AN ACT

Relating to health care providers; creating new provisions; and amending ORS 413.032, 413.037, 413.101, 413.181, 415.013, 415.019 and 415.103.

Be It Enacted by the People of the State of Oregon:

SECTION 1. As used in this section and sections 2 and 3 of this 2021 Act:
(1) “Corporate affiliation” has the meaning prescribed by the Oregon Health Authority by rule, including:
(a) Any relationship between two organizations that reflects, directly or indirectly, a partial or complete controlling interest or partial or complete corporate control; and
(b) Transactions that merge tax identification numbers or corporate governance.
(2) “Essential services” means:
(a) Services that are funded on the prioritized list described in ORS 414.690; and
(b) Services that are essential to achieve health equity.
(3) “Health benefit plan” has the meaning given that term in ORS 743B.005.
(4)(a) “Health care entity” includes:
(A) An individual health professional licensed or certified in this state;
(B) A hospital, as defined in ORS 442.015, or hospital system, as defined by the authority by rule;
(C) A carrier, as defined in ORS 743B.005, that offers a health benefit plan in this state;
(D) A Medicare Advantage plan;
(E) A coordinated care organization or a prepaid managed care health services organization, as both terms are defined in ORS 414.025; and
(F) Any other entity that has as a primary function the provision of health care items or services or that is a parent organization of, or is an entity closely related to, an entity that has as a primary function the provision of health care items or services.
(b) “Health care entity” does not include:
(A) Long term care facilities, as defined in ORS 442.015.
(B) Facilities licensed and operated under ORS 443.400 to 443.455.
(5) “Health equity” has the meaning prescribed by the Oregon Health Policy Board and adopted by the authority by rule.
(6)(a) “Material change transaction” means:
(A) A transaction in which at least one party had average revenue of $25 million or more in the preceding three fiscal years and another party:
   (i) Had an average revenue of at least $10 million in the preceding three fiscal years; or
   (ii) In the case of a new entity, is projected to have at least $10 million in revenue in the first full year of operation at normal levels of utilization or operation as prescribed by the authority by rule.

(B) If a transaction involves a health care entity in this state and an out-of-state entity, a transaction that otherwise qualifies as a material change transaction under this paragraph that may result in increases in the price of health care or limit access to health care services in this state.

(b) “Material change transaction” does not include:
   (A) A clinical affiliation of health care entities formed for the purpose of collaborating on clinical trials or graduate medical education programs.
   (B) A medical services contract or an extension of a medical services contract.
   (C) An affiliation that:
      (i) Does not impact the corporate leadership, governance or control of an entity; and
      (ii) Is necessary, as prescribed by the authority by rule, to adopt advanced value-based payment methodologies to meet the health care cost growth targets under ORS 442.386.
   (D) Contracts under which one health care entity, for and on behalf of a second health care entity, provides patient care and services or provides administrative services relating to, supporting or facilitating the provision of patient care and services, if the second health care entity:
      (i) Maintains responsibility, oversight and control over the patient care and services; and
      (ii) Bills and receives reimbursement for the patient care and services.
   (E) Transactions in which a participant that is a health center as defined in 42 U.S.C. 254b, while meeting all of the participant's obligations, acquires, affiliates with, partners with or enters into any agreement with another entity unless the transaction would result in the participant no longer qualifying as a health center under 42 U.S.C. 254b.
(7)(a) “Medical services contract” means a contract to provide medical or mental health services entered into by:
   (A) A carrier and an independent practice association;
   (B) A carrier, coordinated care organization, independent practice association or network of providers and one or more providers, as defined in ORS 743B.001;
   (C) An independent practice association and an individual health professional or an organization of health care providers;
   (D) Medical, dental, vision or mental health clinics; or
   (E) A medical, dental, vision or mental health clinic and an individual health professional to provide medical, dental, vision or mental health services.
(b) “Medical services contract” does not include a contract of employment or a contract creating a legal entity and ownership of the legal entity that is authorized under ORS chapter 58, 60 or 70 or under any other law authorizing the creation of a professional organization similar to those authorized by ORS chapter 58, 60 or 70, as may be prescribed by the authority by rule.
(8) “Net patient revenue” means the total amount of revenue, after allowance for contractual amounts, charity care and bad debt, received for patient care and services, including:
   (a) Value-based payments;
   (b) Incentive payments;
   (c) Capitation payments or payments under any similar contractual arrangement for the prepayment or reimbursement of patient care and services; and
   (d) Any payment received by a hospital to reimburse a hospital assessment under ORS 414.855.
(9) “Revenue” means:
(a) Net patient revenue; or
(b) The gross amount of premiums received by a health care entity that are derived from health benefit plans.
(10) “Transaction” means:
(a) A merger of a health care entity with another entity;
(b) An acquisition of one or more health care entities by another entity;
(c) New contracts, new clinical affiliations and new contracting affiliations that will eliminate or significantly reduce, as defined by the authority by rule, essential services;
(d) A corporate affiliation involving at least one health care entity; or
(e) Transactions to form a new partnership, joint venture, accountable care organization, parent organization or management services organization, as prescribed by the authority by rule.

