enrollees upon request. The health care plan
12 may place the criteria on a secure, password-protected
13 website so long as the access requirements of the website
14 do not unreasonably restrict access to enrollees or their
15 providers. No restrictions shall be placed upon the
16 enrollee's or treating provider's access right to
17 utilization review criteria obtained under this paragraph
18 at any point in time, including before an initial request
19 for authorization.

20 (4) Track, identify, and analyze how the utilization
21 review criteria are used to certify care, deny care, and
22 support the appeals process.

23 (5) Conduct interrater reliability testing to ensure
24 consistency in utilization review decision-making that
25 covers how medical necessity decisions are made. This
26 assessment shall cover all aspects of utilization review

1 as defined in Section 10.

2 (6) Run interrater reliability reports about how the
3 clinical guidelines are used in conjunction with the
4 utilization review process.

5 (7) Achieve interrater reliability pass rates of at
6 least 90% and, if this threshold is not met, immediately
7 provide for the remediation of poor interrater reliability
8 and interrater reliability testing for all new staff
9 before they can conduct utilization review without
10 supervision.

11 (8) Maintain documentation of interrater reliability
12 testing and the remediation actions taken for those with
13 pass rates lower than 90% and annually submit to the
14 Department of Insurance or, in the case of Medicaid
15 managed care organizations, the Department of Healthcare
16 and Family Services the testing results and a summary of
17 remedial actions. The reports shall be confidential, not
18 subject to subpoena, and not subject to disclosure under
19 the Freedom of Information Act.

20 (g) No utilization review program or any policy, contract,
21 certificate, evidence of coverage, or formulary shall impose
22 step therapy requirements. Nothing in this subsection
23 prohibits a health care plan, by contract, written policy or
24 procedure, or any other agreement or course of conduct, from
25 requiring a pharmacist to effect substitutions of prescription
26 drugs consistent with Section 19.5 of the Pharmacy Practice

1 Act, under which a pharmacist may substitute an
2 interchangeable biologic for a prescribed biologic product,
3 and Section 25 of the Pharmacy Practice Act, under which a
4 pharmacist may select a generic drug determined to be
5 therapeutically equivalent by the United States Food and Drug
6 Administration and in accordance with the Illinois Food, Drug
7 and Cosmetic Act. For health care plans operated or overseen
8 by the Department of Healthcare and Family Services, including
9 Medicaid managed care plans, the prohibition in this
10 subsection does not apply to step therapy requirements for
11 drugs that do not appear on the most recent Preferred Drug List
12 published by the Department of Healthcare and Family Services.

13 (h) Except for subsection (g), this Section does not apply
14 to utilization review concerning diagnosis, prevention, and
15 treatment of mental, emotional, nervous, or substance use
16 disorders or conditions, which shall be governed by Section
17 370c of the Illinois Insurance Code.

18 (i) Nothing in this Section shall be construed to
19 supersede or waive requirements provided under any other State
20 or federal law or federal regulation that any coverage subject
21 to this Section comply with specific utilization review
22 criteria for a specific illness, level of care placement,
23 injury, or condition or its symptoms and comorbidities.

24 Section 6-15. The Health Carrier External Review Act is
25 amended by changing Section 10 as follows:

1 (215 ILCS 180/10)

2 Sec. 10. Definitions. For the purposes of this Act:

3 "Adverse determination" means:

4 (1) a determination by a health carrier or its
5 designee utilization review organization that, based upon
6 the information provided, a request for a benefit under
7 the health carrier's health benefit plan upon application
8 of any utilization review technique does not meet the
9 health carrier's requirements for medical necessity,
10 appropriateness, health care setting, level of care, or
11 effectiveness or is determined to be experimental or
12 investigational and the requested benefit is therefore
13 denied, reduced, or terminated or payment is not provided
14 or made, in whole or in part, for the benefit;

15 (2) the denial, reduction, or termination of or
16 failure to provide or make payment, in whole or in part,
17 for a benefit based on a determination by a health carrier
18 or its designee utilization review organization that a
19 preexisting condition was present before the effective
20 date of coverage; or

21 (3) a rescission of coverage determination, which does
22 not include a cancellation or discontinuance of coverage
23 that is attributable to a failure to timely pay required
24 premiums or contributions towards the cost of coverage.

25 "Authorized representative" means:

1 (1) a person to whom a covered person has given
2 express written consent to represent the covered person
3 for purposes of this Law;

4 (2) a person authorized by law to provide substituted
5 consent for a covered person;

6 (3) a family member of the covered person or the
7 covered person's treating health care professional when
8 the covered person is unable to provide consent;

9 (4) a health care provider when the covered person's
10 health benefit plan requires that a request for a benefit
11 under the plan be initiated by the health care provider;
12 or

13 (5) in the case of an urgent care request, a health
14 care provider with knowledge of the covered person's
15 medical condition.

