

**HOUSE . . . . . No. 1215**

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**The Commonwealth of Massachusetts**

PRESENTED BY:

***John J. Lawn, Jr.***

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*To the Honorable Senate and House of Representatives of the Commonwealth of Massachusetts in General Court assembled:*

The undersigned legislators and/or citizens respectfully petition for the adoption of the accompanying bill:

An Act relative to pharmacy benefit managers.

PETITION OF:

NAME:	DISTRICT/ADDRESS:	DATE ADDED:
<i>John J. Lawn, Jr.</i>	<i>10th Middlesex</i>	<i>1/18/2023</i>
<i>Steven Owens</i>	<i>29th Middlesex</i>	<i>1/31/2023</i>
<i>Michael J. Finn</i>	<i>6th Hampden</i>	<i>1/31/2023</i>
<i>Smitty Pignatelli</i>	<i>3rd Berkshire</i>	<i>1/31/2023</i>
<i>William J. Driscoll, Jr.</i>	<i>7th Norfolk</i>	<i>1/31/2023</i>
<i>Christopher Hendricks</i>	<i>11th Bristol</i>	<i>1/31/2023</i>
<i>Lindsay N. Sabadosa</i>	<i>1st Hampshire</i>	<i>2/2/2023</i>
<i>Vanna Howard</i>	<i>17th Middlesex</i>	<i>2/2/2023</i>
<i>Carmine Lawrence Gentile</i>	<i>13th Middlesex</i>	<i>4/4/2023</i>
<i>Carole A. Fiola</i>	<i>6th Bristol</i>	<i>6/27/2023</i>
<i>Tommy Vitolo</i>	<i>15th Norfolk</i>	<i>6/27/2023</i>
<i>Mindy Domb</i>	<i>3rd Hampshire</i>	<i>10/20/2023</i>

**HOUSE . . . . . No. 1215**

By Representative Lawn of Watertown, a petition (accompanied by bill, House, No. 1215) of John J. Lawn, Jr., and others relative to pharmacy benefit managers. Health Care Financing.

**The Commonwealth of Massachusetts**

**In the One Hundred and Ninety-Third General Court  
(2023-2024)**

An Act relative to pharmacy benefit managers.

*Be it enacted by the Senate and House of Representatives in General Court assembled, and by the authority of the same, as follows:*

1 SECTION 1. Said section 1 of said chapter 6D , as so appearing, is hereby further  
2 amended by inserting after the definition of “Performance penalty” the following 2 definitions:-

3 “Pharmacy benefit manager”, as defined in section 1 of chapter 176X

4 “Pharmacy benefit services”, as defined in section 1 of chapter 176X

5 SECTION 2. Section 4 of said chapter 6D , as so appearing, is hereby amended by  
6 striking out, in lines 6 and 7, the word “manufacturers” and inserting in place thereof the  
7 following words:- pharmacy benefit managers.

8 SECTION 3. Section 6 of said chapter 6D , as so appearing, is hereby amended by adding  
9 the following paragraph:-

10 If the analysis of spending trends with respect to the pharmaceutical or biopharmaceutical  
11 products increases the expenses of the commission, the estimated increases in the commission’s

12 expenses shall be assessed fully to pharmacy benefit managers in the same manner as the  
13 assessment pursuant to section 68 of chapter 118E. A pharmacy benefit manager that is a  
14 surcharge payor subject to the preceding paragraph and administers its own prescription drug,  
15 prescription device or pharmacist services or prescription drug and device and pharmacist  
16 services portion shall not be subject to additional assessment under this paragraph.

17 SECTION 4. Section 8 of said chapter 6D , as so appearing, is hereby amended by  
18 inserting after the word “organization” , in lines 6 and 7, the following words:- , pharmacy  
19 benefit manager.

20 SECTION 5. Said section 8 of said chapter 6D , as so appearing, is hereby further  
21 amended by inserting after the word “organizations”, in line 14, the following words:- ,  
22 pharmacy benefit managers.

23 SECTION 6. Said section 8 of said chapter 6D , as so appearing, is hereby further  
24 amended by striking out, in lines 32 and 33 , the words “and (xi) any witness identified by the  
25 attorney general or the center” and inserting in place thereof the following words:- (xi) 2  
26 pharmacy benefit managers; and (xii) any witness identified by the attorney general or the center.

27 SECTION 7. Subsection (g) of said section 8 of said chapter 6D , as so appearing, is  
28 hereby further amended by striking out the second sentence and inserting in place thereof the  
29 following sentence:- The report shall be based on the commission's analysis of information  
30 provided at the hearings by witnesses, providers, provider organizations, insurers and pharmacy  
31 benefit managers, registration data collected pursuant to section 11, data collected or analyzed by  
32 the center pursuant to sections 8, 9, 10,10A and 10B of chapter 12C and any other available

33 information that the commission considers necessary to fulfill its duties in this section, as defined  
34 in regulations promulgated by the commission.

35 SECTION 8. Section 9 of said chapter 6D , as so appearing, is hereby amended by  
36 inserting after the word “organization”, in line 72, the following words:- , pharmacy benefit  
37 manager.

