

**SENATE**  
**STATE OF MINNESOTA**  
**NINETY-THIRD SESSION**

**S.F. No. 168**

(SENATE AUTHORS: MORRISON, Mann, Boldon, Fateh and Hoffman)

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01/11/2023	142	Introduction and first reading Referred to Health and Human Services
01/19/2023	309	Author added Boldon
01/25/2023	383	Withdrawn and re-referred to Commerce and Consumer Protection
01/27/2023	468	Author added Fateh
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03/15/2023		Withdrawn and re-referred to Health and Human Services

1.1 A bill for an act

1.2 relating to health; prohibiting excessive price increases by manufacturers to generic

1.3 or off-patent drugs; authorizing the attorney general to take action against

1.4 manufacturers for certain price increases; prohibiting withdrawal of certain generic

1.5 or off-patent drugs sales; establishing a prescription drug affordability board and

1.6 prescription drug affordability advisory council; providing for prescription drug

1.7 cost reviews and remedies; providing appointments; imposing civil penalties;

1.8 requiring a report; appropriating money; amending Minnesota Statutes 2022,

1.9 section 151.071, subdivisions 1, 2; proposing coding for new law in Minnesota

1.10 Statutes, chapter 62J.

1.11 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MINNESOTA:

1.12 Section 1. **[62J.841] DEFINITIONS.**

1.13 Subdivision 1. **Scope.** For purposes of sections 62J.841 to 62J.845, the following

1.14 definitions apply.

1.15 Subd. 2. **Consumer Price Index.** "Consumer Price Index" means the Consumer Price

1.16 Index, Annual Average, for All Urban Consumers, CPI-U: U.S. City Average, All Items,

1.17 reported by the United States Department of Labor, Bureau of Labor Statistics, or its

1.18 successor or, if the index is discontinued, an equivalent index reported by a federal authority

1.19 or, if no such index is reported, "Consumer Price Index" means a comparable index chosen

1.20 by the Bureau of Labor Statistics.

1.21 Subd. 3. **Generic or off-patent drug.** "Generic or off-patent drug" means any prescription

1.22 drug for which any exclusive marketing rights granted under the Federal Food, Drug, and

1.23 Cosmetic Act, section 351 of the federal Public Health Service Act, and federal patent law

1.24 have expired, including any drug-device combination product for the delivery of a generic

1.25 drug.

2.1 Subd. 4. **Manufacturer.** "Manufacturer" has the meaning provided in section 151.01,  
2.2 subdivision 14a, but does not include an entity required solely because the entity repackages  
2.3 or relabels drugs.

2.4 Subd. 5. **Prescription drug.** "Prescription drug" means a drug for human use subject  
2.5 to United States Code, title 21, section 353(b)(1).

2.6 Subd. 6. **Wholesale acquisition cost.** "Wholesale acquisition cost" has the meaning  
2.7 provided in United States Code, title 42, section 1395w-3a.

2.8 Subd. 7. **Wholesale distributor.** "Wholesale distributor" has the meaning provided in  
2.9 section 151.441, subdivision 14.

2.10 **Sec. 2. [62J.842] EXCESSIVE PRICE INCREASES PROHIBITED.**

2.11 Subdivision 1. **Prohibition.** No manufacturer shall impose, or cause to be imposed, an  
2.12 excessive price increase, whether directly or through a wholesale distributor, pharmacy, or  
2.13 similar intermediary, on the sale of any generic or off-patent drug sold, dispensed, or  
2.14 delivered to any consumer in the state.

2.15 Subd. 2. **Excessive price increase.** A price increase is excessive for purposes of this  
2.16 section when:

2.17 (1) the price increase, adjusted for inflation utilizing the Consumer Price Index, exceeds:

2.18 (i) 15 percent of the wholesale acquisition cost over the immediately preceding calendar  
2.19 year; or

2.20 (ii) 40 percent of the wholesale acquisition cost over the immediately preceding three  
2.21 calendar years; and

2.22 (2) the price increase, adjusted for inflation utilizing the Consumer Price Index, exceeds  
2.23 \$30 for:

2.24 (i) a 30-day supply of the drug; or

2.25 (ii) a course of treatment lasting less than 30 days.

2.26 Subd. 3. **Exemption.** It is not a violation of this section for a wholesale distributor or  
2.27 pharmacy to increase the price of a generic or off-patent drug if the price increase is directly  
2.28 attributable to additional costs for the drug imposed on the wholesale distributor or pharmacy  
2.29 by the manufacturer of the drug.

3.1 Sec. 3. **[62J.843] REGISTERED AGENT AND OFFICE WITHIN THE STATE.**

3.2 Any manufacturer that sells, distributes, delivers, or offers for sale any generic or  
 3.3 off-patent drug in the state must maintain a registered agent and office within the state.

3.4 Sec. 4. **[62J.844] ENFORCEMENT.**

3.5 Subdivision 1. **Notification.** (a) The commissioner of health shall notify the manufacturer  
 3.6 of a generic or off-patent drug, the attorney general, and the Board of Pharmacy of any price  
 3.7 increase that the commissioner believes may violate section 62J.842.

3.8 (b) The commissioner of management and budget and any other state agency that provides  
 3.9 or purchases a pharmacy benefit except the Department of Human Services, and any entity  
 3.10 under contract with a state agency to provide a pharmacy benefit other than an entity under  
 3.11 contract with the Department of Human Services, may notify the manufacturer of a generic  
 3.12 or off-patent drug, the attorney general, and the Board of Pharmacy of any price increase  
 3.13 that the commissioner or entity believes may violate section 62J.842.