16 "Best evidence" means evidence based on:

17 (1) randomized clinical trials;

18 (2) if randomized clinical trials are not available,
19 then cohort studies or case-control studies;

20 (3) if items (1) and (2) are not available, then
21 case-series; or

22 (4) if items (1), (2), and (3) are not available, then
23 expert opinion.

24 "Case-series" means an evaluation of a series of patients
25 with a particular outcome, without the use of a control group.

26 "Clinical review criteria" means the written screening

1 procedures, decision abstracts, clinical protocols, and
2 practice guidelines used by a health carrier to determine the
3 necessity and appropriateness of health care services.
4 "Clinical review criteria" includes all utilization review
5 criteria as defined in Section 10 of the Managed Care Reform
6 and Patient Rights Act.

7 "Cohort study" means a prospective evaluation of 2 groups
8 of patients with only one group of patients receiving specific
9 intervention.

10 "Concurrent review" means a review conducted during a
11 patient's stay or course of treatment in a facility, the
12 office of a health care professional, or other inpatient or
13 outpatient health care setting.

14 "Covered benefits" or "benefits" means those health care
15 services to which a covered person is entitled under the terms
16 of a health benefit plan.

17 "Covered person" means a policyholder, subscriber,
18 enrollee, or other individual participating in a health
19 benefit plan.

20 "Director" means the Director of the Department of
21 Insurance.

22 "Emergency medical condition" means a medical condition
23 manifesting itself by acute symptoms of sufficient severity,
24 including, but not limited to, severe pain, such that a
25 prudent layperson who possesses an average knowledge of health
26 and medicine could reasonably expect the absence of immediate

1 medical attention to result in:

2 (1) placing the health of the individual or, with
3 respect to a pregnant woman, the health of the woman or her
4 unborn child, in serious jeopardy;

5 (2) serious impairment to bodily functions; or

6 (3) serious dysfunction of any bodily organ or part.

7 "Emergency services" means health care items and services
8 furnished or required to evaluate and treat an emergency
9 medical condition.

10 "Evidence-based standard" means the conscientious,
11 explicit, and judicious use of the current best evidence based
12 on an overall systematic review of the research in making
13 decisions about the care of individual patients.

14 "Expert opinion" means a belief or an interpretation by
15 specialists with experience in a specific area about the
16 scientific evidence pertaining to a particular service,
17 intervention, or therapy.

18 "Facility" means an institution providing health care
19 services or a health care setting.

20 "Final adverse determination" means an adverse
21 determination involving a covered benefit that has been upheld
22 by a health carrier, or its designee utilization review
23 organization, at the completion of the health carrier's
24 internal grievance process procedures as set forth by the
25 Managed Care Reform and Patient Rights Act.

26 "Health benefit plan" means a policy, contract,

1 certificate, plan, or agreement offered or issued by a health
2 carrier to provide, deliver, arrange for, pay for, or
3 reimburse any of the costs of health care services.

4 "Health care provider" or "provider" means a physician,
5 hospital facility, or other health care practitioner licensed,
6 accredited, or certified to perform specified health care
7 services consistent with State law, responsible for
8 recommending health care services on behalf of a covered
9 person.

10 "Health care services" means services for the diagnosis,
11 prevention, treatment, cure, or relief of a health condition,
12 illness, injury, or disease.

13 "Health carrier" means an entity subject to the insurance
14 laws and regulations of this State, or subject to the
15 jurisdiction of the Director, that contracts or offers to
16 contract to provide, deliver, arrange for, pay for, or
17 reimburse any of the costs of health care services, including
18 a sickness and accident insurance company, a health
19 maintenance organization, or any other entity providing a plan
20 of health insurance, health benefits, or health care services.

21 "Health carrier" also means Limited Health Service
22 Organizations (LHSO) and Voluntary Health Service Plans.

23 "Health information" means information or data, whether
24 oral or recorded in any form or medium, and personal facts or
25 information about events or relationships that relate to:

26 (1) the past, present, or future physical, mental, or

1 behavioral health or condition of an individual or a
2 member of the individual's family;

3 (2) the provision of health care services to an
4 individual; or

5 (3) payment for the provision of health care services
6 to an individual.

7 "Independent review organization" means an entity that
8 conducts independent external reviews of adverse
9 determinations and final adverse determinations.