38 SECTION 9. Said Section 9 of said chapter 6D , as so appearing, is hereby further  
39 amended by inserting after the word “organizations”, in line 82, the following words:- ,  
40 pharmacy benefit manager.

41 SECTION 10. Section 1 of chapter 12C of the General Laws, as appearing in the 2018  
42 Official Edition, is hereby amended by inserting after the definition of “Patient-centered medical  
43 home” the following 5 definitions:-

44 “Pharmaceutical manufacturing company”, any entity engaged in the production,  
45 preparation, propagation, compounding, conversion or processing of prescription drugs, either  
46 directly or indirectly, by extraction from substances of natural origin, or independently by means  
47 of chemical synthesis or by a combination of extraction and chemical synthesis, or any entity  
48 engaged in the packaging, repackaging, labeling, relabeling or distribution of prescription drugs;  
49 provided however, that “pharmaceutical manufacturing company” shall not include a wholesale  
50 drug distributor licensed pursuant to section 36A of chapter 112 or a retail pharmacist registered  
51 pursuant to section 38 of said chapter 112.

52 “Pharmacy benefit manager”, any person, business, or entity, however organized, that  
53 administers, either directly or through its subsidiaries, pharmacy benefit services for prescription

54 drugs and devices on behalf of health benefit plan sponsors, including, but not limited to, self-  
55 insured employers, insurance companies and labor unions;

56 “Pharmacy benefit services” shall include, but not be limited to, formulary  
57 administration; drug benefit design; pharmacy network contracting; pharmacy claims processing;  
58 mail and specialty drug pharmacy services; and cost containment, clinical, safety, and adherence  
59 programs for pharmacy services. For the purposes of this section, a health benefit plan that does  
60 not contract with a pharmacy benefit manager shall be a pharmacy benefit manager, unless  
61 specifically exempted.

62 “Wholesale acquisition cost”, the cost of a prescription drug as defined in 42 U.S.C.  
63 §1395w-3a(c)(6)(B).

64 SECTION 11. Section 3 of said chapter 12C , as so appearing, is hereby amended by  
65 inserting after the word “organizations”, in lines 13 and 14, the following words:- , pharmacy  
66 benefit managers.

67 SECTION 12. Section 5 of said chapter 12C , as so appearing, is hereby amended by  
68 inserting after the word “organizations”, in line 11, the following words:- , pharmacy benefit  
69 managers.

70 SECTION 13. Said section 5 of said chapter 12C , as so appearing, is hereby further  
71 amended by inserting after the word “providers”, in line 15, the following words:- , affected  
72 pharmacy benefit managers.

73 SECTION 14. Section 7 of said chapter 12C , as so appearing, is hereby further amended  
74 by adding the following paragraph:-

75 To the extent that the analysis and reporting activities pursuant to section 10A increases  
76 the expenses of the center, the estimated increase in the center's expenses shall be fully assessed  
77 to pharmacy benefit managers in the same manner as the assessment pursuant to section 68 of  
78 chapter 118E.

79 SECTION 15. Said chapter 12C is hereby further amended by inserting after section 10  
80 the following section:-

81 Section 10A . The center shall promulgate regulations necessary to ensure the uniform  
82 analysis of information regarding pharmacy benefit managers that enables the center to analyze:  
83 (1) year-over-year wholesale acquisition cost changes; (2) year-over-year trends in formulary,  
84 maximum allowable costs list and cost-sharing design, including the establishment and  
85 management of specialty product lists; (3) aggregate information regarding discounts,  
86 utilizations limits, rebates, manufacturer administrative fees and other financial incentives or  
87 concessions related to pharmaceutical products or formulary programs; (4) information regarding  
88 the aggregate amount of payments made to pharmacies owned or controlled by the pharmacy  
89 benefit managers and the aggregate amount of payments made to pharmacies that are not owned  
90 or controlled by the pharmacy benefit managers; and (5) additional information deemed  
91 reasonable and necessary by the center as set forth in the center's regulations.

92 SECTION 16. Section 11 of said chapter 12C , as so appearing, is hereby amended by  
93 striking out the first sentence and inserting in place thereof the following sentence:-

94 The center shall ensure the timely reporting of information required pursuant to sections  
95 8, 9, 10 and 10A.

96 SECTION 17. Said section 11 of said chapter 12C , as so appearing, is hereby further  
97 amended by striking out, in line 11, the figure “\$1,000” and inserting in place thereof the  
98 following figure:- \$5,000.

99 SECTION 18. Said section 11 of said chapter 12C , as so appearing, is hereby further  
100 amended by striking out, in line 16, the figure “\$50,000” and inserting in place thereof the  
101 following figure:- \$200,000.

