# 2022 South Dakota Legislature

# Senate Bill 163

## AMENDMENT 163A FOR THE INTRODUCED BILL

## 1 An Act to address transparency in prescription drug pricing.

## 2 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF SOUTH DAKOTA:

#### 3 Section 1. That § 58-29E-1 be AMENDED:

#### 58-29E-1. Terms used in this chapter mean:

- 5 "Covered entity," a nonprofit hospital or medical service corporation, health insurer, (1)6 health benefit plan, or health maintenance organization; a health program 7 administered by a department or the state in the capacity of provider of health 8 coverage; or an employer, labor union, or other group of persons organized in the 9 state that provides health coverage to covered individuals who are employed or 10 reside in the state. The term does not include a self-funded plan that is exempt 11 from state regulation pursuant to ERISA, a plan issued for coverage for federal 12 employees, or a health plan that provides coverage only for accidental injury, 13 specified disease, hospital indemnity, medicare supplement, disability income, 14 long-term care, or other limited benefit health insurance policies and contracts;
- (2) "Covered individual," a member, participant, enrollee, contract holder, policy
   holder, or beneficiary of a covered entity third-party payor who is provided health
   coverage by the covered entity third-party payor. The term includes a dependent
   or other person provided health coverage through a policy, contract, or plan for a
   covered individual;
- 20 (3)(2) "Director," the director of the Division of Insurance;
- (4)(3) "Generic drug," a chemically equivalent copy of a brand-name drug with an expired
   patent;
- (5)(4) "Labeler," an entity or person that receives prescription drugs from a manufacturer
   or wholesaler and repackages those drugs for later retail sale and that has a labeler
   code from the federal Food and Drug Administration under 21 C.F.R. § 270.20
   (1999);

1	(6)(5) "Maximum allowable cost list," any listing of pharmaceutical products, or method
2	for calculating reimbursement amounts, used by a pharmacy benefit manager,
3	directly or indirectly, to establish the maximum allowable cost on which
4	reimbursement payment, to a pharmacy or pharmacist, may be based for
5	dispensing a prescription pharmaceutical product, including:
6	(a) Average acquisition cost;
7	(b) Average manufacturer price;
8	(c) Average wholesale price;
9	(d) Brand effective rate or generic effective rate;
10	(e) Discount indexing;
11	(f) Federal upper limits;
12	(g) National average drug acquisition cost;
13	(h) Wholesale acquisition cost; and
14	(i) Any other factor used by a pharmacy benefit manager or a third-party payor
15	to establish reimbursement rates to a pharmacy or pharmacist for
16	pharmaceutical products;
17	(6) "National Drug Code," a unique, three-segment numeric identifier assigned to each
18	medication in accordance with the Federal Food, Drug, and Cosmetic Act, 21 U.S.C.
19	<u>§ 360 (as of January 1, 2022);</u>
20	(7) "Pharmaceutical product," a generic drug, brand-name drug, biologic, or other
21	prescription drug, vaccine, or device;
22	(8) "Pharmaceutical wholesaler," a person who:
23	(a) Sells and distributes, directly or indirectly, pharmaceutical products and
24	over-the-counter pharmaceuticals; and
25	(b) Offers regular or private delivery to a pharmacy;
26	(9) "Pharmacy acquisition cost," the amount that a pharmaceutical wholesaler charges
27	for a pharmaceutical product, as listed on the pharmacy's billing invoice;
28	(10) "Pharmacy benefits benefit management," the procurement of prescription drugs at
29	a negotiated rate for dispensation within this state to covered individuals, the
30	administration or management of prescription drug benefits provided by a <del>covered</del>
31	<del>entity <u>third-party payor</u>for the benefit of <mark>covered</mark>individuals, or any of the</del>
32	following services provided with regard to the administration of the following
33	pharmacy benefits:
34	(a) Mail service pharmacy;