10 "Medical or scientific evidence" means evidence found in
11 the following sources:

12 (1) peer-reviewed scientific studies published in or
13 accepted for publication by medical journals that meet
14 nationally recognized requirements for scientific
15 manuscripts and that submit most of their published
16 articles for review by experts who are not part of the
17 editorial staff;

18 (2) peer-reviewed medical literature, including
19 literature relating to therapies reviewed and approved by
20 a qualified institutional review board, biomedical
21 compendia, and other medical literature that meet the
22 criteria of the National Institutes of Health's Library of
23 Medicine for indexing in Index Medicus (Medline) and
24 Elsevier Science Ltd. for indexing in Excerpta Medicus
25 (EMBASE);

26 (3) medical journals recognized by the Secretary of

1 Health and Human Services under Section 1861(t)(2) of the
2 federal Social Security Act;

3 (4) the following standard reference compendia:

4 (a) The American Hospital Formulary Service-Drug
5 Information;

6 (b) Drug Facts and Comparisons;

7 (c) The American Dental Association Accepted
8 Dental Therapeutics; and

9 (d) The United States Pharmacopoeia-Drug
10 Information;

11 (5) findings, studies, or research conducted by or
12 under the auspices of federal government agencies and
13 nationally recognized federal research institutes,
14 including:

15 (a) the federal Agency for Healthcare Research and
16 Quality;

17 (b) the National Institutes of Health;

18 (c) the National Cancer Institute;

19 (d) the National Academy of Sciences;

20 (e) the Centers for Medicare & Medicaid Services;

21 (f) the federal Food and Drug Administration; and

22 (g) any national board recognized by the National
23 Institutes of Health for the purpose of evaluating the
24 medical value of health care services; or

25 (6) any other medical or scientific evidence that is
26 comparable to the sources listed in items (1) through (5).

1 "Person" means an individual, a corporation, a
2 partnership, an association, a joint venture, a joint stock
3 company, a trust, an unincorporated organization, any similar
4 entity, or any combination of the foregoing.

5 "Prospective review" means a review conducted prior to an
6 admission or the provision of a health care service or a course
7 of treatment in accordance with a health carrier's requirement
8 that the health care service or course of treatment, in whole
9 or in part, be approved prior to its provision.

10 "Protected health information" means health information
11 (i) that identifies an individual who is the subject of the
12 information; or (ii) with respect to which there is a
13 reasonable basis to believe that the information could be used
14 to identify an individual.

15 "Randomized clinical trial" means a controlled prospective
16 study of patients that have been randomized into an
17 experimental group and a control group at the beginning of the
18 study with only the experimental group of patients receiving a
19 specific intervention, which includes study of the groups for
20 variables and anticipated outcomes over time.

21 "Retrospective review" means any review of a request for a
22 benefit that is not a concurrent or prospective review
23 request. "Retrospective review" does not include the review of
24 a claim that is limited to veracity of documentation or
25 accuracy of coding.

26 "Utilization review" has the meaning provided by the

1 Managed Care Reform and Patient Rights Act.

2 "Utilization review organization" means a utilization
3 review program as defined in the Managed Care Reform and
4 Patient Rights Act.

5 (Source: P.A. 97-574, eff. 8-26-11; 97-813, eff. 7-13-12;
6 98-756, eff. 7-16-14.)

7 Section 6-20. The Prior Authorization Reform Act is
8 amended by changing Sections 15 and 20 as follows:

9 (215 ILCS 200/15)

10 Sec. 15. Definitions. As used in this Act:

11 "Adverse determination" has the meaning given to that term
12 in Section 10 of the Health Carrier External Review Act.

13 "Appeal" means a formal request, either orally or in
14 writing, to reconsider an adverse determination.

15 "Approval" means a determination by a health insurance
16 issuer or its contracted utilization review organization that
17 a health care service has been reviewed and, based on the
18 information provided, satisfies the health insurance issuer's
19 or its contracted utilization review organization's
20 requirements for medical necessity and appropriateness.

21 "Clinical review criteria" has the meaning given to that
22 term in Section 10 of the Health Carrier External Review Act.

23 "Department" means the Department of Insurance.

24 "Emergency medical condition" has the meaning given to

1 that term in Section 10 of the Managed Care Reform and Patient
2 Rights Act.

3 "Emergency services" has the meaning given to that term in
4 federal health insurance reform requirements for the group and
5 individual health insurance markets, 45 CFR 147.138.

6 "Enrollee" has the meaning given to that term in Section
7 10 of the Managed Care Reform and Patient Rights Act.