3.14 Subd. 2. **Submission of drug cost statement and other information by manufacturer;**  
 3.15 **investigation by attorney general.** (a) Within 45 days of receiving a notice under subdivision  
 3.16 1, the manufacturer of the generic or off-patent drug shall submit a drug cost statement to  
 3.17 the attorney general. The statement must:

3.18 (1) itemize the cost components related to production of the drug;

3.19 (2) identify the circumstances and timing of any increase in materials or manufacturing  
 3.20 costs that caused any increase during the preceding calendar year, or preceding three calendar  
 3.21 years as applicable, in the price of the drug; and

3.22 (3) provide any other information that the manufacturer believes to be relevant to a  
 3.23 determination of whether a violation of section 62J.842 has occurred.

3.24 (b) The attorney general may investigate whether a violation of section 62J.842 has  
 3.25 occurred, in accordance with section 8.31, subdivision 2.

3.26 Subd. 3. **Petition to court.** (a) On petition of the attorney general, a court may issue an  
 3.27 order:

3.28 (1) compelling the manufacturer of a generic or off-patent drug to:

3.29 (i) provide the drug cost statement required under subdivision 2, paragraph (a); and

3.30 (ii) answer interrogatories, produce records or documents, or be examined under oath,  
 3.31 as required by the attorney general under subdivision 2, paragraph (b);

4.1 (2) restraining or enjoining a violation of sections 62J.841 to 62J.845, including issuing  
 4.2 an order requiring that drug prices be restored to levels that comply with section 62J.842;

4.3 (3) requiring the manufacturer to provide an accounting to the attorney general of all  
 4.4 revenues resulting from a violation of section 62J.842;

4.5 (4) requiring the manufacturer to repay to all Minnesota consumers, including any  
 4.6 third-party payers, any money acquired as a result of a price increase that violates section  
 4.7 62J.842;

4.8 (5) notwithstanding section 16A.151, requiring that all revenues generated from a  
 4.9 violation of section 62J.842 be remitted to the state and deposited into a special fund, to be  
 4.10 used for initiatives to reduce the cost to consumers of acquiring prescription drugs, if a  
 4.11 manufacturer is unable to determine the individual transactions necessary to provide the  
 4.12 repayments described in clause (4);

4.13 (6) imposing a civil penalty of up to \$10,000 per day for each violation of section 62J.842;

4.14 (7) providing for the attorney general's recovery of costs and disbursements incurred in  
 4.15 bringing an action against a manufacturer found in violation of section 62J.842, including  
 4.16 the costs of investigation and reasonable attorney's fees; and

4.17 (8) providing any other appropriate relief, including any other equitable relief as  
 4.18 determined by the court.

4.19 (b) For purposes of paragraph (a), clause (6), every individual transaction in violation  
 4.20 of section 62J.842 is considered a separate violation.

4.21 Subd. 4. **Private right of action.** Any action brought pursuant to section 8.31, subdivision  
 4.22 3a, by a person injured by a violation of section 62J.842 is for the benefit of the public.

4.23 Sec. 5. **[62J.845] PROHIBITION ON WITHDRAWAL OF GENERIC OR**  
 4.24 **OFF-PATENT DRUGS FOR SALE.**

4.25 Subdivision 1. **Prohibition.** A manufacturer of a generic or off-patent drug is prohibited  
 4.26 from withdrawing that drug from sale or distribution within this state for the purpose of  
 4.27 avoiding the prohibition on excessive price increases under section 62J.842.

4.28 Subd. 2. **Notice to board and attorney general.** Any manufacturer that intends to  
 4.29 withdraw a generic or off-patent drug from sale or distribution within the state shall provide  
 4.30 a written notice of withdrawal to the Board of Pharmacy and the attorney general, at least  
 4.31 90 days prior to the withdrawal.

5.1 Subd. 3. **Financial penalty.** The attorney general shall assess a penalty of \$500,000 on  
5.2 any manufacturer of a generic or off-patent drug that the attorney general determines has  
5.3 failed to comply with the requirements of this section.

5.4 Sec. 6. [62J.846] SEVERABILITY.

5.5 If any provision of sections 62J.841 to 62J.845 or the application thereof to any person  
5.6 or circumstance is held invalid for any reason in a court of competent jurisdiction, the  
5.7 invalidity does not affect other provisions or any other application of sections 62J.841 to  
5.8 62J.845 that can be given effect without the invalid provision or application.

5.9 Sec. 7. [62J.85] CITATION.

5.10 Sections 62J.85 to 62J.95 may be cited as the "Prescription Drug Affordability Act."

5.11 Sec. 8. [62J.86] DEFINITIONS.

5.12 Subdivision 1. **Definitions.** For the purposes of sections 62J.85 to 62J.95, the following  
5.13 terms have the meanings given them.

5.14 Subd. 2. **Advisory council.** "Advisory council" means the Prescription Drug Affordability  
5.15 Advisory Council established under section 62J.88.

5.16 Subd. 3. **Biologic.** "Biologic" means a drug that is produced or distributed in accordance  
5.17 with a biologics license application approved under Code of Federal Regulations, title 42,  
5.18 section 447.502.

5.19 Subd. 4. **Biosimilar.** "Biosimilar" has the meaning provided in section 62J.84, subdivision  
5.20 2, paragraph (b).

5.21 Subd. 5. **Board.** "Board" means the Prescription Drug Affordability Board established  
5.22 under section 62J.87.

5.23 Subd. 6. **Brand name drug.** "Brand name drug" means a drug that is produced or  
5.24 distributed pursuant to:

5.25 (1) a new drug application approved under United States Code, title 21, section 355(c),  
5.26 except for a generic drug as defined under Code of Federal Regulations, title 42, section  
5.27 447.502; or

5.28 (2) a biologics license application approved under United States Code, title 45, section  
5.29 262(a)(c).