SECTION 2. (1) The purpose of this section is to promote the public interest and to advance the goals set forth in ORS 414.018 and the goals of the Oregon Integrated and Coordinated Health Care Delivery System described in ORS 414.570.
(2) In accordance with subsection (1) of this section, the Oregon Health Authority shall adopt by rule criteria approved by the Oregon Health Policy Board for the consideration of requests by health care entities to engage in a material change transaction and procedures for the review of material change transactions under this section.
(3)(a) A notice of a material change transaction involving the sale, merger or acquisition of a domestic health insurer shall be submitted to the Department of Consumer and Business Services as an addendum to filings required by ORS 732.517 to 732.546 or 732.576. The department shall provide to the authority the notice submitted under this subsection to enable the authority to conduct a review in accordance with subsections (5) and (7) of this section. The authority shall notify the department of the outcome of the authority's review.
(b) The department shall make the final determination in material change transactions involving the sale, merger or acquisition of a domestic health insurer and shall coordinate with the authority to incorporate the authority's review into the department's final determination.
(4) An entity shall submit to the authority a notice of a material change transaction, other than a transaction described in subsection (3) of this section, in the form and manner prescribed by the authority, no less than 180 days before the date of the transaction and shall pay a fee prescribed in section 4 of this 2021 Act.
(5) No later than 30 days after receiving a notice described in subsections (3) and (4) of this section, the authority shall conduct a preliminary review to determine if the transaction has the potential to have a negative impact on access to affordable health care in this state and meets the criteria in subsection (9) of this section.
(6) Following a preliminary review, the authority or the department shall approve a transaction or approve a transaction with conditions designed to further the goals described in subsection (1) of this section based on criteria prescribed by the authority by rule, including but not limited to:
(a) If the transaction is in the interest of consumers and is urgently necessary to maintain the solvency of an entity involved in the transaction; or
(b) If the authority determines that the transaction does not have the potential to have a negative impact on access to affordable health care in this state or the transaction is likely to meet the criteria in subsection (9) of this section.
(7)(a) Except as provided in paragraph (b) of this subsection, if a transaction does not meet the criteria in subsection (6) of this section, the authority shall conduct a comprehensive review and may appoint a review board of stakeholders to conduct a comprehensive review and make recommendations as provided in subsections (11) to (18) of this section. The
authority shall complete the comprehensive review no later than 180 days after receipt of the notice unless the parties to the transaction agree to an extension of time.

(b) The authority or the department may intervene in a transaction described in section 1 (6)(a)(C) of this 2021 Act in which the final authority rests with another state and, if the transaction is approved by the other state, may place conditions on health care entities operating in this state with respect to the insurance or health care industry market in this state, prices charged to patients residing in this state and the services available in health care facilities in this state, to serve the public good.

