1	H.766
2	Senator Brock moves that the Senate propose to the House that the bill be
3	amended by striking out all after the enacting clause and inserting in lieu
4	thereof the following:
5	Sec. 1. 8 V.S.A. § 4089i(e) is amended to read:
6	(e)(1) A health insurance or other health benefit plan offered by a health
7	insurer or by a pharmacy benefit manager on behalf of a health insurer that
8	provides coverage for prescription drugs and uses step-therapy protocols shall:
9	(A) not require failure, including discontinuation due to lack of
10	efficacy or effectiveness, diminished effect, or an adverse event, on the same
11	medication on more than one occasion for continuously enrolled members or
12	subscribers insureds who are continuously enrolled in a plan offered by the
13	insurer or its pharmacy benefit manager; and
14	(B) grant an exception to its step-therapy protocols upon request of
15	an insured or the insured's treating health care professional under the same
16	time parameters as set forth for prior authorization requests in 18 V.S.A.
17	§ 9418b(g)(4) if any one or more of the following conditions apply:
18	(i) the prescription drug required under the step-therapy protocol
19	is contraindicated or will likely cause an adverse reaction or physical or mental
20	harm to the insured;

1	(ii) the prescription drug required under the step-therapy protocol
2	is expected to be ineffective based on the insured's known clinical history,
3	condition, and prescription drug regimen;
4	(iii) the insured has already tried the prescription drugs on the
5	protocol, or other prescription drugs in the same pharmacologic class or with
6	the same mechanism of action, which have been discontinued due to lack of
7	efficacy or effectiveness, diminished effect, or an adverse event, regardless of
8	whether the insured was covered at the time on a plan offered by the current
9	insurer or its pharmacy benefit manager;
10	(iv) the insured is stable on a prescription drug selected by the
11	insured's treating health care professional for the medical condition under
12	consideration; or
13	(v) the step-therapy protocol or a prescription drug required under
14	the protocol is not in the patient's best interests because it will:
15	(I) pose a barrier to adherence;
16	(II) likely worsen a comorbid condition; or
17	(III) likely decrease the insured's ability to achieve or maintain
18	reasonable functional ability.
19	(2) Nothing in this subsection shall be construed to prohibit the use of
20	tiered co-payments for members or subscribers not subject to a step-therapy
21	protocol.

1	(3) Notwithstanding <u>any provision of</u> subdivision (1) of this subsection
2	to the contrary, a health insurance or other health benefit plan offered by an
3	insurer or by a pharmacy benefit manager on behalf of a health insurer that
4	provides coverage for prescription drugs shall not utilize a step-therapy, "fail
5	first," or other protocol that requires documented trials of a medication,
6	including a trial documented through a "MedWatch" (FDA Form 3500), before
7	approving a prescription for the treatment of substance use disorder.
8	Sec. 2. 18 V.S.A. § 9418a is amended to read:
9	§ 9418a. PROCESSING CLAIMS, DOWNCODING, AND ADHERENCE
10	TO CODING RULES
11	(a) Health plans, contracting entities, covered entities, and payers shall
12	accept and initiate the processing of all health care claims submitted by a
13	health care provider pursuant to and consistent with the current version of the
14	American Medical Association's Current Procedural Terminology (CPT)
15	codes, reporting guidelines, and conventions; the Centers for Medicare and
16	Medicaid Services Healthcare Common Procedure Coding System (HCPCS);
17	American Society of Anesthesiologists; the National Correct Coding Initiative
18	(NCCI); the National Council for Prescription Drug Programs coding; or other
19	appropriate nationally recognized standards, guidelines, or conventions
20	approved by the Commissioner.