6.1 Subd. 7. **Generic drug.** "Generic drug" has the meaning provided in section 62J.84,  
6.2 subdivision 2, paragraph (e).

6.3 Subd. 8. **Group purchaser.** "Group purchaser" has the meaning given in section 62J.03,  
6.4 subdivision 6, and includes pharmacy benefit managers as defined in section 62W.02,  
6.5 subdivision 15.

6.6 Subd. 9. **Manufacturer.** "Manufacturer" means an entity that:

6.7 (1) engages in the manufacture of a prescription drug product or enters into a lease with  
6.8 another manufacturer to market and distribute a prescription drug product under the entity's  
6.9 own name; and

6.10 (2) sets or changes the wholesale acquisition cost of the prescription drug product it  
6.11 manufactures or markets.

6.12 Subd. 10. **Prescription drug product.** "Prescription drug product" means a brand name  
6.13 drug, a generic drug, a biologic, or a biosimilar.

6.14 Subd. 11. **Wholesale acquisition cost or WAC.** "Wholesale acquisition cost" or "WAC"  
6.15 has the meaning given in United States Code, title 42, section 1395W-3a(c)(6)(B).

6.16 Sec. 9. **[62J.87] PRESCRIPTION DRUG AFFORDABILITY BOARD.**

6.17 Subdivision 1. **Establishment.** The commissioner of commerce shall establish the  
6.18 Prescription Drug Affordability Board, which shall be governed as a board under section  
6.19 15.012, paragraph (a), to protect consumers, state and local governments, health plan  
6.20 companies, providers, pharmacies, and other health care system stakeholders from  
6.21 unaffordable costs of certain prescription drugs.

6.22 Subd. 2. **Membership.** (a) The Prescription Drug Affordability Board consists of eleven  
6.23 members appointed as follows:

6.24 (1) seven voting members appointed by the governor;

6.25 (2) one nonvoting member appointed by the majority leader of the senate;

6.26 (3) one nonvoting member appointed by the minority leader of the senate;

6.27 (4) one nonvoting member appointed by the speaker of the house; and

6.28 (5) one nonvoting member appointed by the minority leader of the house of  
6.29 representatives.

6.30 (b) All members appointed must have knowledge and demonstrated expertise in  
6.31 pharmaceutical economics and finance or health care economics and finance. A member

7.1 must not be an employee of, a board member of, or a consultant to a manufacturer or trade  
7.2 association for manufacturers or a pharmacy benefit manager or trade association for  
7.3 pharmacy benefit managers.

7.4 (c) Initial appointments must be made by January 1, 2024.

7.5 Subd. 3. **Terms.** (a) Board appointees shall serve four-year terms, except that initial  
7.6 appointees shall serve staggered terms of two, three, or four years as determined by lot by  
7.7 the secretary of state. A board member shall serve no more than two consecutive terms.

7.8 (b) A board member may resign at any time by giving written notice to the board.

7.9 Subd. 4. **Chair; other officers.** (a) The governor shall designate an acting chair from  
7.10 the members appointed by the governor.

7.11 (b) The board shall elect a chair to replace the acting chair at the first meeting of the  
7.12 board by a majority of the members. The chair shall serve for one year.

7.13 (c) The board shall elect a vice-chair and other officers from its membership as it deems  
7.14 necessary.

7.15 Subd. 5. **Staff; technical assistance.** (a) The board shall hire an executive director and  
7.16 other staff, who shall serve in the unclassified service. The executive director must have  
7.17 knowledge and demonstrated expertise in pharmacoeconomics, pharmacology, health policy,  
7.18 health services research, medicine, or a related field or discipline.

7.19 (b) The commissioner of health shall provide technical assistance to the board. The board  
7.20 may also employ or contract for professional and technical assistance as the board deems  
7.21 necessary to perform the board's duties.

7.22 (c) The attorney general shall provide legal services to the board.

7.23 Subd. 6. **Compensation.** The board members shall not receive compensation but may  
7.24 receive reimbursement for expenses as authorized under section 15.059, subdivision 3.

7.25 Subd. 7. **Meetings.** (a) Meetings of the board are subject to chapter 13D. The board shall  
7.26 meet publicly at least every three months to review prescription drug product information  
7.27 submitted to the board under section 62J.90. If there are no pending submissions, the chair  
7.28 of the board may cancel or postpone the required meeting. The board may meet in closed  
7.29 session when reviewing proprietary information as determined under the standards developed  
7.30 in accordance with section 62J.91, subdivision 3.

8.1 (b) The board shall announce each public meeting at least three weeks prior to the  
 8.2 scheduled date of the meeting. Any materials for the meeting shall be made public at least  
 8.3 two weeks prior to the scheduled date of the meeting.

8.4 (c) At each public meeting, the board shall provide the opportunity for comments from  
 8.5 the public, including the opportunity for written comments to be submitted to the board  
 8.6 prior to a decision by the board.

8.7 **Sec. 10. [62J.88] PRESCRIPTION DRUG AFFORDABILITY ADVISORY**  
 8.8 **COUNCIL.**

8.9 Subdivision 1. **Establishment.** The governor shall appoint a 17-member stakeholder  
 8.10 advisory council to provide advice to the board on drug cost issues and to represent  
 8.11 stakeholders' views. The governor shall appoint the members of the advisory council based  
 8.12 on the members' knowledge and demonstrated expertise in one or more of the following  
 8.13 areas: the pharmaceutical business; practice of medicine; patient perspectives; health care  
 8.14 cost trends and drivers; clinical and health services research; and the health care marketplace.