102 SECTION 19. Section 12 of said chapter 12C, as so appearing, is hereby amended by  
103 striking out, in line 2, the words “9 and 10” and inserting in place thereof the following words:-  
104 9, 10 and 10A

105 SECTION 20. Subsection (a) of section 16 of said chapter 12C , as so appearing, is  
106 hereby amended by striking out the first sentence and inserting in place thereof the following  
107 sentence:- The center shall publish an annual report based on the information submitted pursuant  
108 to sections 8, 9, 10, 10A and 10B concerning health care provider, provider organization,  
109 pharmacy benefit manager and private and public health care payer costs and cost and price  
110 trends, pursuant to section 13 of chapter 6D relative to market impact reviews and pursuant to  
111 section 15 relative to quality data.

112 SECTION 21. Chapter 94C is hereby further amended by inserting after section 21B the  
113 following section:-

114 Section 21C. (a) For the purposes of this section, the following words shall, unless the  
115 context clearly requires otherwise, have the following meanings:-

116 “Cost sharing”, amounts owed by a consumer under the terms of the consumer’s health  
117 benefit plan as defined in section 1 of chapter 176O or as required by a pharmacy benefit  
118 manager as defined in section 1 of chapter 6D.

119 “Pharmacy retail price”, the amount an individual would pay for a prescription  
120 medication at a pharmacy if the individual purchased that prescription medication at that  
121 pharmacy without using a health benefit plan as defined in section 1 of chapter 176O or any  
122 other prescription medication benefit or discount.

123 “Registered pharmacist”, a pharmacist who holds a valid certificate of registration issued  
124 by the board of registration in pharmacy pursuant to section 24 of chapter 112.

125 (b) A pharmacy shall post a notice informing consumers that a consumer may request, at  
126 the point of sale, the current pharmacy retail price for each prescription medication the consumer  
127 intends to purchase. If the consumer’s cost-sharing amount for a prescription medication exceeds  
128 the current pharmacy retail price, the pharmacist, or an authorized individual at the direction of a  
129 pharmacist, shall notify the consumer that the pharmacy retail price is less than the patient’s cost-  
130 sharing amount. The pharmacist shall charge the consumer the applicable cost-sharing amount or  
131 the current pharmacy retail price for that prescription medication, as directed by the consumer.

132 A pharmacist shall not be subject to a penalty by the board of registration in pharmacy or  
133 a third party for failure to comply with this section.

134 (c) A contractual obligation shall not prohibit a pharmacist from complying with this  
135 section; provided however, that a pharmacist shall submit a claim to the consumer’s health  
136 benefit plan or its pharmacy benefit manager if the pharmacist has knowledge that the  
137 prescription medication is covered under the consumer’s health benefit plan.



138 (d) Failure to post notice pursuant to subsection (b) shall be an unfair or deceptive act of  
139 practice under chapter 93A.

140 SECTION 22 Section 226 of chapter 175 is hereby repealed.

141 SECTION 23. The General Laws are hereby amended by inserting after Chapter 176W  
142 the following chapter:-

143 Chapter 176X

144 Section 1 . As used in this chapter, the following words shall have the following  
145 meanings, unless the context clearly requires otherwise:-

146 “Carrier”, as defined in section 1 of chapter 176O “Commissioner”, the commissioner of  
147 the division of insurance.

148 “Cost-sharing requirement”, any copayment, coinsurance, deductible, or annual limitation  
149 on cost-sharing (including a limitation subject to 42 U.S.C. §§ 18022(c) and 300gg-6(b)),  
150 required by or on behalf of an insured in order to receive specific health care services, including  
151 a prescription drug, covered by a health benefit plan .

152 “Division”, the division of insurance.

153 “Health benefit plan”, as defined in section 1 of chapter 176O

154 “Health care services”, supplies, care and services of a medical, surgical, optometric,  
155 dental, podiatric, chiropractic, psychiatric, therapeutic, diagnostic, preventative, rehabilitative,  
156 supportive, or geriatric nature including, but not limited to, inpatient and outpatient acute  
157 hospital care and services, services provided by a community health center or by a sanatorium, as

158 included in the definition of “hospital” in Title XVIII of the federal Social Security Act, and  
159 treatment and care compatible with such services or by a health maintenance organization.

160 “Insured”, an enrollee, covered person, insured, member, policyholder or subscriber of a  
161 carrier, including an individual whose eligibility as an insured of a carrier is in dispute or under  
162 review, or any other individual whose care may be subject to review by a utilization review  
163 program or entity as described under other provisions of this chapter.

164 “Mail order pharmacy”, a pharmacy whose primary business is to receive prescriptions  
165 by mail, telefax or through electronic submissions and to dispense medication to insureds  
166 through the use of the United States mail or other common or contract carrier services and that  
167 provides any consultation with patients electronically rather than face to face. “Network”, as  
168 defined in section 1 of chapter 176O.

169 “Network pharmacy”, a retail or other licensed pharmacy provider that contracts with a  
170 pharmacy benefit manager.

171 “Person”, a natural person, corporation, mutual company, unincorporated association,  
172 partnership, joint venture, limited liability company, trust, estate, foundation, not-for-profit  
173 corporation, unincorporated organization, government or governmental subdivision or agency.