1	(b)(1) When Except as provided in subsection (c) of this section, when
2	editing claims, health plans, contracting entities, covered entities, and payers
3	shall adhere to require not more than the following edit standards, processes,
4	and guidelines except as provided in subsection (c) of this section:
5	(1)(A) the CPT, HCPCS, and for claims for outpatient and professional
6	services, the NCCI as in effect for Medicare;
7	(2)(B) national specialty society edit standards for facility claims, the
8	Medicare Code Editor as in effect for Medicare; or
9	(3)(C) for pharmacy claims, appropriate nationally recognized edit
10	standards, guidelines, or conventions; and
11	(D) for any other claim not addressed by subdivision (A), (B), or (C)
12	of this subdivision (1), other appropriate nationally recognized edit standards,
13	guidelines, or conventions approved by the Commissioner.
14	(2) For outpatient services, professional services, and facility claims, a
15	health plan, contracting entity, covered entity, or payer shall apply the relevant
16	edit standards, processes, and guidelines from NCCI or Medicare Code Editor
17	pursuant to subdivisions (1)(A) and (B) of this subsection that were in effect
18	for Medicare on the date of the claim submission; provided, however, that if
19	Medicare has changed an applicable edit standard, process, or guideline within
20	90 days prior to the date of the claim submission, the health plan, contracting
21	entity, covered entity, or payer may use the version of the edit standard,

1	process, or guideline that Medicare had applied prior to the most recent change
2	if the health plan, contracting entity, covered entity, or payer has not yet
3	released an updated version of its edits in accordance with subsection (d) of
4	this section.
5	(c) Adherence to the edit standards in subdivision (b)(1) or (2) subsection
6	(b) of this section is not required:
7	(1) when necessary to comply with State or federal laws, rules,
8	regulations, or coverage mandates; or
9	(2) for edits that the payer determines are more favorable to providers
10	than the edit standards in subdivisions (b)(1) through (3) subsection (b) of this
11	section or to address new codes not yet incorporated by a payer's edit
12	management software, provided the edit standards are:
13	(A) developed with input from the relevant Vermont provider
14	community and national provider organizations;
15	(B) clearly supported by nationally recognized standards, guidelines,
16	or conventions approved by the Commissioner of Financial Regulation; and
17	(C) provided the edits are available to providers on the plan's
18	websites and in their its newsletters or equivalent electronic communications.
19	(d) Health plans, contracting entities, covered entities, and payers shall not
20	release edits more than quarterly, to take effect on January 1, April 1, July 1, or
21	October 1, as applicable, and the edits shall not be implemented without filing

1	with the Commissioner of Financial Regulation to ensure consistency with
2	nationally recognized standards guidelines, and conventions, and at least 30
3	days' advance notice to providers. Whenever Medicare changes an edit
4	standard, process, or guideline that it applies to outpatient service, professional
5	service, or facility claims, each health plan, contracting entity, covered entity,
6	or payer shall incorporate those modifications into its next quarterly release of
7	edits.
8	(e)(1) Except as otherwise provided in subdivision (2) of this subsection,
9	no health plan, contracting entity, covered entity, or payer shall subject any
10	health care provider to prepayment coding validation edit review. As used in
11	this subsection, "prepayment coding validation edit review" means any action
12	by the health plan, contracting entity, covered entity, or payer, or by a
13	contractor, assignee, agent, or other entity acting on its behalf, requiring a
14	health care provider to provide medical record documentation in conjunction
15	with or after submission of a claim for payment for health care services
16	delivered, but before the claim has been adjudicated.
17	(2) Nothing in this subsection shall be construed to prohibit targeted
18	prepayment coding validation edit review of a specific provider, provider
19	group, or facility under certain circumstances, including evaluating high-dollar
20	claims; verifying complex financial arrangements; investigating member
21	questions; conducting post-audit monitoring; addressing a reasonable belief of

1	fraud, waste, or abuse; or other circumstances determined by the
2	Commissioner through a bulletin or guidance.
3	(f) Nothing in this section shall preclude a health plan, contracting entity,
4	covered entity, or payer from determining that any such claim is not eligible
5	for payment in full or in part, based on a determination that:
6	* * *
7	(e)(g) Nothing in this section shall be deemed to require a health plan,
8	contracting entity, covered entity, or payer to pay or reimburse a claim, in full
9	or in part, or to dictate the amount of a claim to be paid by a health plan,
10	contracting entity, covered entity, or payer to a health care provider.
11	(f)(h) No health plan, contracting entity, covered entity, or payer shall
12	automatically reassign or reduce the code level of evaluation and management
13	codes billed for covered services (downcoding), except that a health plan,
14	contracting entity, covered entity, or payer may reassign a new patient visit
15	code to an established patient visit code based solely on CPT codes, CPT
16	guidelines, and CPT conventions.
17	$\frac{(g)(i)}{g}$ Notwithstanding the provisions of subsection $\frac{(d)(f)}{g}$ of this section,
18	and other than the edits contained in the conventions in subsections (a) and (b)
19	of this section, health plans, contracting entities, covered entities, and payers
20	shall continue to have the right to deny, pend, or adjust claims for services on
21	other bases and shall have the right to reassign or reduce the code level for