8.15 Subd. 2. **Membership.** The council's membership shall consist of the following:

8.16 (1) two members representing patients and health care consumers;

8.17 (2) two members representing health care providers;

8.18 (3) one member representing health plan companies;

8.19 (4) two members representing employers, with one member representing large employers  
 8.20 and one member representing small employers;

8.21 (5) one member representing government employee benefit plans;

8.22 (6) one member representing pharmaceutical manufacturers;

8.23 (7) one member who is a health services clinical researcher;

8.24 (8) one member who is a pharmacologist;

8.25 (9) one member representing the commissioner of health with expertise in health  
 8.26 economics;

8.27 (10) one member representing pharmaceutical wholesalers;

8.28 (11) one member representing pharmacy benefit managers;

8.29 (12) one member from the Rare Disease Advisory Council;

8.30 (13) one member representing generic drug manufacturers; and



9.1 (14) one member representing pharmaceutical distributors.

9.2 Subd. 3. **Terms.** (a) The initial appointments to the advisory council must be made by  
 9.3 January 1, 2024. The initial appointed advisory council members shall serve staggered terms  
 9.4 of two, three, or four years determined by lot by the secretary of state. Following the initial  
 9.5 appointments, the advisory council members shall serve four-year terms.

9.6 (b) Removal and vacancies of advisory council members shall be governed by section  
 9.7 15.059.

9.8 Subd. 4. **Compensation.** Advisory council members may be compensated according to  
 9.9 section 15.059.

9.10 Subd. 5. **Meetings.** Meetings of the advisory council are subject to chapter 13D. The  
 9.11 advisory council shall meet publicly at least every three months to advise the board on drug  
 9.12 cost issues related to the prescription drug product information submitted to the board under  
 9.13 section 62J.90.

9.14 Subd. 6. **Exemption.** Notwithstanding section 15.059, the advisory council shall not  
 9.15 expire.

9.16 **Sec. 11. [62J.89] CONFLICTS OF INTEREST.**

9.17 Subdivision 1. **Definition.** For purposes of this section, "conflict of interest" means a  
 9.18 financial or personal association that has the potential to bias or have the appearance of  
 9.19 biasing a person's decisions in matters related to the board, the advisory council, or in the  
 9.20 conduct of the board's or council's activities. A conflict of interest includes any instance in  
 9.21 which a person, a person's immediate family member, including a spouse, parent, child, or  
 9.22 other legal dependent, or an in-law of any of the preceding individuals, has received or  
 9.23 could receive a direct or indirect financial benefit of any amount deriving from the result  
 9.24 or findings of a decision or determination of the board. For purposes of this section, a  
 9.25 financial benefit includes honoraria, fees, stock, the value of the member's, immediate family  
 9.26 member's, or in-law's stock holdings, and any direct financial benefit deriving from the  
 9.27 finding of a review conducted under sections 62J.85 to 62J.95. Ownership of securities is  
 9.28 not a conflict of interest if the securities are: (1) part of a diversified mutual or exchange  
 9.29 traded fund; or (2) in a tax-deferred or tax-exempt retirement account that is administered  
 9.30 by an independent trustee.

9.31 Subd. 2. **General.** (a) Prior to the acceptance of an appointment or employment, or prior  
 9.32 to entering into a contractual agreement, a board or advisory council member, board staff  
 9.33 member, or third-party contractor must disclose to the appointing authority or the board

10.1 any conflicts of interest. The information disclosed must include the type, nature, and  
 10.2 magnitude of the interests involved.

10.3 (b) A board member, board staff member, or third-party contractor with a conflict of  
 10.4 interest with regard to any prescription drug product under review must recuse themselves  
 10.5 from any discussion, review, decision, or determination made by the board relating to the  
 10.6 prescription drug product.

10.7 (c) Any conflict of interest must be disclosed in advance of the first meeting after the  
 10.8 conflict is identified or within five days after the conflict is identified, whichever is earlier.

10.9 Subd. 3. **Prohibitions.** Board members, board staff, or third-party contractors are  
 10.10 prohibited from accepting gifts, bequeaths, or donations of services or property that raise  
 10.11 the specter of a conflict of interest or have the appearance of injecting bias into the activities  
 10.12 of the board.

10.13 Sec. 12. **[62J.90] PRESCRIPTION DRUG PRICE INFORMATION; DECISION**  
 10.14 **TO CONDUCT COST REVIEW.**

10.15 Subdivision 1. **Drug price information from the commissioner of health and other**  
 10.16 **sources.** (a) The commissioner of health shall provide to the board the information reported  
 10.17 to the commissioner by drug manufacturers under section 62J.84, subdivisions 3, 4, and 5.  
 10.18 The commissioner shall provide this information to the board within 30 days of the date the  
 10.19 information is received from drug manufacturers.

10.20 (b) The board may subscribe to one or more prescription drug pricing files, such as  
 10.21 Medispan or FirstDatabank, or as otherwise determined by the board.