174 “Pharmacy”, a facility, either physical or electronic, under the direction or supervision of  
175 a registered pharmacist which is authorized to dispense prescription drugs and has entered into a  
176 network contract with a pharmacy benefit manager or a carrier.

177 “Pharmacy benefit manager”, a person, business, or other entity that, pursuant to a  
178 contract or under an employment relationship with a carrier, a self-insurance plan, or other third-

179 party payer, either directly or through an intermediary, manages the prescription drug coverage  
180 provided by the carrier, self-insurance plan, or other third-party payer including, but not limited  
181 to, the processing and payment of claims for prescription drugs, the performance of drug  
182 utilization review, the processing of drug prior authorization requests, the adjudication of appeals  
183 or grievances related to prescription drug coverage, contracting with network pharmacies, and  
184 controlling the cost of covered prescription drugs.

185 “Pharmacy benefit services” shall include, but not be limited to, formulary  
186 administration; drug benefit design; pharmacy network contracting; pharmacy claims processing;  
187 mail and specialty drug pharmacy services; and cost containment, clinical, safety, adherence  
188 programs for pharmacy services, and any other pharmacy benefit service that the commissioner  
189 deems appropriate. For the purposes of the chapter, a health benefit plan that does not contract  
190 with a pharmacy benefit manager shall be a pharmacy benefit manager.

191 “Rebates or fees”, all fees or price concessions paid by a manufacturer to a pharmacy  
192 benefit manager or carrier, including rebates, discounts, and other price concessions that are  
193 based on actual or estimated utilization of a prescription drug. Rebates also include price  
194 concessions based on the effectiveness a drug as in a value-based or performance-based contract.

195 “Retail pharmacy”, as defined in section 39D of chapter 112.

196 "Spread pricing" means the practice of a pharmacy benefit manager retaining an  
197 additional amount of money in addition to the amount paid to the pharmacy to fill a prescription.

198 "Steering", a practice employed by a pharmacy benefit manager or carrier that channels a  
199 prescription to a pharmacy in which a pharmacy benefit manager or carrier has an ownership  
200 interest, and includes but is not limited to retail, mail-order, or specialty pharmacies.

201           Section 2. (a) Any pharmacy benefit manager contracting with a pharmacy that operates  
202 in the commonwealth shall comply with the provisions of this chapter.

203           (b) A pharmacy benefit manager shall receive a license from the division before  
204 conducting business in the commonwealth. A license granted pursuant to this section is not  
205 transferable.

206           (c) A license may be granted only when the division is satisfied that the entity possesses  
207 the necessary organization, background expertise, and financial integrity to supply the services  
208 sought to be offered.

209           (d) The division may issue a license subject to restrictions or limitations upon the  
210 authorization, including the type of services that may be supplied or the activities in which the  
211 entity may be engaged.

212           (e) A license shall be valid for a period of three years. The commissioner shall charge  
213 application and renewal fees in the amount of \$25,000

214           (f) The division shall develop an application for licensure that includes at least the  
215 following information: (i) the name of the pharmacy benefit manager; (ii) the address and contact  
216 telephone number for the pharmacy benefit manager; (iii) the name and address of the pharmacy  
217 benefit manager's agent for service of process in the commonwealth; (iv) the name and address  
218 of each person beneficially interested in the pharmacy benefit manager; and (v) the name and  
219 address of each person with management or control over the pharmacy benefit manager.

220           (g) The division may suspend, revoke, or place on probation a pharmacy benefit manager  
221 license under any of the following circumstances: (i) the pharmacy benefit manager has engaged

222 in fraudulent activity that constitutes a violation of state or federal law; (ii) the division received  
223 consumer complaints that justify an action under this chapter to protect the safety and interests of  
224 consumers; (iii) the pharmacy benefit manager fails to pay an application fee for the license; or  
225 (iv) the pharmacy benefit manager fails to comply with a requirement set forth in this chapter.

226 (h) If an entity performs the functions of pharmacy benefit manager acts without  
227 registering, it will be subject to a fine of \$5,000 per day for the period they are found to be in  
228 violation.

### 229 Section 3

230 (a) (i) The pharmacy benefit manager shall have a duty and obligation to perform  
231 pharmacy benefit services with care, skill, prudence, diligence, and professionalism.

232 (ii) In addition to the duties as may be prescribed by regulation:

233 (1) A pharmacy benefit manager interacting with a covered individual shall have the  
234 same duty to a covered individual as the health plan for whom it is performing pharmacy benefit  
235 services.

236 (2) A pharmacy benefit manager shall have a duty of good faith and fair dealing with all  
237 parties, including but not limited to covered individuals and pharmacies, with whom it interacts  
238 in the performance of pharmacy benefit services.

### 239 Section 4

240 (a) A pharmacy benefit manager shall provide a reasonably adequate and accessible  
241 pharmacy benefit manager network for the provision of prescription drugs, which provides for  
242 convenient patient access to pharmacies within a reasonable distance from a patient's residence.