(8) The authority shall prescribe by rule:

(a) Criteria to exempt an entity from the requirements of subsection (4) of this section if there is an emergency situation that threatens immediate care services and the transaction is urgently needed to protect the interest of consumers;

(b) Provision for the authority's failure to complete a review under subsection (5) of this section within 30 days; and

(c) Criteria for when to conduct a comprehensive review and appoint a review board under subsection (7) of this section that must include, but is not limited to:

(A) The potential loss or change in access to essential services;

(B) The potential to impact a large number of residents in this state; or

(C) A significant change in the market share of an entity involved in the transaction.

(9) A health care entity may engage in a material change transaction if, following a comprehensive review conducted by the authority and recommendations by a review board appointed under subsection (7) of this section, the authority determines that the transaction meets the criteria adopted by the department by rule under subsection (2) of this section and:

(a) (A) The parties to the transaction demonstrate that the transaction will benefit the public good and communities by:

(i) Reducing the growth in patient costs in accordance with the health care cost growth targets established under ORS 442.386 or maintain a rate of cost growth that exceeds the target that the entity demonstrates is the best interest of the public;

(ii) Increasing access to services in medically underserved areas; or

(iii) Rectifying historical and contemporary factors contributing to a lack of health equities or access to services; or

(B) The transaction will improve health outcomes for residents of this state; and

(b) There is no substantial likelihood of anticompetitive effects from the transaction that outweigh the benefits of the transaction in increasing or maintaining services to underserved populations.

(10) The authority may suspend a proposed material change transaction if necessary to conduct an examination and complete an analysis of whether the transaction is consistent with subsection (9) of this section and the criteria adopted by rule under subsection (2) of this section.

(11)(a) A review board convened by the authority under subsection (7) of this section must consist of members of the affected community, consumer advocates and health care experts. No more than one-third of the members of the review board may be representatives of institutional health care providers. The authority may not appoint to a review board an individual who is employed by an entity that is a party to the transaction that is under review or is employed by a competitor that is of a similar size to an entity that is a party to the transaction.

(b) A member of a review board shall file a notice of conflict of interest and the notice shall be made public.

(12) The authority may request additional information from an entity that is a party to the material change transaction, and the entity shall promptly reply using the form of

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communication requested by the authority and verified by an officer of the entity if required by the authority.

(13)(a) An entity may not refuse to provide documents or other information requested under subsection (4) or (12) of this section on the grounds that the information is confidential.

(b) Material that is privileged or confidential may not be publicly disclosed if:
(A) The authority determines that disclosure of the material would cause harm to the public;
(B) The material may not be disclosed under ORS 192.311 to 192.478; or
(C) The material is not subject to disclosure under ORS 705.137.

(c) The authority shall maintain the confidentiality of all confidential information and documents that are not publicly available that are obtained in relation to a material change transaction and may not disclose the information or documents to any person, including a member of the review board, without the consent of the person who provided the information or document. Information and documents described in this paragraph are exempt from disclosure under ORS 192.311 to 192.478.

(14) The authority or the Department of Justice may retain actuaries, accountants or other professionals independent of the authority who are qualified and have expertise in the type of material change transaction under review as necessary to assist the authority in conducting the analysis of a proposed material change transaction. The authority or the Department of Justice shall designate the party or parties to the material change transaction that shall bear the reasonable and actual cost of retaining the professionals.

(15) A review board may hold up to two public hearings to seek public input and otherwise engage the public before making a determination on the proposed transaction. A public hearing must be held in the service area or areas of the health care entities that are parties to the material change transaction. At least 10 days prior to the public hearing, the authority shall post to the authority's website information about the public hearing and materials related to the material change transaction, including:
(a) A summary of the proposed transaction;
(b) An explanation of the groups or individuals likely to be impacted by the transaction;
(c) Information about services currently provided by the health care entity, commitments by the health care entity to continue such services and any services that will be reduced or eliminated;
(d) Details about the hearings and how to submit comments, in a format that is easy to find and easy to read; and
(e) Information about potential or perceived conflicts of interest among executives and members of the board of directors of health care entities that are parties to the transaction.