(b) Claims processing, retail network management, and payment of claims to
pharmacies for prescription drugs dispensed to covered individuals;
(c) Clinical formulary development and management services;
(d) Rebate contracting and administration;
(e) Certain patient compliance, therapeutic intervention, and generic substitution
programs; and
(f) Disease management programs involving prescription drug utilization;
<del>(7)<u>(11)</u> "Pharmacy <u>benefits</u> <u>benefit</u> manager," <del>an entity that</del>a person who performs</del>
pharmacy <del>benefits <u>benefit</u> management<mark>. The term includesa <u>:</u></mark></del>
(a) <u>A person or entity acting for a pharmacy benefits benefit</u> manager, in a
contractual or employment relationship, in the performance of pharmacy
benefits benefit management for a covered entity andincludes
mail <u>third-party payor; and</u>
(b) <u>A mail-</u> service pharmacy.
The term does not include a health carrier licensed pursuant to Title 58when , if
the health carrier or its subsidiary is providing pharmacy benefits management to
its own insureds; or a public self-funded pool or a private single employer
self-funded plan that provides such benefits or services directly to its beneficiaries
<del>for a third-party payor</del> ;
(8)(12) "Pharmacy benefit manager affiliate," a pharmacy that or a pharmacist who,
directly or indirectly, through one or more intermediaries:
(a) Owns or controls a pharmacy benefit manager;
(b) Is owned or controlled by a pharmacy benefit manager; or
(c) Is under common ownership or control with a pharmacy benefit manager;
(13) "Pharmacy benefit plan or program," a plan or program that pays for, reimburses,
covers the cost of, or otherwise provides for pharmaceutical products to individuals
who reside in, or are employed in, this state;
(14) "Pharmacy service administrative organization," an organization that has the
authority to contract with a pharmacy benefit manager on behalf of multiple
independently owned pharmacies;
(15) "Proprietary information," information on pricing, costs, revenue, taxes, market
share, negotiating strategies, customers, and personnel held by private entities
and used for that private entity's business purposes;
<del>(9)</del> (16)"Third-party payor," any person involved in the financing of a pharmacy benefit
plan or program, other than:

1		<u>(a)</u>	The patient;
2		<u>(b)</u>	A health care provider; or
3		<u>(c)</u>	The sponsor of a plan that is subject to regulation under Medicare Part D,
4			42 U.S.C. § 1395w-101, et seq., as of January 1, 2022; or
5		<u>(d)</u>	A plan administered by South Dakota Medicaid;
6	<u>(17)</u>	"Trade	secret," information, including a formula, pattern, compilation, program,
7		device	, method, technique, or process, that:
8		(a)	Derives independent economic value, actual or potential, from not being
9			generally known to, and not being readily ascertainable by proper means
10			by, other persons who can obtain economic value from its disclosure or use;
11			and
12		(b)	Is the subject of efforts that are reasonable under the circumstances to
13			maintain its secrecy <u>; and</u>
14	<u>(18)</u>	"340B	entity," an entity participating in the federal drug discount program, as
15		<u>describ</u>	bed in section 340B of the Public Health Service Act, 42 U.S.C. § 256b, as of
16		<u>Januar</u>	<u>y 1, 2022</u> .
4 7		<b>7</b>	
17	Section	2. Inat	§ 58-29E-3 be AMENDED:
18		58-29	E-3. Each pharmacy benefits benefit manager shall perform its duties
18 19	exerc		<b>E-3.</b> Each pharmacy benefits <u>benefit</u> manager shall perform its duties of faith and fair dealing toward the covered entitythird-party payor.
19		ising goo	od faith and fair dealing toward the covered entitythird-party payor.
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19 20		ising goo <b>3. That</b>	od faith and fair dealing toward the <del>covered entity<u>third-party payor</u>.</del>
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19 20	Section : placer	ising goo <b>3. That</b> <u>Before</u> ment of	od faith and fair dealing toward the covered entitythird-party payor. chapter 58-29E be amended with a NEW SECTION: a pharmaceutical benefit manager places or provides for the continued f a pharmaceutical product on a maximum allowable cost list, the
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19 20 21 22 23 24 25 26 27 28 29	Section : placer pharm	ising goo <b>3. That</b> <u>Before</u> <u>ment of</u> <u>naceutica</u> <u>The pro-</u> <u>(a)</u>	od faith and fair dealing toward the <del>covered entity</del> <u>third-party payor</u> . <b>chapter 58-29E be amended with a NEW SECTION:</b> a pharmaceutical benefit manager places or provides for the continued f a pharmaceutical product on a maximum allowable cost list, the al benefit manager shall ensure that: oduct: Is listed as therapeutically equivalent and pharmaceutically equivalent A- or B-rated in the United States Food and Drug Administration's most recent edition of Approved Drug Products with Therapeutic Equivalence Evaluations or on the United States Food and Drug Administration's most recent list of approved animal drug products; or
19 20 21 22 23 24 25 26 27 28 29 30	Section : placer pharm	ising goo <b>3. That</b> <u>Before</u> <u>ment of</u> <u>naceutica</u> <u>The product</u> <u>(a)</u>	od faith and fair dealing toward the <del>covered entity</del> <u>third-party payor</u> . <b>chapter 58-29E be amended with a NEW SECTION:</b> <u>a pharmaceutical benefit manager places or provides for the continued</u> <u>a pharmaceutical product on a maximum allowable cost list, the</u> <u>al benefit manager shall ensure that:</u> <u>oduct:</u> <u>Is listed as therapeutically equivalent and pharmaceutically equivalent A- or B-rated in the United States Food and Drug Administration's most recent <u>edition of Approved Drug Products with Therapeutic Equivalence Evaluations</u> <u>or on the United States Food and Drug Administration's most recent list of</u> <u>approved animal drug products; or</u> <u>Has an NR rating, an NA rating, or a similar rating by a nationally recognized</u></u>
19 20 21 22 23 24 25 26 27 28 29 30 31	Section : placer pharm (1)	ising goo <b>3. That</b> <u>Before</u> <u>ment of</u> <u>naceutica</u> <u>The pro-</u> (a) <u>(b)</u>	od faith and fair dealing toward the <del>covered entitythird-party payor</del> . <b>chapter 58-29E be amended with a NEW SECTION:</b> <u>a pharmaceutical benefit manager places or provides for the continued</u> <u>f a pharmaceutical product on a maximum allowable cost list, the</u> <u>al benefit manager shall ensure that:</u> <u>oduct:</u> <u>Is listed as therapeutically equivalent and pharmaceutically equivalent A-</u> <u>or B-rated in the United States Food and Drug Administration's most recent</u> <u>edition of Approved Drug Products with Therapeutic Equivalence Evaluations</u> <u>or on the United States Food and Drug Administration's most recent list of</u> <u>approved animal drug products; or</u> <u>Has an NR rating, an NA rating, or a similar rating by a nationally recognized</u> <u>drug compendia provider;</u>