243 (b) A pharmacy benefit manager may not deny a pharmacy the opportunity to participate  
244 in a pharmacy benefit manager network at preferred participation status if the pharmacy is  
245 willing to accept the terms and conditions that the pharmacy benefit manager has established for  
246 other pharmacies as a condition of preferred network participation status.

247 (c) A mail-order pharmacy shall not be included in the calculations for determining  
248 pharmacy benefit manager network adequacy under this section.

249 Section 5.

250 (a) After the date of receipt of a clean claim for payment made by a pharmacy, a  
251 pharmacy benefit manager shall not retroactively reduce payment on the claim, either directly or  
252 indirectly, through aggregated effective rate, direct or indirect remuneration, quality assurance  
253 program or otherwise, except if the claim is found not to be a clean claim during the course of a  
254 routine audit performed pursuant to an agreement between the pharmacy benefit manager and the  
255 pharmacy. When a pharmacy adjudicates a claim at the point of sale, the reimbursement amount  
256 provided to the pharmacy by the pharmacy benefit manager shall constitute a final  
257 reimbursement amount. Nothing in this section shall be construed to prohibit any retroactive  
258 increase in payment to a pharmacy pursuant to a contract between the pharmacy benefit manager  
259 or a pharmacy.

260 (b) For the purpose of this section, "clean claim" means a claim that has no defect or  
261 impropriety, including a lack of any required substantiating documentation, or other  
262 circumstance requiring special treatment, including, but not limited to, those listed in subsection  
263 (d) of this section, that prevents timely payment from being made on the claim.

264 (c) A pharmacy benefit manager shall not recoup funds from a pharmacy in connection  
265 with claims for which the pharmacy has already been paid unless the recoupment is:

266 (1) otherwise permitted or required by law; or

267 (2) the result of an audit, performed pursuant to a contract between the pharmacy benefit  
268 manager and the pharmacy; or

269 (d) The provisions of this section shall not apply to an investigative audit of pharmacy  
270 records when:

271 (1) fraud, waste, abuse or other intentional misconduct is indicated by physical review or  
272 review of claims data or statements; or

273 (2) other investigative methods indicate a pharmacy is or has been engaged in criminal  
274 wrongdoing, fraud or other intentional or willful misrepresentation.

275 (e) No pharmacy benefit manager shall charge or collect from an individual a copayment  
276 that exceeds the total submitted charges by the pharmacy for which the pharmacy is paid. If an  
277 individual pays a copayment, the pharmacy shall retain the adjudicated costs and the pharmacy  
278 benefit manager shall not redact or recoup the adjudicated cost.

279 Section 6.

280 (a) As used in this section:

281 (1) “Generically equivalent drug”, a drug that is pharmaceutically and therapeutically  
282 equivalent to the drug prescribed;

283 (2)(A) “Maximum allowable cost list”, a listing of drugs or other methodology used by a  
284 pharmacy benefit manager, directly or indirectly, setting the maximum allowable payment to a  
285 pharmacy or pharmacist for a generic drug, brand-name drug, biologic product, or other  
286 prescription drug.

287 (B) Maximum allowable cost list includes without limitation:

288 (i) Average acquisition cost, including national average drug acquisition cost;

289 (ii) Average manufacturer price;

290 (iii) Average wholesale price;

291 (iv) Brand effective rate or generic effective rate;

292 (v) Discount indexing;

293 (vi) Federal upper limits;

294 (vii) Wholesale acquisition cost; and

295 (viii) Any other term that a pharmacy benefit manager or a carrier may use to establish  
296 reimbursement rates to a pharmacist or pharmacy for pharmacist services;

297 (3) “Pharmaceutical wholesaler”, as defined in section 36A of chapter 112;

298 (4) “Pharmacist”, a pharmacist who, pursuant to the provisions of M.G.L. c. 112, § 24, is  
299 registered by the Board to practice pharmacy;



300 (5) “Pharmacist services”, products, goods, and services, or any combination of products,  
301 goods, and services, provided as a part of the practice of pharmacy as defined in section 39D of  
302 chapter 112;

303 (6) “Pharmacy”, shall have the same meaning as defined in section 39D of chapter 112;

304 (7) “Pharmacy acquisition cost” means the amount that a pharmaceutical wholesaler  
305 charges for a pharmaceutical product as listed on the pharmacy's billing invoice;

306 (8) “Pharmacy benefit manager”, as defined in section 1 of chapter 176X;

307 (9) “Pharmacy benefit manager affiliate”, a pharmacy or pharmacist that directly or  
308 indirectly, through one (1) or more intermediaries, owns or controls, is owned or controlled by,  
309 or is under common ownership or control with a pharmacy benefits manager; and

310 (10) “Pharmacy benefit plan or program”, a plan or program that pays for, reimburses,  
311 covers the cost of, or otherwise provides for pharmacist services to individuals who reside in or  
312 are employed in the commonwealth.