21

1	selected claims for services based on a review of the clinical information
2	provided at the time the service was rendered for the particular claim or a
3	review of the information derived from a health plan's fraud or abuse billing
4	detection programs that create a reasonable belief of fraudulent or abusive
5	billing practices, provided that the decision to reassign or reduce is based
6	primarily on a review of clinical information.
7	(h)(j) Every If adding an edit pursuant to subsection (b) or subdivision
8	(c)(1) or (2) of this section, a health plan, contracting entity, covered entity,
9	and or payer shall publish on its provider website and in its provider newsletter
10	if applicable or equivalent electronic provider communications:
11	(1) the name of any commercially available claims editing software
12	product that the health plan, contracting entity, covered entity, or payer
13	utilizes;
14	(2) the <u>specific</u> standard or standards , pursuant to subsection (b) of this
15	section, that the entity uses for claim edits and how those claim edits are
16	supported by those specific standards;
17	(3) the payment percentages for modifiers; and
18	(4) any significant the specific edit or edits, as determined by the health
19	plan, contracting entity, covered entity, or payer, added to the claims software
20	product after the effective date of this section, which are made at the request of

the health plan, contracting entity, covered entity, or payer.

1	$\frac{(1)(K)}{(1)}$ Upon written request, the health plan, contracting entity, covered
2	entity, or payer shall also directly provide the information in subsection (h)(j)
3	of this section to a health care provider who is a participating member in the
4	health plan's, contracting entity's, covered entity's, or payer's provider
5	network.
6	(j)(1) For purposes of this section, "health plan" includes a workers'
7	compensation policy of a casualty insurer licensed to do business in Vermont.
8	(k)(m) BlueCross BlueShield of Vermont and the Vermont Medical
9	Society are requested to continue convening a work group consisting of There
10	is established a working group comprising the health plans, contracting
11	entities, covered entities, and payers subject to the reporting requirement in
12	subsection 9414a(b) of this title; representatives of hospitals and health care
13	providers; representatives of the Department of Financial Regulation and of
14	other relevant State agencies; and other interested parties to study the edit
15	standards in subsection (b) of this section, the edit standards in national class
16	action settlements, and edit standards and edit transparency standards
17	established by other states to determine the most appropriate way to ensure that
18	health care providers can access information about the edit standards
19	applicable to the health care services they provide trends in coding and billing
20	that health plans, contracting entities, covered entities, or payers, or a
21	combination of them, seek to address through claim editing. The work

1	working group is requested to shall provide an annual a progress report to the
2	House Committee on Health Care and the Senate Committees on Health and
3	Welfare and on Finance upon request.
4	(1)(n) With respect to the work working group established under subsection
5	(k)(m) of this section and to the extent required to avoid violations of federal
6	antitrust laws, the Department shall facilitate and supervise the participation of
7	members of the work working group.
8	Sec. 3. 18 V.S.A. § 9418b(c) and (d) are amended to read:
9	(c) A health plan shall furnish, upon request from a health care provider, a
10	current list of services and supplies requiring prior authorization.
11	(1) It is the intent of the General Assembly to reduce variability in prior
12	authorization requirements by aligning to the greatest extent possible with the
13	prior authorization requirements in Vermont's Medicaid program.
14	(2) A health plan shall not impose any prior authorization requirement
15	for any admission, item, service, treatment, or procedure that is more
16	restrictive than the prior authorization requirements that the Department of
17	Vermont Health Access would apply for the same admission, item, service,
18	treatment, or procedure under Vermont's Medicaid program.
19	(3) Each health plan shall review the prior authorization requirements in
20	effect in Vermont's Medicaid program at least once every six months to ensure