1	<u>(3)</u>	The product is not obsolete.			
2		For purposes of this section, the term, NR, means not rated, and the term, NA,			
3	mean	ans not available.			
4	Section 4	4. That chapter 58-29E be amended with a NEW SECTION:			
5		<u>A pharmacy benefit manager shall:</u>			
6	<u>(1)</u>	Provide each pharmacy that is subject to the maximum allowable cost list with			
7		notification of any changes to the list;			
8	<u>(2)</u>	Provide each pharmacy that is subject to the maximum allowable cost list with			
9		access to the list; and			
10	<u>(3)</u>	Update the maximum allowable cost list within seven calendar days if:			
11		(a) Pharmacy acquisition costs from at least sixty percent of the pharmaceutical			
12		wholesalers doing business in the state increase by ten percent or more			
13		over the previously listed cost;			
14		(b) There is a change in the methodology on which the maximum allowable cost			
15		<u>list is based; or</u>			
16		(c) There is a change in the value of a variable involved in the methodology.			
17	Section !	5. That chapter 58-29E be amended with a NEW SECTION:			
18		A pharmacy benefit manager shall establish an administrative procedure by which			
19	<u>a pha</u>	armacy may appeal determinations regarding the maximum allowable costs and			
20	reimb	ursements for a specific pharmaceutical product as:			
21	<u>(1)</u>	Not meeting the requirements set forth in this chapter; or			
22	<u>(2)</u>	Being below the pharmacy acquisition cost.			
23	Section	6. That chapter 58-29E be amended with a NEW SECTION:			
24		The administrative procedure required under section 5 of this Act must:			
25	<u>(1)</u>	Provide a telephone number, email address, and website, for initiating an appeal;			
26	<u>(2)</u>	Provide that an appeal may be filed directly with the pharmacy benefit manger or			
27		through a pharmacy service administrative organization; and			
28	<u>(3)</u>	Establish a period within which any appeal is to be filed, provided the period is at			
29		least seven days.			

# **Section 7. That chapter 58-29E be amended with a NEW SECTION:**

1	If an appeal is filed in accordance with the administrative procedure set forth in			
2	section 5 of this Act, the pharmacy benefit manager shall, within seven days of receipt:			
3	(1) Find that the appeal is merited and:			
4	(a) Make the change in the maximum allowable cost;			
5	(b) Permit the appealing pharmacy or pharmacist to reverse and re-bill the			
6	claim in question;			
7	(c) Provide to the pharmacy or pharmacist the National Drug Code on which			
8	the change is based; and			
9	(d) Ensure that the change made under this subsection is effective for each			
10	similarly situated pharmacy, as defined by the payor, subject to the			
11	maximum allowable cost list; or			
12	(2) Find that the appeal is not merited and provide to the appealing pharmacy or			
13	pharmacist the National Drug Code and the name of the national or regional			
14	pharmaceutical wholesalers who are operating in this state and have the drug in			
15	stock at a price below that on the maximum allowable cost list.			
16	If the National Drug Code provided by the pharmacy benefit manager is not			
17	available below the pharmacy acquisition cost of the pharmaceutical wholesaler from			
18	whom the pharmacy or pharmacist purchases the majority of prescription drugs for resale,			
19	the pharmacy benefit manager shall adjust the maximum allowable cost, as listed on the			
20	maximum allowable cost list, above the appealing pharmacy's acquisition cost and permit			
21	the appealing pharmacy to reverse and re-bill each claim affected by the inability to			
22	procure the drug at a cost that is equal to or less than the previously appealed maximum			
23	allowable cost.			
24	Section 8. That chapter 58-29E be amended with a NEW SECTION:			
25	A pharmacy benefit manager may not reimburse a pharmacy or pharmacist in the			
26	state an amount less than the amount that the pharmacy benefit manager reimburses a			
27	pharmacy benefit manager affiliate for providing the same pharmacist services.			
28	The amount must be calculated on a per-unit basis, using the same generic product			
29	identifier or generic code number.			
30	Section 9. That chapter 58-29E be amended with a NEW SECTION:			
31	A pharmacy or pharmacist may decline to provide a pharmaceutical product to a			
32	patient or pharmacy benefit manager if, as a result of a maximum allowable cost list, a			

# pharmacy or pharmacist is to be paid less than the pharmacy acquisition cost of the pharmacy providing the pharmaceutical product.

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#### 4 Section 10. That chapter 58-29E be amended with a NEW SECTION:

- 5 <u>A pharmacy benefit manager shall pay a pharmacy a professional dispensing fee at a rate not</u>
- 6 less than that paid in accordance with the current South Dakota Medicaid pharmacy
- 7 dispensing fee schedule, for each prescription pharmaceutical product that is dispensed to the
- 8 patient, by the pharmacy, on a per-unit basis, based on the same generic product identifier
- 9 <u>or generic code number.</u>
- 10 The dispensing fee must be in addition to any amount that the pharmacy benefit manager
- 11 reimburses a pharmacy, consistent with this chapter, for the cost of the pharmaceutical
- 12 product dispensed to the patient.