313 (b) Before a pharmacy benefit manager places or continues a particular drug on a  
314 maximum allowable cost list, the drug:

315 (1) If the drug is a generically equivalent drug, it shall be listed as therapeutically  
316 equivalent and pharmaceutically equivalent A or B rated in the United States Food and Drug  
317 Administration's most recent version of the Orange Book or Green Book or have an NR or NA  
318 rating by Medi-Span, Gold Standard, or a similar rating by a nationally recognized reference;

319 (2) Shall be available for purchase by each pharmacy in the state from national or  
320 regional wholesalers operating in the commonwealth; and

321 (3) Shall not be obsolete.

322 (c ) A pharmacy benefit manager shall:

323 (1) Provide access to its maximum allowable cost list to each pharmacy subject to the  
324 maximum allowable cost list;

325 (2) Update its maximum allowable cost list on a timely basis, but in no event longer than  
326 seven (7) calendar days from an increase of ten per cent or more in the pharmacy acquisition cost  
327 from sixty per cent or more of the pharmaceutical wholesalers doing business in the state or a  
328 change in the methodology on which the maximum allowable cost list is based or in the value of  
329 a variable involved in the methodology;

330 (3) Provide a process for each pharmacy subject to the maximum allowable cost list to  
331 receive prompt notification of an update to the maximum allowable cost list; and

332 (4)(A)(i) Provide a reasonable administrative appeal procedure to allow pharmacies to  
333 challenge maximum allowable cost list and reimbursements made under a maximum allowable  
334 cost list for a specific drug or drugs as:

335 (a) Not meeting the requirements of this section; or

336 (b) Being below the pharmacy acquisition cost.

337 (ii) The reasonable administrative appeal procedure shall include the following:

338 (a) A dedicated telephone number, email address, and website for the purpose of  
339 submitting administrative appeals;

340 (b) The ability to submit an administrative appeal directly to the pharmacy benefit  
341 manager regarding the pharmacy benefits plan or program or through a pharmacy service  
342 administrative organization; and

343 (c) No less than thirty business days to file an administrative appeal.

344 (B) The pharmacy benefit manager shall respond to the challenge under subdivision  
345 (c)(4)(A) of this section within thirty business days after receipt of the challenge.

346 (C) If a challenge is made under subdivision (c)(4)(A) of this section, the pharmacy  
347 benefit manager shall within thirty business days after receipt of the challenge either:

348 (i) If the appeal is upheld:

349 (a) Make the change in the maximum allowable cost list payment to at least the pharmacy  
350 acquisition cost;

351 (b) Permit the challenging pharmacy or pharmacist to reverse and rebill the claim in  
352 question;

353 (c) Provide the National Drug Code that the increase or change is based on to the  
354 pharmacy or pharmacist; and

355 (d) Make the change under subdivision (c)(4)(C)(i)(a) of this section effective for each  
356 similarly situated pharmacy as defined by the payor subject to the maximum allowable cost list;

357 (ii) If the appeal is denied, provide the challenging pharmacy or pharmacist the National  
358 Drug Code and the name of the national or regional pharmaceutical wholesalers operating in the

359 commonwealth that have the drug currently in stock at a price below the maximum allowable  
360 cost as listed on the maximum allowable cost list; or

361 (iii) If the National Drug Code provided by the pharmacy benefit manager is not available  
362 below the pharmacy acquisition cost from the pharmaceutical wholesaler from whom the  
363 pharmacy or pharmacist purchases the majority of prescription drugs for resale, then the  
364 pharmacy benefit manager shall adjust the maximum allowable cost as listed on the maximum  
365 allowable cost list above the challenging pharmacy's pharmacy acquisition cost and permit the  
366 pharmacy to reverse and rebill each claim affected by the inability to procure the drug at a cost  
367 that is equal to or less than the previously challenged maximum allowable cost.

368 (d)(1) A pharmacy benefit manager shall not reimburse a pharmacy or pharmacist in the  
369 commonwealth an amount less than the amount that the pharmacy benefit manager reimburses a  
370 pharmacy benefit manager affiliate for providing the same pharmacist services.

371 (2) The amount shall be calculated on a per unit basis based on the same generic product  
372 identifier or generic code number.

373 (e) A pharmacy or pharmacist may decline to provide the pharmacist services to a patient  
374 or pharmacy benefit manager if, as a result of a maximum allowable cost list, a pharmacy or  
375 pharmacist is to be paid less than the pharmacy acquisition cost of the pharmacy providing  
376 pharmacist services.

377 (f) This section does not apply to a maximum allowable cost list maintained by  
378 MassHealth or the division of insurance.

379 (g)(1)A violation of this section shall constitute an unfair or deceptive act or practice  
380 pursuant to chapter 93A.

381 Section 7.

382 (a) No pharmacy benefit manager or representative of a pharmacy benefit manager shall  
383 conduct spread pricing in the commonwealth.

384 (b) A pharmacy benefit manager or representative of a pharmacy benefit manager that  
385 violates this section shall be subject to the surcharge under section 8 of chapter 176X.