1	that the health plan is maintaining the prior authorization alignment required
2	by subdivision (2) of this subsection.
3	(4) Nothing in this subsection shall be construed to:
4	(A) require prior authorization alignment with Vermont Medicaid for
5	prescription drugs;
6	(B) prohibit prior authorization requirements for any admission, item,
7	service, treatment, or procedure that is not covered by Vermont Medicaid;
8	(C) prohibit prior authorization requirements for an admission, item,
9	service, treatment, or procedure that is provided out-of-network; or
10	(D) require a health plan to maintain the same provider network as
11	Vermont Medicaid.
12	(d)(1) A health plan shall furnish, upon request from a health care provider,
13	a current list of services and supplies requiring prior authorization.
14	(2) A health plan shall post make a current list of services and supplies
15	requiring prior authorization <u>available</u> to <u>the public on</u> the insurer's website.
16	Sec. 4. 18 V.S.A. § 9418b(g)(4) is amended to read:
17	(4) A health plan shall respond to a completed prior authorization
18	request from a prescribing health care provider within 48 hours after receipt for
19	urgent requests and within two business days after receipt for nonurgent
20	requests. The health plan shall notify a health care provider of or make
21	available to a health care provider a receipt of the request for prior

1	authorization and any needed missing information within 24 hours after
2	receipt.
3	(A)(i) For urgent prior authorization requests, a A health plan shall
4	approve, deny, or inform the insured or health care provider if any information
5	is missing from a prior authorization request from an insured or a prescribing
6	health care provider within 24 hours following receipt for all prior
7	authorization requests and for all appeals of denial of prior authorization
8	<u>requests.</u>
9	(ii) If a health plan informs an insured or a health care provider
10	that more information is necessary for the health plan to make a determination
11	on the request, the health plan shall have 24 hours to approve or deny the
12	request upon receipt of the necessary information.
13	(B) For nonurgent prior authorization requests:
14	(i) A health plan shall approve or deny a completed prior
15	authorization request from an insured or a prescribing health care provider
16	within two business days following receipt.
17	(ii) A health plan shall acknowledge receipt of the prior
18	authorization request within 24 hours following receipt and shall inform the
19	insured or health care provider at that time if any information is missing that is
20	necessary for the health plan to make a determination on the request.

1	(iii) If a health plan notifies an insured or a health care provider
2	that more information is necessary pursuant to subdivision (ii) of this
3	subdivision (4)(B), the health plan shall have 24 hours to approve or deny the
4	request upon receipt of the necessary information.
5	(B) If a health plan does not, within the time limits set forth in this
6	section subdivision (A) of this subdivision (4), respond to a completed prior
7	authorization request, acknowledge receipt of the request for prior
8	authorization, or request missing information, the prior authorization request
9	shall be deemed to have been granted.
10	(C) Prior authorization approval for a prescribed or ordered
11	treatment, service, or course of medication shall be valid for the duration of the
12	prescribed or ordered treatment, service, or course of medication or one year,
13	whichever is longer; provided, however, that for a prescribed or ordered
14	treatment, service, or course of medication that continues for more than one
15	year, a health plan shall not require renewal of the prior authorization approval
16	more frequently than once every five years.
17	(D) For an insured who is stable on a treatment, service, or course of
18	medication, as determined by a health care provider, that was approved for
19	coverage under a previous health plan, a health plan shall not restrict coverage
20	of that treatment, service, or course of medication for at least 90 days upon the
21	insured's enrollment in the new health plan.