#### 13 Section 10. That chapter 58-29E be amended with a NEW SECTION:

- 14 <u>A pharmacy benefit manager may not:</u>
- (1) Assess, charge, or collect any form of remuneration or fees from a pharmacy or
   pharmacist, including brand effective rate fees, claim processing fees,
   credentialling fees, dispensing fee effective rate fees, generic effective rate fees,
   network participation fees, and performance-based fees; or
- 19 (2) Directly or indirectly deny or reduce a claim after the claim has been adjudicated,
   20 unless:
- 21 (a) The original claim was submitted fraudulently; or
- (b) The original claim payment was incorrect because the pharmacy or
   pharmacist had already been paid for the pharmaceutical product.
- 24 Section 11. That chapter 58-29E be amended with a NEW SECTION:

25		<u>A pharmacy benefit manager may not:</u>
26	(1)	Take any action that prevents a 340B entity from dispensing drugs purchased
27		under section 340B of the Public Health Service Act, 42 U.S.C. § 256b, as of
28		January 1, 2022, to patients of the 340B entity;
29	<u>(2)</u>	Refuse to contract with a 340B entity or impose on a 340B entity any contracting
30		standards that differ from those imposed on a non-340B entity;

1	<u>(3)</u>	By cor	ntract, provider manual, or any other means:
2		<u>(a)</u>	Modify the definition of a pharmacy, as set forth in chapter 36-11;
3		<u>(b)</u>	Provide a lower reimbursement for a drug purchased under section 340B
4			than that provided for the same drug if purchased by a non-340B entity
5			pharmacy in the same class of trade;
6		<u>(c)</u>	Impose, on a 340B entity, any fee, chargeback, financial or other
7			adjustment, or claims-related information, which is not imposed, in the
8			same manner, on a non-340B entity;
9		<u>(d)</u>	Prevent or otherwise interfere with the ability of covered individuals to
10			receive drugs from a 340B entity of the individual's choice, including
11			through mail order pharmacy services; or
12		<u>(e)</u>	Require or compel the submission of ingredient costs, pricing data, or any
13			other data pertaining to drugs purchased under section 340B.

#### 14 Section 12. That § 58-29E-4 be AMENDED:

**58-29E-4.** A <u>covered entitythird-party payor</u> may request that any pharmacy <u>benefits benefit</u> manager with which it has a pharmacy <u>benefits benefit</u> management services contract disclose to <u>the covered entity it</u>, the amount of all rebate revenues and the nature, type, and amounts of all other revenues that the pharmacy <u>benefits benefit</u> manager receives from each pharmaceutical manufacturer or labeler with whom the pharmacy <u>benefits benefit</u> manager has a contract. The pharmacy <u>benefits benefit</u> manager shall disclose in writing:

- (1) The aggregate amount, and for a list of drugs to be specified in the contract, the
   specific amount, of all rebates and other retrospective utilization discounts received
   by the pharmacy <u>benefits benefit</u> manager, directly or indirectly, from each
   pharmaceutical manufacturer or labeler<del>that</del>, which are earned in connection with
   the dispensing of prescription drugs to covered individuals of the health benefit
   plans issued by the covered entity third-party payor, or for which the covered entity
   third-party payor is the designated administrator;
- (2) The nature, type, and amount of all other revenue received by the pharmacy
  benefits benefit manager, directly or indirectly, from each pharmaceutical
  manufacturer or labeler for any other products or services provided to the
  pharmaceutical manufacturer or labeler by the pharmacy benefits benefit manager,
  with respect to programs that the covered entity third-party payor offers or
  provides to its enrollees; and

(3) Any prescription drug utilization information requested by the covered entity third <u>party payor</u>, relating to covered individuals.