(16) The authority shall post the information described in subsection (15)(a) to (d) of this section to the authority's website in the languages spoken in the area affected by the material change transaction and in a culturally sensitive manner.

(17) The authority shall provide the information described in subsection (15)(a) to (d) of this section to:
(a) At least one newspaper of general circulation in the area affected by the material change transaction;
(b) Health facilities in the area affected by the material change transaction for posting by the health facilities; and
(c) Local officials in the area affected by the material change transaction.

(18) A review board shall make recommendations to the authority to approve the material change transaction, disapprove the material change transaction or approve the material change transaction subject to conditions, based on subsection (9) of this section and the criteria adopted by rule under subsection (2) of this section. The authority shall issue a proposed order and allow the parties and the public a reasonable opportunity to make written
exceptions to the proposed order. The authority shall consider the parties' and the public's written exceptions and issue a final order setting forth the authority's findings and rationale for adopting or modifying the recommendations of the review board. If the authority modifies the recommendations of the review board, the authority shall explain the modifications in the final order and the reasons for the modifications. A party to the material change transaction may contest the final order as provided in ORS chapter 183.

(19) A health care entity that is a party to an approved material change transaction shall notify the authority upon the completion of the transaction in the form and manner prescribed by the authority. One year, two years and five years after the material change transaction is completed, the authority shall analyze:

(a) The health care entities' compliance with conditions placed on the transaction, if any;
(b) The cost trends and cost growth trends of the parties to the transaction; and
(c) The impact of the transaction on the health care cost growth target established under ORS 442.386.

(20) The authority shall publish the authority's analyses and conclusions under subsection (19) of this section and shall incorporate the authority's analyses and conclusions under subsection (19) of this section in the report described in ORS 442.386 (6).

(21) This section does not impair, modify, limit or supersede the applicability of ORS 65.800 to 65.815, 646.605 to 646.652 or 646.705 to 646.805.

(22) Whenever it appears to the Director of the Oregon Health Authority that any person has committed or is about to commit a violation of this section or any rule or order issued by the authority under this section, the director may apply to the Circuit Court for Marion County for an order enjoining the person, and any director, officer, employee or agent of the person, from the violation, and for such other equitable relief as the nature of the case and the interest of the public may require.

(23) The remedies provided under this section are in addition to any other remedy, civil or criminal, that may be available under any other provision of law.

(24) The authority may adopt rules necessary to carry out the provisions of this section.

SECTION 3. (1) An officer or employee of the Oregon Health Authority who is delegated responsibilities in the enforcement of section 2 of this 2021 Act or rules adopted pursuant to section 2 of this 2021 Act may not:

(a) Be a director, officer or employee of or be financially interested in an entity that is a party to a proposed material change transaction except as an enrollee or patient of a health care entity or by reason of rights vested in compensation or benefits related to services performed prior to affiliation with the authority; or
(b) Be engaged in any other business or occupation interfering with or inconsistent with the duties of the authority.

(2) This section does not permit any conduct, affiliation or interest that is otherwise prohibited by public policy.

SECTION 4. (1) The Oregon Health Authority shall prescribe by rule a fee to be paid under section 2 (3) of this 2021 Act, proportionate to the size of the parties to the transaction, sufficient to reimburse the costs of administering section 2 of this 2021 Act.

(2) Moneys received by the authority under this section shall be deposited to the Oregon Health Authority Fund established in ORS 413.101 to be used for carrying out section 2 of this 2021 Act.

SECTION 5. (1) In addition to any other penalty imposed by law, the Director of the Oregon Health Authority may impose a civil penalty, as determined by the director, for a violation of ORS 413.037 or section 2 of this 2021 Act. The amount of the civil penalty may not exceed $10,000 for each offense. The civil penalty imposed on an individual health professional may not exceed $1,000 for each offense.

(2) Civil penalties shall be imposed and enforced in accordance with ORS 183.745.
(3) Moneys received by the Oregon Health Authority under this section shall be paid to the State Treasury and credited to the General Fund.