8 "Health care professional" has the meaning given to that
9 term in Section 10 of the Managed Care Reform and Patient
10 Rights Act.

11 "Health care provider" has the meaning given to that term
12 in Section 10 of the Managed Care Reform and Patient Rights
13 Act, except that facilities licensed under the Nursing Home
14 Care Act and long-term care facilities as defined in Section
15 1-113 of the Nursing Home Care Act are excluded from this Act.

16 "Health care service" means any services or level of
17 services included in the furnishing to an individual of
18 medical care or the hospitalization incident to the furnishing
19 of such care, as well as the furnishing to any person of any
20 other services for the purpose of preventing, alleviating,
21 curing, or healing human illness or injury, including
22 behavioral health, mental health, home health, and
23 pharmaceutical services and products.

24 "Health insurance issuer" has the meaning given to that
25 term in Section 5 of the Illinois Health Insurance Portability
26 and Accountability Act.

1 "Medically necessary" has the meaning given to that term
2 in Section 10 of the Managed Care Reform and Patient Rights
3 Act. ~~means a health care professional exercising prudent~~
4 ~~clinical judgment would provide care to a patient for the~~
5 ~~purpose of preventing, diagnosing, or treating an illness,~~
6 ~~injury, disease, or its symptoms and that are: (i) in~~
7 ~~accordance with generally accepted standards of medical~~
8 ~~practice; (ii) clinically appropriate in terms of type,~~
9 ~~frequency, extent, site, and duration and are considered~~
10 ~~effective for the patient's illness, injury, or disease; and~~
11 ~~(iii) not primarily for the convenience of the patient,~~
12 ~~treating physician, other health care professional, caregiver,~~
13 ~~family member, or other interested party, but focused on what~~
14 ~~is best for the patient's health outcome.~~

15 "Physician" means a person licensed under the Medical
16 Practice Act of 1987 or licensed under the laws of another
17 state to practice medicine in all its branches.

18 "Prior authorization" means the process by which health
19 insurance issuers or their contracted utilization review
20 organizations determine the medical necessity and medical
21 appropriateness of otherwise covered health care services
22 before the rendering of such health care services. "Prior
23 authorization" includes any health insurance issuer's or its
24 contracted utilization review organization's requirement that
25 an enrollee, health care professional, or health care provider
26 notify the health insurance issuer or its contracted

1 utilization review organization before, at the time of, or
2 concurrent to providing a health care service.

3 "Urgent health care service" means a health care service
4 with respect to which the application of the time periods for
5 making a non-expedited prior authorization that in the opinion
6 of a health care professional with knowledge of the enrollee's
7 medical condition:

8 (1) could seriously jeopardize the life or health of
9 the enrollee or the ability of the enrollee to regain
10 maximum function; or

11 (2) could subject the enrollee to severe pain that
12 cannot be adequately managed without the care or treatment
13 that is the subject of the utilization review.

14 "Urgent health care service" does not include emergency
15 services.

16 "Utilization review organization" has the meaning given to
17 that term in 50 Ill. Adm. Code 4520.30.

18 (Source: P.A. 102-409, eff. 1-1-22.)

19 (215 ILCS 200/20)

20 Sec. 20. Disclosure and review of prior authorization
21 requirements.

22 (a) A health insurance issuer shall maintain a complete
23 list of services for which prior authorization is required,
24 including for all services where prior authorization is
25 performed by an entity under contract with the health

1 insurance issuer. The health insurance issuer shall publish
2 this list on its public website without requiring a member of
3 the general public to create any account or enter any
4 credentials to access it. The list described in this
5 subsection is not required to contain the clinical review
6 criteria applicable to these services.

7 (b) A health insurance issuer shall make any current prior
8 authorization requirements and restrictions, including the
9 written clinical review criteria, readily accessible and
10 conspicuously posted on its website to enrollees, health care
11 professionals, and health care providers. Content published by
12 a third party and licensed for use by a health insurance issuer
13 or its contracted utilization review organization may be made
14 available through the health insurance issuer's or its
15 contracted utilization review organization's secure,
16 password-protected website so long as the access requirements
17 of the website do not unreasonably restrict access.
18 Requirements shall be described in detail, written in easily
19 understandable language, and readily available to the health
20 care professional and health care provider at the point of
21 care. The website shall indicate for each service subject to
22 prior authorization:

23 (1) when prior authorization became required for
24 policies issued or delivered in Illinois, including the
25 effective date or dates and the termination date or dates,
26 if applicable, in Illinois;

1 (2) the date the Illinois-specific requirement was
2 listed on the health insurance issuer's or its contracted
3 utilization review organization's website;

4 (3) where applicable, the date that prior
5 authorization was removed for Illinois; and

6 (4) where applicable, access to a standardized
7 electronic prior authorization request transaction
8 process.