10.22 Subd. 2. **Identification of certain prescription drug products.** (a) The board, in  
 10.23 consultation with the advisory council, shall identify selected prescription drug products  
 10.24 based on the following criteria:

10.25 (1) brand name drugs or biologics for which the WAC increases by more than 15 percent  
 10.26 or by more than \$3,000 during any 12-month period or course of treatment if less than 12  
 10.27 months, after adjusting for changes in the consumer price index (CPI);

10.28 (2) brand name drugs or biologics with a WAC of \$60,000 or more per calendar year  
 10.29 or per course of treatment;

10.30 (3) biosimilar drugs that have a WAC that is not at least 20 percent lower than the  
 10.31 referenced brand name biologic at the time the biosimilar is introduced; and

10.32 (4) generic drugs for which the WAC:

11.1 (i) is \$100 or more, after adjusting for changes in the CPI, for:

11.2 (A) a 30-day supply lasting a patient for 30 consecutive days based on the recommended  
 11.3 dosage approved for labeling by the United States Food and Drug Administration (FDA);

11.4 (B) a supply lasting a patient for fewer than 30 days based on recommended dosage  
 11.5 approved for labeling by the FDA; or

11.6 (C) one unit of the drug if the labeling approved by the FDA does not recommend a  
 11.7 finite dosage; and

11.8 (ii) is increased by 200 percent or more during the immediate preceding 12-month period,  
 11.9 as determined by the difference between the resulting WAC and the average of the WAC  
 11.10 reported over the preceding 12 months, after adjusting for changes in the CPI.

11.11 (b) The board, in consultation with the advisory council and the commissioner of health,  
 11.12 may identify prescription drug products not described in paragraph (a) that may impose  
 11.13 costs that create significant affordability challenges for the state health care system or for  
 11.14 patients, including but not limited to drugs to address public health emergencies.

11.15 (c) The board shall make available to the public the names and related price information  
 11.16 of the prescription drug products identified under this subdivision, with the exception of  
 11.17 information determined by the board to be proprietary under the standards developed by  
 11.18 the board under section 62J.91, subdivision 3, and information provided by the commissioner  
 11.19 of health classified as not public data under section 13.02, subdivision 8a, or as trade secret  
 11.20 information under section 13.37, subdivision 1, paragraph (b), or as trade secret information  
 11.21 under the Defend Trade Secrets Act of 2016, United States Code, title 18, section 1836, as  
 11.22 amended.

11.23 Subd. 3. **Determination to proceed with review.** (a) The board may initiate a cost  
 11.24 review of a prescription drug product identified by the board under this section.

11.25 (b) The board shall consider requests by the public for the board to proceed with a cost  
 11.26 review of any prescription drug product identified under this section.

11.27 (c) If there is no consensus among the members of the board on whether to initiate a  
 11.28 cost review of a prescription drug product, any member of the board may request a vote to  
 11.29 determine whether to review the cost of the prescription drug product.

11.30 Sec. 13. **[62J.91] PRESCRIPTION DRUG PRODUCT REVIEWS.**

11.31 Subdivision 1. **General.** Once a decision by the board has been made to proceed with  
 11.32 a cost review of a prescription drug product, the board shall conduct the review and make

12.1 a determination as to whether appropriate utilization of the prescription drug under review,  
12.2 based on utilization that is consistent with the United States Food and Drug Administration  
12.3 (FDA) label or standard medical practice, has led or will lead to affordability challenges  
12.4 for the state health care system or for patients.

12.5 Subd. 2. **Review considerations.** In reviewing the cost of a prescription drug product,  
12.6 the board may consider the following factors:

12.7 (1) the price at which the prescription drug product has been and will be sold in the state;

12.8 (2) manufacturer monetary price concessions, discounts, or rebates, and drug-specific  
12.9 patient assistance;

12.10 (3) the price of therapeutic alternatives;

12.11 (4) the cost to group purchasers based on patient access consistent with the FDA-labeled  
12.12 indications and standard medical practice;

12.13 (5) measures of patient access, including cost-sharing and other metrics;

12.14 (6) the extent to which the attorney general or a court has determined that a price increase  
12.15 for a generic or off-patent prescription drug product was excessive under sections 62J.842  
12.16 and 62J.844;

12.17 (7) any information a manufacturer chooses to provide; and

12.18 (8) any other factors as determined by the board.

12.19 Subd. 3. **Public data; proprietary information.** (a) Any submission made to the board  
12.20 related to a drug cost review must be made available to the public with the exception of  
12.21 information determined by the board to be proprietary and information provided by the  
12.22 commissioner of health classified as not public data under section 13.02, subdivision 8a, or  
12.23 as trade secret information under section 13.37, subdivision 1, paragraph (b), or as trade  
12.24 secret information under the Defend Trade Secrets Act of 2016, United States Code, title  
12.25 18, section 1836, as amended.

12.26 (b) The board shall establish the standards for the information to be considered proprietary  
12.27 under paragraph (a) and section 62J.90, subdivision 2, including standards for heightened  
12.28 consideration of proprietary information for submissions for a cost review of a drug that is  
12.29 not yet approved by the FDA.

12.30 (c) Prior to the board establishing the standards under paragraph (b), the public shall be  
12.31 provided notice and the opportunity to submit comments.

13.1 (d) The establishment of standards under this subdivision is exempt from the rulemaking  
13.2 requirements under chapter 14, and section 14.386 does not apply.

13.3 **Sec. 14. [62J.92] DETERMINATIONS; COMPLIANCE; REMEDIES.**

13.4 Subdivision 1. **Upper payment limit.** (a) In the event the board finds that the spending  
13.5 on a prescription drug product reviewed under section 62J.91 creates an affordability  
13.6 challenge for the state health care system or for patients, the board shall establish an upper  
13.7 payment limit after considering:

13.8 (1) extraordinary supply costs, if applicable;

13.9 (2) the range of prices at which the drug is sold in the United States according to one or  
13.10 more pricing files accessed under section 62J.90, subdivision 1, and the range at which  
13.11 pharmacies are reimbursed in Canada; and

13.12 (3) any other relevant pricing and administrative cost information for the drug.