1	Sec. 5. 18 V.S.A. § 9418c is amended to read:
2	§ 9418c. FAIR CONTRACT STANDARDS
3	(a) Required information.
4	(1) Each contracting entity shall provide and each health care contract
5	shall obligate the contracting entity to provide participating health care
6	providers information sufficient for the participating provider to determine the
7	compensation or payment terms for health care services, including all of the
8	following:
9	(A) The manner of payment, such as fee-for-service, capitation, case
10	rate, or risk.
11	(B) On request, the fee-for-service dollar amount allowable for each
12	CPT code for those CPT codes that a provider in the same specialty typically
13	uses or that the requesting provider actually bills. Fee schedule information
14	may be provided by CD-ROM or electronically, at the election of the
15	contracting entity, but a provider may elect to receive a hard copy of the fee
16	schedule information instead of the CD-ROM or electronic version.
17	(C) A clearly understandable, readily available mechanism, such as a
18	specific website address, that includes the following information:
19	(i) the name of the commercially available claims editing software
20	product that the health plan, contracting entity, covered entity, or payer uses;

1	(ii) the <u>specific</u> standard or standards from subsection 9418a(c) of
2	this title that the entity uses for claim edits and how those claim edits are
3	supported by those specific standards;
4	(iii) payment percentages for modifiers; and
5	(iv) any significant edits, as determined by the health plan,
6	contracting entity, covered entity, or payer, added to the claims software
7	product, which are made at the request of the health plan, contracting entity,
8	covered entity, or payer, and which have been approved by the Commissioner
9	pursuant to subsection 9418a(b) or (c) of this title.
10	(D) Any policies for prepayment or postpayment audits, or both,
11	including whether the policies include limits on the number of medical records
12	a contracting entity may request for audit in any calendar year.
13	* * *
14	(5)(A) If a contracting entity uses policies or manuals to augment the
15	content of the contract with a health care provider, the contracting entity shall
16	ensure that those policies or manuals contain sufficient information to allow
17	providers to understand and comply with the content.
18	(B) For any new policy or manual, or any change to an existing
19	policy or manual, the contracting entity shall do all of the following:
20	(i) Provide notice of the new policy, manual, or change to each
21	participating provider in writing not fewer than 60 days prior to the effective

1	date of the policy, manual, or change, which notice shall be conspicuously
2	entitled "Notice of Policy Change" and shall include:
3	(I) a summary of the new policy, manual, or change;
4	(II) an explanation of the policy, manual, or change;
5	(III) the effective date of the policy, manual, or change; and
6	(IV) a notice of the right to object in writing to the policy,
7	manual, or change, along with a timeframe for objection and where and how to
8	send the objection.
9	(ii) Provide the participating provider 60 days after receiving the
10	notice and summary to object in writing to the new policy, manual, or change.
11	If the participating provider objects to the new policy, manual, or change, the
12	contracting entity shall provide an initial substantive response to the objection
13	within 30 days following the contracting entity's receipt of the written
14	objection, and the contracting entity shall work together with the provider to
15	achieve a reasonable resolution to the objection within 60 days following the
16	provider's receipt of contracting entity's initial substantive response. If the
17	provider is not satisfied with the proposed resolution, the provider may pursue
18	any remedy available to the provider under the health care contract or under
19	applicable law.
20	* * *

1	Sec. 6. DEPARTMENT OF FINANCIAL REGULATION; ACTUARIAL
2	ANALYSIS; REPORT
3	(a) The Department of Financial Regulation shall arrange for an
4	actuarial analysis of the anticipated costs and potential benefits of aligning
5	health insurance plans' claim edit requirements with those in Medicare, as
6	set forth in Sec. 2 of this act, and of aligning health insurance plans' prior
7	authorization requirements with those in Vermont Medicaid, as set forth
8	in Sec. 3 of this act.
9	(b) On or before December 1, 2024, the Department of Financial
10	Regulation shall provide the results of the actuarial analysis to the House
11	Committee on Health Care and the Senate Committees on Health and
12	Welfare and on Finance.
13	Sec. 7. PRIOR AUTHORIZATION; INSURER IMPACT REPORTS
14	On or before January 15, 2027, each health insurer with at least 2,000
15	covered lives in Vermont shall report to the House Committee on Health Care
16	and the Senate Committees on Health and Welfare and on Finance regarding
17	the impact of the prior authorization provisions of this act on the following
18	during plan years 2025 and 2026:
19	(1) utilization of health care services covered by the insurer's plans;
20	(2) development of the insurer's premium rates for future plan years;
21	<u>and</u>