386 (c) A pharmacy benefit manager shall report to the commissioner on a quarterly basis for  
387 each healthcare insurer the following information:

388 (A) The aggregate number of rebates received by the pharmacy benefit manager;

389 (B) The aggregate number of rebates distributed to the appropriate healthcare insurer;

390 (C) The aggregate number of rebates passed on to an insured of each healthcare insurer at  
391 the point of sale that reduced the insured's applicable deductible, copayment, coinsurance, or  
392 other cost-sharing amount;

393 (D) The individual and aggregate amount paid by the healthcare insurer to the pharmacy  
394 benefit manager for pharmacist services itemized by pharmacy, by product, and by goods and  
395 services; and

396 (E) The individual and aggregate amount a pharmacy benefit manager paid for  
397 pharmacist services itemized by pharmacy, by product, and by goods and services.

398 (d) The commissioner, in consultation with the health policy commission and the center  
399 for health information and analysis, shall annually report on the rebates and amounts reported  
400 under subsection (c), which shall be public record.

401 Section 8.

402 (a) A pharmacy benefits manager that engages in the practices of (i) spread pricing; (ii)  
403 steering; or (iii) imposing point-of-sale fees or retroactive fees shall be subject to a surcharge  
404 payable to the division of 10 percent on the aggregate dollar amount it reimbursed pharmacies in  
405 the previous calendar year for prescription drugs in the commonwealth.

406 (b) By March 1 of each year, a pharmacy benefit manager shall provide a letter to the  
407 commissioner attesting as to whether or not, in the previous calendar year, it engaged in the any  
408 of the practices under subsection (a). The pharmacy benefit manager shall also submit to the  
409 commissioner, in a form and manner and by a date specified by the commissioner, data detailing  
410 all prescription drug claims it administered in the commonwealth for insured residents on behalf  
411 of each health plan client and any other data the commissioner deems necessary to evaluate  
412 whether a pharmacy benefit manager may be engaged in any of the practices under subsection  
413 (a)

414 (c) By April 1 of each year, a pharmacy benefit manager shall pay into the general fund  
415 the surcharge owed, if any, as contained in the report submitted pursuant to subsection (b) of this  
416 section.

417 (d) Nothing in this section shall be construed to authorize the practices of steering or  
418 imposing point-of-sale fees or retroactive fees where otherwise prohibited by law.

419 (e) The commissioner, in consultation with the health policy commission and the center  
420 for health information and analysis, shall prepare an aggregate report reflecting the total number  
421 of prescriptions administered by the reporting pharmacy benefit manager with the total sum due  
422 to the commonwealth, which shall be public record.

423 Section 9.

424 (a) Any person operating a health plan whose contracted pharmacy benefits manager  
425 engages in the practices of (i) spread pricing; (ii) steering; or (iii) imposing point-of-sale fees or  
426 retroactive fees in connection with its health plans shall be subject to a surcharge payable to the  
427 division of 10 percent on the aggregate dollar amount its pharmacy benefit manager reimbursed  
428 pharmacies on its behalf in the previous calendar year for prescription drugs in the  
429 commonwealth.

430 (b) By March 1 of each year, any person operating a health plan and licensed in the  
431 commonwealth that utilizes a contracted pharmacy benefit manager shall provide a letter to the  
432 commissioner attesting as to whether or not, in the previous calendar year, its contracted  
433 pharmacy benefit manager engaged in any of the practices under subsection (a) in connection  
434 with its health plans. The health plan shall also submit to the commissioner, in a form and  
435 manner and by a date specified by the commissioner, data detailing all prescription drug claims  
436 its contracted pharmacy benefit manager administered in the commonwealth for insured  
437 residents and any other data the commissioner deems necessary to evaluate whether a health  
438 plan's pharmacy benefit manager may be engaged in any of the practices under subsection (a).

439 (c) By April 1 of each year, any person operating a health plan and licensed under this  
440 title shall pay into the general fund the surcharge owed, if any, as contained in the report  
441 submitted pursuant to subsection (b) of this section.

442 (d) Nothing in this section shall be construed to authorize the practices of steering or  
443 imposing point-of-sale fees or retroactive fees where otherwise prohibited by law.

444 (e) The commissioner, in consultation with the health policy commission and the center  
445 for health information and analysis, shall prepare an aggregate report reflecting the total number  
446 of prescriptions administered by the reporting health plan along with the total sum due to the  
447 commonwealth, which shall be public record.

448 Section 10.

449 When calculating an insured's contribution to any applicable cost sharing requirement, a  
450 pharmacy benefit manager shall include any cost-sharing amounts paid by the insured or on  
451 behalf of the insured by another person.

452 Section 11.

453 (a) A pharmacy benefit manager shall conduct an audit of the records of a pharmacy in  
454 accordance with paragraphs (1) to (13), inclusive.

455 (1) The contract between a pharmacy and a pharmacy benefit manager shall identify and  
456 describe the audit procedures in detail.

457 (2) With the exception of an investigative fraud audit, the auditor shall give the pharmacy  
458 written notice at least 2 weeks prior to conducting the initial on-site audit for each audit cycle.



459 (3) A pharmacy benefit manager shall not audit claims beyond 2 years prior to the date of  
460 audit.