13.13 (b) An upper payment limit applies to all purchases of, and payer reimbursements for,  
13.14 a prescription drug that is dispensed or administered to individuals in the state in person,  
13.15 by mail, or by other means, and for which an upper payment limit has been established.

13.16 Subd. 2. **Implementation and administration of the upper payment limit.** (a) An  
13.17 upper payment limit may take effect no sooner than 120 days following the date of its public  
13.18 release by the board.

13.19 (b) When setting an upper payment limit for a drug subject to the Medicare maximum  
13.20 fair price under United States Code, title 42, section 1191(c), the board shall set the upper  
13.21 payment limit at the Medicare maximum fair price.

13.22 (c) Health plan companies and pharmacy benefit managers shall report annually to the  
13.23 board, in the form and manner specified by the board, on how cost savings resulting from  
13.24 the establishment of an upper payment limit have been used by the health plan company or  
13.25 pharmacy benefit manager to benefit enrollees, including but not limited to reducing enrollee  
13.26 cost-sharing.

13.27 Subd. 3. **Noncompliance.** (a) The board shall, and other persons may, notify the Office  
13.28 of the Attorney General of a potential failure by an entity subject to an upper payment limit  
13.29 to comply with that limit.

13.30 (b) If the Office of the Attorney General finds that an entity was noncompliant with the  
13.31 upper payment limit requirements, the attorney general may pursue remedies consistent  
13.32 with chapter 8 or appropriate criminal charges if there is evidence of intentional profiteering.

14.1 (c) An entity who obtains price concessions from a drug manufacturer that result in a  
14.2 lower net cost to the stakeholder than the upper payment limit established by the board is  
14.3 not considered noncompliant.

14.4 (d) The Office of the Attorney General may provide guidance to stakeholders concerning  
14.5 activities that could be considered noncompliant.

14.6 Subd. 4. Appeals. (a) Persons affected by a decision of the board may request an appeal  
14.7 of the board's decision within 30 days of the date of the decision. The board shall hear the  
14.8 appeal and render a decision within 60 days of the hearing.

14.9 (b) All appeal decisions are subject to judicial review in accordance with chapter 14.

14.10 Sec. 15. **[62J.93] REPORTS.**

14.11 Beginning March 1, 2024, and each March 1 thereafter, the board shall submit a report  
14.12 to the governor and legislature on general price trends for prescription drug products and  
14.13 the number of prescription drug products that were subject to the board's cost review and  
14.14 analysis, including the result of any analysis as well as the number and disposition of appeals  
14.15 and judicial reviews.

14.16 Sec. 16. **[62J.94] ERISA PLANS AND MEDICARE DRUG PLANS.**

14.17 (a) Nothing in sections 62J.85 to 62J.95 shall be construed to require ERISA plans or  
14.18 Medicare Part D plans to comply with decisions of the board. ERISA plans or Medicare  
14.19 Part D plans are free to choose to exceed the upper payment limit established by the board  
14.20 under section 62J.92.

14.21 (b) Providers who dispense and administer drugs in the state must bill all payers no more  
14.22 than the upper payment limit without regard to whether an ERISA plan or Medicare Part  
14.23 D plan chooses to reimburse the provider in an amount greater than the upper payment limit  
14.24 established by the board.

14.25 (c) For purposes of this section, an ERISA plan or group health plan is an employee  
14.26 welfare benefit plan established by or maintained by an employer or an employee  
14.27 organization, or both, that provides employer sponsored health coverage to employees and  
14.28 the employee's dependents and is subject to the Employee Retirement Income Security Act  
14.29 of 1974 (ERISA).

15.1 Sec. 17. **[62J.95] SEVERABILITY.**

15.2 If any provision of sections 62J.85 to 62J.94 or the application thereof to any person or  
15.3 circumstance is held invalid for any reason in a court of competent jurisdiction, the invalidity  
15.4 does not affect other provisions or any other application of sections 62J.85 to 62J.94 that  
15.5 can be given effect without the invalid provision or application.

15.6 Sec. 18. Minnesota Statutes 2022, section 151.071, subdivision 1, is amended to read:

15.7 Subdivision 1. **Forms of disciplinary action.** When the board finds that a licensee,  
15.8 registrant, or applicant has engaged in conduct prohibited under subdivision 2, it may do  
15.9 one or more of the following:

15.10 (1) deny the issuance of a license or registration;

15.11 (2) refuse to renew a license or registration;

15.12 (3) revoke the license or registration;

15.13 (4) suspend the license or registration;

15.14 (5) impose limitations, conditions, or both on the license or registration, including but  
15.15 not limited to: the limitation of practice to designated settings; the limitation of the scope  
15.16 of practice within designated settings; the imposition of retraining or rehabilitation  
15.17 requirements; the requirement of practice under supervision; the requirement of participation  
15.18 in a diversion program such as that established pursuant to section 214.31 or the conditioning  
15.19 of continued practice on demonstration of knowledge or skills by appropriate examination  
15.20 or other review of skill and competence;

15.21 (6) impose a civil penalty not exceeding \$10,000 for each separate violation, except that  
15.22 a civil penalty not exceeding \$25,000 may be imposed for each separate violation of section  
15.23 62J.842, the amount of the civil penalty to be fixed so as to deprive a licensee or registrant  
15.24 of any economic advantage gained by reason of the violation, to discourage similar violations  
15.25 by the licensee or registrant or any other licensee or registrant, or to reimburse the board  
15.26 for the cost of the investigation and proceeding, including but not limited to, fees paid for  
15.27 services provided by the Office of Administrative Hearings, legal and investigative services  
15.28 provided by the Office of the Attorney General, court reporters, witnesses, reproduction of  
15.29 records, board members' per diem compensation, board staff time, and travel costs and  
15.30 expenses incurred by board staff and board members; and

15.31 (7) reprimand the licensee or registrant.