1	(3) the insurer's estimated avoided costs, including:
2	(A) the specific methodologies that the insurer uses to determine the
3	amount of "savings" from avoided costs;
4	(B) the costs of the alternative tests, procedures, medications, and
5	other items or services ordered for insureds as a result of the insurer's denials
6	of requests for prior authorizations; and
7	(C) the costs of emergency department visits and inpatient stays,
8	including stays in intensive care units, as a result of the insurer's denials of
9	requests for prior authorizations.
10	Sec. 8. PRIOR AUTHORIZATION; PROVIDER IMPACT REPORTS
11	(a) The General Assembly requests that organizations representing
12	Vermont's hospital-employed, federally qualified health center-employed, and
13	independent health care providers who are affected by the prior authorization
14	provisions of this act gather information from their members on or before
15	January 1, 2025 and on or before July 1, 2026 regarding current circumstances
16	and the impact of the prior authorization provisions of this act on their provider
17	members and the members' practices. To the extent practicable, the
18	information gathered should align with survey questions published by
19	nationally recognized provider organizations and include information
20	regarding the impact of prior authorization processes and requirements on care
21	delivery, quality of care, and staffing.

1	(b) On or before January 15, 2027, each provider organization that gathered
2	information from its members in accordance with subsection (a) of this section
3	is requested to summarize and report on that information to the House
4	Committee on Health Care and the Senate Committees on Health and Welfare
5	and on Finance, including providing a summary of the impact of the prior
6	authorization provisions of this act on the organization's members' practices.
7	Sec. 9. REPEAL
8	18 V.S.A. § 9418(m) and (n) (claims edit working group) are repealed on
9	<u>January 1, 2028.</u>
10	Sec. 10. EFFECTIVE DATES
11	(a) Secs. 6 (Department of Financial Regulation; actuarial analysis;
11 12	(a) Secs. 6 (Department of Financial Regulation; actuarial analysis; report), 7 (prior authorization; insurer impact reports), and 8 (prior
12	report), 7 (prior authorization; insurer impact reports), and 8 (prior
12 13	report), 7 (prior authorization; insurer impact reports), and 8 (prior authorization; provider impact reports) and this section shall take effect on
12 13 14	report), 7 (prior authorization; insurer impact reports), and 8 (prior authorization; provider impact reports) and this section shall take effect on passage.
12 13 14 15	report), 7 (prior authorization; insurer impact reports), and 8 (prior authorization; provider impact reports) and this section shall take effect on passage. (b) Sec. 4 (18 V.S.A. § 9418b(g)(4); prior authorization time frames) shall
12 13 14 15 16	report), 7 (prior authorization; insurer impact reports), and 8 (prior authorization; provider impact reports) and this section shall take effect on passage. (b) Sec. 4 (18 V.S.A. § 9418b(g)(4); prior authorization time frames) shall take effect on January 1, 2025, except that a health plan that must modify its
12 13 14 15 16 17	report), 7 (prior authorization; insurer impact reports), and 8 (prior authorization; provider impact reports) and this section shall take effect on passage. (b) Sec. 4 (18 V.S.A. § 9418b(g)(4); prior authorization time frames) shall take effect on January 1, 2025, except that a health plan that must modify its technology in order to continue administering its own internal utilization

1	(c) In Sec. 2 (18 V.S.A. § 9418a), subsections (b)–(d) shall take effect on
2	January 1, 2026 and shall apply to all health plans issued on and after that
3	date, to all health care provider contracts entered into or renewed on and
4	after that date, and to all claims processed on and after that date.
5	(d) Sec. 3 (18 V.S.A. § 9418b(c) and (d)) shall take effect on January 1,
6	2026 and shall apply to all health plans issued on and after that date, to all
7	health care provider contracts entered into or renewed on and after that
8	date, and to all claims processed on and after that date.
9	(e) The remaining sections shall take effect on January 1, 2025 and shall
10	apply to all health plans issued on and after that date, to all health care provider
11	contracts entered into or renewed on and after that date, and to all claims
12	processed on and after that date.