A pharmacy <u>benefits benefit</u> manager shall provide <u>such the</u> information requested by the <u>covered entity third-party payor</u> for <u>such</u> disclosure within thirty days of receipt of the request. If requested, the information <u>shall must</u> be provided no less than once each year. The contract entered into between the pharmacy <u>benefits benefit</u> manager and the covered entity shall third-party payor must set forth any fees to be charged for drug utilization reports requested by the <u>covered entity third-party payor</u>.

#### 9 Section 13. That § 58-29E-5 be AMENDED:

58-29E-5. A pharmacy benefits manager, unless authorized pursuant to the terms
 of its contract with a covered entitythird-party payor, may not contact any covered
 individual without express written permission of the covered entitythird-party payor.

#### 13 Section 14. That § 58-29E-6 be AMENDED:

**58-29E-6.** Except for utilization information, a <u>covered entity third-party payor</u> shall maintain any information disclosed in response to a request pursuant to § 58-29E-4 as confidential and proprietary information, and may not use such information for any other purpose, or disclose <u>such that information</u> to any other person, except as provided in this chapter, or in the pharmacy <u>benefits benefit</u> management services contract between the parties.<del>Any covered entity who</del>

<u>A third-party payor that discloses information in violation of this section is subject</u>
 to an action for injunctive relief and is liable for any damages which that are the direct
 and proximate result of such the disclosure.

Nothing in this section prohibits a covered entity third-party payor from disclosing
 confidential or proprietary information to the director, upon request. Any such-information
 obtained by the director is confidential and privileged and is not open to public inspection
 or disclosure.

#### 27 Section 15. That § 58-29E-7 be AMENDED:

58-29E-7. The covered entity<u>A third-party payor</u> may have the pharmacy benefits
 benefit manager's books and records related to the rebates or other information described
 in subdivisions 58-29E-4(1), (2), and (3)§ 58-29E-4, to the extent the information relates
 directly or indirectly to such covered entity'sthe third-party payor's contract, audited in

accordance with the terms of the pharmacy <u>benefits benefit</u> management services contract
 between the parties. <u>However, ifIf</u> the parties have not expressly provided for audit rights
 and the pharmacy <u>benefits benefit</u> manager has advised the <u>covered entity third-party</u>
 <u>payor</u> that other reasonable options are available and subject to negotiation, the <u>covered</u>
 <u>entity third-party payor</u> may have <u>such the</u> books and records audited as follows:

- 6 (1) <u>Such The audits may be conducted no more frequently than once in each</u>
  7 twelve-month period, upon not less than<u>at least</u> thirty business days' written notice
  8 to the pharmacy <u>benefits benefit manager;</u>
- 9 (2) The covered entity third-party payor may select an independent firm to conduct such the audit, and such. The independent firm shall sign a confidentiality 10 agreement with the covered entity third-party payor and the pharmacy benefits 11 benefit manager, ensuring that all information obtained during such the audit will 12 13 be treated as confidential. The firm may not use, disclose, or otherwise reveal any 14 such of the information, in any manner or form, to any person or entity, except as 15 otherwise permitted under the confidentiality agreement. The covered entity third-16 party payor shall treat all information obtained as a result of the audit as 17 confidential, and may not use or disclose such that information, except as may be 18 otherwise permitted under the terms of the contract between the covered entity third-party payor and the pharmacy benefits benefit manager, or if ordered by a 19 20 court of competent jurisdiction, for good cause shown;
- (3) Any such<u>An</u> audit shall <u>under this section must</u> be conducted at the pharmacy
   benefits <u>benefit</u> manager's office where such the records are located, during normal
   business hours, without undue interference with the pharmacy <u>benefits benefit</u>
   manager's business activities, and in accordance with reasonable audit procedures.
- 25