9 (c) The clinical review criteria must:

10 (1) be based on nationally recognized, generally
11 accepted standards except where State law provides its own
12 standard;

13 (2) be developed in accordance with the current
14 standards of a national medical accreditation entity;

15 (3) ensure quality of care and access to needed health
16 care services;

17 (4) be evidence-based;

18 (5) be sufficiently flexible to allow deviations from
19 norms when justified on a case-by-case basis; and

20 (6) be evaluated and updated, if necessary, at least
21 annually.

22 (d) A health insurance issuer shall not deny a claim for
23 failure to obtain prior authorization if the prior
24 authorization requirement was not in effect on the date of
25 service on the claim.

26 (e) A health insurance issuer or its contracted

1 utilization review organization shall not deem as incidental
2 or deny supplies or health care services that are routinely
3 used as part of a health care service when:

4 (1) an associated health care service has received
5 prior authorization; or

6 (2) prior authorization for the health care service is
7 not required.

8 (f) If a health insurance issuer intends either to
9 implement a new prior authorization requirement or restriction
10 or amend an existing requirement or restriction, the health
11 insurance issuer shall provide contracted health care
12 professionals and contracted health care providers of
13 enrollees written notice of the new or amended requirement or
14 amendment no less than 60 days before the requirement or
15 restriction is implemented. The written notice may be provided
16 in an electronic format, including email or facsimile, if the
17 health care professional or health care provider has agreed in
18 advance to receive notices electronically. The health
19 insurance issuer shall ensure that the new or amended
20 requirement is not implemented unless the health insurance
21 issuer's or its contracted utilization review organization's
22 website has been updated to reflect the new or amended
23 requirement or restriction.

24 (g) Entities using prior authorization shall make
25 statistics available regarding prior authorization approvals
26 and denials on their website in a readily accessible format.

1 The statistics must be updated annually and include all of the
2 following information:

3 (1) a list of all health care services, including
4 medications, that are subject to prior authorization;

5 (2) the total number of prior authorization requests
6 received;

7 (3) the number of prior authorization requests denied
8 during the previous plan year by the health insurance
9 issuer or its contracted utilization review organization
10 with respect to each service described in paragraph (1)
11 and the top 5 reasons for denial;

12 (4) the number of requests described in paragraph (3)
13 that were appealed, the number of the appealed requests
14 that upheld the adverse determination, and the number of
15 appealed requests that reversed the adverse determination;

16 (5) the average time between submission and response;
17 and

18 (6) any other information as the Director determines
19 appropriate.

20 (Source: P.A. 102-409, eff. 1-1-22.)

21 Section 6-25. The Illinois Public Aid Code is amended by
22 changing Section 5-16.12 as follows:

23 (305 ILCS 5/5-16.12)

24 Sec. 5-16.12. Managed Care Reform and Patient Rights Act.

1 The medical assistance program and other programs administered
2 by the Department are subject to the provisions of the Managed
3 Care Reform and Patient Rights Act. The Department may adopt
4 rules to implement those provisions. These rules shall require
5 compliance with that Act in the medical assistance managed
6 care programs and other programs administered by the
7 Department. The medical assistance fee-for-service program is
8 not subject to the provisions of the Managed Care Reform and
9 Patient Rights Act, except for Sections 85 and 87 of the
10 Managed Care Reform and Patient Rights Act and for any
11 definition in Section 10 of the Managed Care Reform and
12 Patient Rights Act that applies to Sections 85 and 87 of the
13 Managed Care Reform and Patient Rights Act.

14 Nothing in the Managed Care Reform and Patient Rights Act
15 shall be construed to mean that the Department is a health care
16 plan as defined in that Act simply because the Department
17 enters into contractual relationships with health care plans;
18 provided that this clause shall not defeat the applicability
19 of Sections 10, 85, and 87 of the Managed Care Reform and
20 Patient Rights Act to the fee-for-service program.

21 (Source: P.A. 91-617, eff. 1-1-00.)

22 Article 99.

23 Section 99-95. No acceleration or delay. Where this Act
24 makes changes in a statute that is represented in this Act by

1 text that is not yet or no longer in effect (for example, a
2 Section represented by multiple versions), the use of that
3 text does not accelerate or delay the taking effect of (i) the
4 changes made by this Act or (ii) provisions derived from any
5 other Public Act.

6 Section 99-99. Effective date. This Act takes effect
7 January 1, 2025.".