SECTION 6. Every four years, the Oregon Health Authority shall commission a study of the impact of health care consolidation in this state. The study must review consolidation occurring during the previous four-year period and include an analysis of:

(1) The impact on costs to consumers for health care either to the benefit or the detriment of consumers; and
(2) Any increases or decreases in the quality of care, including:
   (a) Improvement or reductions in morbidity;
   (b) Improvement or reductions in the management of population health;
   (c) Changes to health and patient outcomes, particularly for underserved and uninsured individuals, recipients of medical assistance and other low-income individuals and individuals living in rural areas, as measured by nationally recognized measures of the quality of health care, such as measures used or endorsed by the National Committee for Quality Assurance, the National Quality Forum, the Physician Consortium for Performance Improvement or the Agency for Healthcare Research and Quality.

SECTION 6a. The Oregon Health Authority shall commission the first study under section 6 of this 2021 Act no later than September 15, 2026.

SECTION 7. ORS 413.101 is amended to read:

413.101. The Oregon Health Authority Fund is established in the State Treasury, separate and distinct from the General Fund. Interest earned by the Oregon Health Authority Fund shall be credited to the fund. Moneys in the fund are continuously appropriated to the Oregon Health Authority for carrying out the duties, functions and powers of the authority under ORS 413.032 and 431A.183 and section 2 of this 2021 Act.

SECTION 8. ORS 413.032 is amended to read:

413.032. (1) The Oregon Health Authority is established. The authority shall:
   (a) Carry out policies adopted by the Oregon Health Policy Board;
   (b) Administer the Oregon Integrated and Coordinated Health Care Delivery System established in ORS 414.570;
   (c) Administer the Oregon Prescription Drug Program;
   (d) Develop the policies for and the provision of publicly funded medical care and medical assistance in this state;
   (e) Develop the policies for and the provision of mental health treatment and treatment of addictions;
   (f) Assess, promote and protect the health of the public as specified by state and federal law;
   (g) Provide regular reports to the board with respect to the performance of health services contractors serving recipients of medical assistance, including reports of trends in health services and enrollee satisfaction;
   (h) Guide and support, with the authorization of the board, community-centered health initiatives designed to address critical risk factors, especially those that contribute to chronic disease;
   (i) Be the state Medicaid agency for the administration of funds from Titles XIX and XXI of the Social Security Act and administer medical assistance under ORS chapter 414;
   (j) In consultation with the Director of the Department of Consumer and Business Services, periodically review and recommend standards and methodologies to the Legislative Assembly for:
      (A) Review of administrative expenses of health insurers;
      (B) Approval of rates; and
      (C) Enforcement of rating rules adopted by the Department of Consumer and Business Services;
      (k) Structure reimbursement rates for providers that serve recipients of medical assistance to reward comprehensive management of diseases, quality outcomes and the efficient use of resources and to promote cost-effective procedures, services and programs including, without limitation, preventive health, dental and primary care services, web-based office visits, telephone consultations and telemedicine consultations;
(L) Guide and support community three-share agreements in which an employer, state or local government and an individual all contribute a portion of a premium for a community-centered health initiative or for insurance coverage;

(m) Develop, in consultation with the Department of Consumer and Business Services, one or more products designed to provide more affordable options for the small group market;

(n) Implement policies and programs to expand the skilled, diverse workforce as described in ORS 414.018 (4); and

(o) Implement a process for collecting the health outcome and quality measure data identified by the Health Plan Quality Metrics Committee and report the data to the Oregon Health Policy Board.

(2) The Oregon Health Authority is authorized to:

(a) Create an all-claims, all-payer database to collect health care data and monitor and evaluate health care reform in Oregon and to provide comparative cost and quality information to consumers, providers and purchasers of health care about Oregon’s health care systems and health plan networks in order to provide comparative information to consumers.