#### Section 16. That § 58-29E-8 be AMENDED:

- 58-29E-8. With regard to the dispensation of a substitute prescription drug for a
   prescribed drug to a covered individual, when the pharmacy <u>benefits benefit</u> manager
   requests a substitution, the following provisions apply:
- (1) The pharmacy <u>benefits</u> <u>benefit</u> manager may request the substitution of a
   lower- priced generic and therapeutically equivalent drug for a higher-priced
   prescribed drug;
- With regard to substitutions in which the substitute drug's net cost is more for the
   covered individual or the covered entity-third-party payor than the prescribed drug,
   the substitution must be made only for medical reasons that benefit the covered

- individual. If a substitution is being requested pursuant to this subdivision, the
   pharmacy benefits benefit manager shall obtain the approval of the prescribing
   health professional.
- 4 Nothing in this section permits the substitution of an equivalent drug product
  5 contrary to § 36-11-46.2
- 6 Section 17. That § 58-29E-8.1 be AMENDED:
- 58-29E-8.1. A pharmacy benefits benefit manager may neither prohibit nor not
   restrict or penalize a pharmacy or pharmacist or pharmacy for providing cost sharing
   information on the amount a covered individual may pay for a particular prescription drug
   for informing a patient about:
- 11 (1) The cost of a prescription pharmaceutical product;
- 12 (2) The amount of reimbursement that the pharmacy will receive for dispensing the
   13 prescription pharmaceutical product;
- 14 (3) The cost and clinical efficacy of a more affordable alternative pharmaceutical
   15 product, if one is available; or
- (4) Any differential between the amount the patient would pay under the patient's
   prescription benefit plan or program and a lower price the patient would pay for
   the prescription pharmaceutical product, if the patient obtained the pharmaceutical
   product without making a claim for benefits on the patient's prescription benefit
   plan or program.
- 21 Section 18. That § 58-29E-9 be AMENDED:
- 58-29E-9. The Division of Insurance shall promulgate rules, pursuant to chapter
   1-26, to carry out the issuance of the license required by § 58-29E-2 and the enforcement
   provisions of this chapter. The rules may must include the following:
- 25 (1) Definition of terms;
- 26 (2) Use of prescribed forms;
- 27 (3) Reporting requirements;
- 28 (4) Enforcement procedures; and
- 29 (5) Protection of proprietary information and trade secrets.

#### 30 Section 19. That § 58-29E-10 be AMENDED:

58-29E-10. Any covered entity third-party payor may bring a civil action to
 enforce the provisions of this chapter or to seek civil damages for the <u>a</u> violation of its
 provisionsthis chapter.

#### 4 Section 20. That § 58-29E-11 be AMENDED:

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**58-29E-11.** The provisions of Except as otherwise provided in this section, this chapter apply applies only to pharmacy benefits benefit management services contracts entered into or renewed after June 30, 2004.

8 <u>Sections 3 to 110, inclusive, of this Act, apply only to pharmacy benefit</u>
 9 <u>management service contracts entered into or renewed after June 30, 2022.</u>

#### 10 Section 21. That § 58-29E-12 be AMENDED:

**58-29E-12.** <u>No A</u> pharmacy benefit manager <u>shall may not</u> contractually require a pharmacy, who is a participating provider in a health plan provided by a <del>covered</del> <del>entitythird-party payor</del>, to charge or collect, from an insured, a cost share for a prescription or pharmacy service that exceeds the amount retained, by the pharmacist or pharmacy, from all payment sources, for the filling of the prescription or providing the pharmacy service.

#### 17 Section 22. That § 58-29E-13 be AMENDED:

**58-29E-13.** No <u>A</u> pharmacy benefit manager contracting with a covered entity
 shall third-party payor may not retroactively adjust a claim for reimbursement submitted
 by a pharmacy for a prescription drug, unless the adjustment is a result of either of the
 following:

- 22 (1) A pharmacy audit conducted in accordance with chapter 58-29F; or
- 23 (2) A technical billing error.