16.1 Sec. 19. Minnesota Statutes 2022, section 151.071, subdivision 2, is amended to read:

16.2 Subd. 2. **Grounds for disciplinary action.** The following conduct is prohibited and is  
16.3 grounds for disciplinary action:

16.4 (1) failure to demonstrate the qualifications or satisfy the requirements for a license or  
16.5 registration contained in this chapter or the rules of the board. The burden of proof is on  
16.6 the applicant to demonstrate such qualifications or satisfaction of such requirements;

16.7 (2) obtaining a license by fraud or by misleading the board in any way during the  
16.8 application process or obtaining a license by cheating, or attempting to subvert the licensing  
16.9 examination process. Conduct that subverts or attempts to subvert the licensing examination  
16.10 process includes, but is not limited to: (i) conduct that violates the security of the examination  
16.11 materials, such as removing examination materials from the examination room or having  
16.12 unauthorized possession of any portion of a future, current, or previously administered  
16.13 licensing examination; (ii) conduct that violates the standard of test administration, such as  
16.14 communicating with another examinee during administration of the examination, copying  
16.15 another examinee's answers, permitting another examinee to copy one's answers, or  
16.16 possessing unauthorized materials; or (iii) impersonating an examinee or permitting an  
16.17 impersonator to take the examination on one's own behalf;

16.18 (3) for a pharmacist, pharmacy technician, pharmacist intern, applicant for a pharmacist  
16.19 or pharmacy license, or applicant for a pharmacy technician or pharmacist intern registration,  
16.20 conviction of a felony reasonably related to the practice of pharmacy. Conviction as used  
16.21 in this subdivision includes a conviction of an offense that if committed in this state would  
16.22 be deemed a felony without regard to its designation elsewhere, or a criminal proceeding  
16.23 where a finding or verdict of guilt is made or returned but the adjudication of guilt is either  
16.24 withheld or not entered thereon. The board may delay the issuance of a new license or  
16.25 registration if the applicant has been charged with a felony until the matter has been  
16.26 adjudicated;

16.27 (4) for a facility, other than a pharmacy, licensed or registered by the board, if an owner  
16.28 or applicant is convicted of a felony reasonably related to the operation of the facility. The  
16.29 board may delay the issuance of a new license or registration if the owner or applicant has  
16.30 been charged with a felony until the matter has been adjudicated;

16.31 (5) for a controlled substance researcher, conviction of a felony reasonably related to  
16.32 controlled substances or to the practice of the researcher's profession. The board may delay  
16.33 the issuance of a registration if the applicant has been charged with a felony until the matter  
16.34 has been adjudicated;



17.1 (6) disciplinary action taken by another state or by one of this state's health licensing  
17.2 agencies:

17.3 (i) revocation, suspension, restriction, limitation, or other disciplinary action against a  
17.4 license or registration in another state or jurisdiction, failure to report to the board that  
17.5 charges or allegations regarding the person's license or registration have been brought in  
17.6 another state or jurisdiction, or having been refused a license or registration by any other  
17.7 state or jurisdiction. The board may delay the issuance of a new license or registration if an  
17.8 investigation or disciplinary action is pending in another state or jurisdiction until the  
17.9 investigation or action has been dismissed or otherwise resolved; and

17.10 (ii) revocation, suspension, restriction, limitation, or other disciplinary action against a  
17.11 license or registration issued by another of this state's health licensing agencies, failure to  
17.12 report to the board that charges regarding the person's license or registration have been  
17.13 brought by another of this state's health licensing agencies, or having been refused a license  
17.14 or registration by another of this state's health licensing agencies. The board may delay the  
17.15 issuance of a new license or registration if a disciplinary action is pending before another  
17.16 of this state's health licensing agencies until the action has been dismissed or otherwise  
17.17 resolved;

17.18 (7) for a pharmacist, pharmacy, pharmacy technician, or pharmacist intern, violation of  
17.19 any order of the board, of any of the provisions of this chapter or any rules of the board or  
17.20 violation of any federal, state, or local law or rule reasonably pertaining to the practice of  
17.21 pharmacy;

17.22 (8) for a facility, other than a pharmacy, licensed by the board, violations of any order  
17.23 of the board, of any of the provisions of this chapter or the rules of the board or violation  
17.24 of any federal, state, or local law relating to the operation of the facility;

17.25 (9) engaging in any unethical conduct; conduct likely to deceive, defraud, or harm the  
17.26 public, or demonstrating a willful or careless disregard for the health, welfare, or safety of  
17.27 a patient; or pharmacy practice that is professionally incompetent, in that it may create  
17.28 unnecessary danger to any patient's life, health, or safety, in any of which cases, proof of  
17.29 actual injury need not be established;

17.30 (10) aiding or abetting an unlicensed person in the practice of pharmacy, except that it  
17.31 is not a violation of this clause for a pharmacist to supervise a properly registered pharmacy  
17.32 technician or pharmacist intern if that person is performing duties allowed by this chapter  
17.33 or the rules of the board;

18.1 (11) for an individual licensed or registered by the board, adjudication as mentally ill  
18.2 or developmentally disabled, or as a chemically dependent person, a person dangerous to  
18.3 the public, a sexually dangerous person, or a person who has a sexual psychopathic  
18.4 personality, by a court of competent jurisdiction, within or without this state. Such  
18.5 adjudication shall automatically suspend a license for the duration thereof unless the board  
18.6 orders otherwise;