(b) Develop uniform contracting standards for the purchase of health care, including the following:

(A) Uniform quality standards and performance measures;

(B) Evidence-based guidelines for major chronic disease management and health care services with unexplained variations in frequency or cost;

(C) Evidence-based effectiveness guidelines for select new technologies and medical equipment;

(D) A statewide drug formulary that may be used by publicly funded health benefit plans; and

(E) Standards that accept and consider tribal-based practices for mental health and substance abuse prevention, counseling and treatment for persons who are Native American or Alaska Native as equivalent to evidence-based practices.

(3) The enumeration of duties, functions and powers in this section is not intended to be exclusive nor to limit the duties, functions and powers imposed on or vested in the Oregon Health Authority by ORS 413.006 to 413.042, 415.012 to 415.430 and 741.340 and section 2 of this 2021 Act or by other statutes.

SECTION 9. ORS 413.037 is amended to read:

413.037. (1) The Director of the Oregon Health Authority, each deputy director and authorized representatives of the director may administer oaths, take depositions and issue subpoenas to compel the attendance of witnesses and the production of documents or other written information necessary to carry out the provisions of ORS 413.006 to 413.042, 415.012 to 415.430 and 741.340 and section 2 of this 2021 Act.

(2) If any person fails to comply with a subpoena issued under this section or refuses to testify on matters on which the person lawfully may be interrogated, the director, deputy director or authorized representative may follow the procedure set out in ORS 183.440 to compel obedience.

SECTION 10. ORS 413.181 is amended to read:

413.181. (1) The Department of Consumer and Business Services and the Oregon Health Authority may enter into agreements governing the disclosure of information reported to the department by insurers with certificates of authority to transact insurance in this state and the disclosure of information reported to the Oregon Health Authority by coordinated care organizations.

(2) The authority may use information disclosed under subsection (1) of this section for the purpose of carrying out ORS 413.032, 414.572, 414.591, 414.605, 414.609, 414.638 and 415.012 to 415.430 and section 2 of this 2021 Act.

SECTION 11. ORS 415.013 is amended to read:

415.013. (1) The Oregon Health Authority shall enforce the provisions of ORS 415.012 to 415.430 and section 2 of this 2021 Act and rules adopted pursuant to ORS 415.011 and 415.012 to 415.430 and section 2 of this 2021 Act for the public good.
(2) The authority has the powers and authority expressly conferred by or reasonably implied from the provisions of ORS 415.012 to 415.430 and section 2 of this 2021 Act and rules adopted pursuant to ORS 415.011 and 415.012 to 415.430 and section 2 of this 2021 Act.

(3) The authority may conduct examinations and investigations of matters concerning the regulation of coordinated care organizations as the authority considers proper to determine whether any person has violated any provision of ORS 415.012 to 415.430 or rules adopted pursuant to ORS 415.011 or to secure information useful in the lawful administration of any of ORS 415.011 the provisions and require the production of books, records, accounts, papers, documents and computer and other recordings the authority considers necessary to administer and enforce ORS 415.012 to 415.430 or section 2 of this 2021 Act and any rules adopted pursuant to ORS 415.011 or 415.012 to 415.430 or section 2 of this 2021 Act.

SECTION 12. ORS 415.019 is amended to read:

415.019. (1) The Oregon Health Authority shall hold a contested case hearing upon written request for a hearing by a person aggrieved by any act, threatened act or failure of the authority to act under ORS 415.012 to 415.430 or section 2 of this 2021 Act or rules adopted pursuant to ORS 415.011 or 415.012 to 415.430 or section 2 of this 2021 Act.

(2) The provisions of ORS chapter 183 govern the hearing procedures and any judicial review of a final order issued in a contested case hearing.

SECTION 13. ORS 415.103 is amended to read:

415.103. A person may not file or cause to be filed with the Oregon Health Authority any article, certificate, report, statement, application or other information required or permitted to be filed under ORS 415.012 to 415.430 or section 2 of this 2021 Act or rules adopted pursuant to ORS 415.011 or 415.012 to 415.430 or section 2 of this 2021 Act that is known by the person to be false or misleading in any material respect.

SECTION 14. Section 2 of this 2021 Act becomes operative on March 1, 2022.