18.7 (12) for a pharmacist or pharmacy intern, engaging in unprofessional conduct as specified  
18.8 in the board's rules. In the case of a pharmacy technician, engaging in conduct specified in  
18.9 board rules that would be unprofessional if it were engaged in by a pharmacist or pharmacist  
18.10 intern or performing duties specifically reserved for pharmacists under this chapter or the  
18.11 rules of the board;

18.12 (13) for a pharmacy, operation of the pharmacy without a pharmacist present and on  
18.13 duty except as allowed by a variance approved by the board;

18.14 (14) for a pharmacist, the inability to practice pharmacy with reasonable skill and safety  
18.15 to patients by reason of illness, use of alcohol, drugs, narcotics, chemicals, or any other type  
18.16 of material or as a result of any mental or physical condition, including deterioration through  
18.17 the aging process or loss of motor skills. In the case of registered pharmacy technicians,  
18.18 pharmacist interns, or controlled substance researchers, the inability to carry out duties  
18.19 allowed under this chapter or the rules of the board with reasonable skill and safety to  
18.20 patients by reason of illness, use of alcohol, drugs, narcotics, chemicals, or any other type  
18.21 of material or as a result of any mental or physical condition, including deterioration through  
18.22 the aging process or loss of motor skills;

18.23 (15) for a pharmacist, pharmacy, pharmacist intern, pharmacy technician, medical gas  
18.24 dispenser, or controlled substance researcher, revealing a privileged communication from  
18.25 or relating to a patient except when otherwise required or permitted by law;

18.26 (16) for a pharmacist or pharmacy, improper management of patient records, including  
18.27 failure to maintain adequate patient records, to comply with a patient's request made pursuant  
18.28 to sections 144.291 to 144.298, or to furnish a patient record or report required by law;

18.29 (17) fee splitting, including without limitation:

18.30 (i) paying, offering to pay, receiving, or agreeing to receive, a commission, rebate,  
18.31 kickback, or other form of remuneration, directly or indirectly, for the referral of patients;

18.32 (ii) referring a patient to any health care provider as defined in sections 144.291 to  
18.33 144.298 in which the licensee or registrant has a financial or economic interest as defined

19.1 in section 144.6521, subdivision 3, unless the licensee or registrant has disclosed the  
19.2 licensee's or registrant's financial or economic interest in accordance with section 144.6521;  
19.3 and

19.4 (iii) any arrangement through which a pharmacy, in which the prescribing practitioner  
19.5 does not have a significant ownership interest, fills a prescription drug order and the  
19.6 prescribing practitioner is involved in any manner, directly or indirectly, in setting the price  
19.7 for the filled prescription that is charged to the patient, the patient's insurer or pharmacy  
19.8 benefit manager, or other person paying for the prescription or, in the case of veterinary  
19.9 patients, the price for the filled prescription that is charged to the client or other person  
19.10 paying for the prescription, except that a veterinarian and a pharmacy may enter into such  
19.11 an arrangement provided that the client or other person paying for the prescription is notified,  
19.12 in writing and with each prescription dispensed, about the arrangement, unless such  
19.13 arrangement involves pharmacy services provided for livestock, poultry, and agricultural  
19.14 production systems, in which case client notification would not be required;

19.15 (18) engaging in abusive or fraudulent billing practices, including violations of the  
19.16 federal Medicare and Medicaid laws or state medical assistance laws or rules;

19.17 (19) engaging in conduct with a patient that is sexual or may reasonably be interpreted  
19.18 by the patient as sexual, or in any verbal behavior that is seductive or sexually demeaning  
19.19 to a patient;

19.20 (20) failure to make reports as required by section 151.072 or to cooperate with an  
19.21 investigation of the board as required by section 151.074;

19.22 (21) knowingly providing false or misleading information that is directly related to the  
19.23 care of a patient unless done for an accepted therapeutic purpose such as the dispensing and  
19.24 administration of a placebo;

19.25 (22) aiding suicide or aiding attempted suicide in violation of section 609.215 as  
19.26 established by any of the following:

19.27 (i) a copy of the record of criminal conviction or plea of guilty for a felony in violation  
19.28 of section 609.215, subdivision 1 or 2;

19.29 (ii) a copy of the record of a judgment of contempt of court for violating an injunction  
19.30 issued under section 609.215, subdivision 4;

19.31 (iii) a copy of the record of a judgment assessing damages under section 609.215,  
19.32 subdivision 5; or

20.1 (iv) a finding by the board that the person violated section 609.215, subdivision 1 or 2.  
20.2 The board must investigate any complaint of a violation of section 609.215, subdivision 1  
20.3 or 2;

20.4 (23) for a pharmacist, practice of pharmacy under a lapsed or nonrenewed license. For  
20.5 a pharmacist intern, pharmacy technician, or controlled substance researcher, performing  
20.6 duties permitted to such individuals by this chapter or the rules of the board under a lapsed  
20.7 or nonrenewed registration. For a facility required to be licensed under this chapter, operation  
20.8 of the facility under a lapsed or nonrenewed license or registration; ~~and~~

20.9 (24) for a pharmacist, pharmacist intern, or pharmacy technician, termination or discharge  
20.10 from the health professionals services program for reasons other than the satisfactory  
20.11 completion of the program; and

20.12 (25) for a manufacturer, a violation of section 62J.842 or section 62J.845.

20.13 Sec. 20. **APPROPRIATION.**

20.14 \$..... in fiscal year 2024 and \$..... in fiscal year 2025 are appropriated from the general  
20.15 fund to the Prescription Drug Affordability Board established under Minnesota Statutes,  
20.16 section 62J.87, for implementation of the Prescription Drug Affordability Act.