461 (4) The auditor shall not interfere with the delivery of pharmacist services to a patient and  
462 shall make a reasonable effort to minimize the inconvenience and disruption to the pharmacy  
463 operations during the audit process.

464 (5) Any audit that involves clinical or professional judgment shall be conducted by, or in  
465 consultation with, a licensed pharmacist from any state.

466 (6) A finding of an overpayment or underpayment shall be based on the actual  
467 overpayment or underpayment. A statistically sound calculation for overpayment or  
468 underpayment may be used to determine recoupment as part of a settlement as agreed to by the  
469 pharmacy.

470 (7) The auditor shall audit each pharmacy under the same standards and parameters with  
471 which they audit other similarly situated pharmacies.

472 (8) An audit shall not be initiated or scheduled during the first 5 calendar days of any  
473 month for any pharmacy that averages more than 600 prescriptions per week without the  
474 pharmacy's consent.

475 (9) A preliminary audit report shall be delivered to the pharmacy not later than 30 days  
476 after the conclusion of the audit.

477 (10) The preliminary audit report shall be signed and shall include the signature of any  
478 pharmacist participating in the audit.

479 (11) A pharmacy benefit manager shall not withhold payment to a pharmacy for  
480 reimbursement claims as a means to recoup money until after the final internal disposition of an  
481 audit, including the appeals process, as provided in subsection (b), unless fraud or  
482 misrepresentation is reasonably suspected or the discrepant amount exceeds \$15,000.

483 (12) The auditor shall provide a copy of the final audit report to the pharmacy and plan  
484 sponsor within 30 days following the pharmacy's receipt of the signed preliminary audit report or  
485 the completion of the appeals process, as provided in subsection (b), whichever is later.

486 (13) No auditing company or agent shall receive payment based upon a percentage of the  
487 amount recovered or other financial incentive tied to the findings of the audit.

488 (b)(1) Each auditor shall establish an appeals process under which a pharmacy may  
489 appeal findings in a preliminary audit.

490 (2) To appeal a finding, a pharmacy may use the records of a hospital, physician, or other  
491 authorized prescriber to validate the record with respect to orders or refills of prescription drugs  
492 or devices.

493 (3) A pharmacy shall have 30 days to appeal any discrepancy found during the  
494 preliminary audit.

495 (4) The National Council for Prescription Drug Programs or any other recognized  
496 national industry standard shall be used to evaluate claims submission and product size disputes.

497 (5) If an audit results in the identification of any clerical or record-keeping errors in a  
498 required document or record, the pharmacy shall not be subject to recoupment of funds by the  
499 pharmacy benefit manager; provided, that the pharmacy may provide proof that the patient

500 received the medication billed to the plan via patient signature logs or other acceptable methods,  
501 unless there is financial harm to the plan or errors that exceed the normal course of business.

502 (c) This section shall not apply to any audit or investigation of a pharmacy that involves  
503 potential fraud, willful misrepresentation or abuse, including, but not limited to, investigative  
504 audits or any other statutory or regulatory provision which authorizes investigations relating to  
505 insurance fraud.

506 (d) This section shall not apply to a public health care payer, as defined in section 1 of  
507 chapter 12C.

508 (e) The commissioner shall promulgate regulations to enforce this section.

509 Section 12.

510 (a) The commissioner may make an examination of the affairs of a Pharmacy Benefit  
511 Manager when the commissioner deems prudent but not less frequently than once every 3 years.  
512 The focus of the examination shall be to ensure that a pharmacy benefit manager is able to meet  
513 its responsibilities under contracts with licensed carriers. The examination shall be conducted  
514 according to the procedures set forth in subsection (6) of section 4 of chapter 175.

515 (b) The commissioner, a deputy or an examiner may conduct an on-site examination of  
516 each pharmacy benefit manager in the commonwealth to thoroughly inspect and examine its  
517 affairs.

518 (c) The charge for each such examination shall be determined annually according to the  
519 procedures set forth in subsection (6) of section 4 of chapter 175.

520 (d) Not later than 60 days following completion of the examination, the examiner in  
521 charge shall file with the commissioner a verified written report of examination under oath.  
522 Upon receipt of the verified report, the commissioner shall transmit the report to the pharmacy  
523 benefit manager examined with a notice which shall afford the pharmacy benefit manager  
524 examined a reasonable opportunity of not more than 30 days to make a written submission or  
525 rebuttal with respect to any matters contained in the examination report. Within 30 days of the  
526 end of the period allowed for the receipt of written submissions or rebuttals, the commissioner  
527 shall consider and review the reports together with any written submissions or rebuttals and any  
528 relevant portions of the examiner's work papers and enter an order:

529 (i) adopting the examination report as filed with modifications or corrections and, if the  
530 examination report reveals that the pharmacy benefit manager is operating in violation of this  
531 section or any regulation or prior order of the commissioner, the commissioner may order the  
532 pharmacy benefit manager to take any action the commissioner considered necessary and  
533 appropriate to cure such violation;

534 (ii) rejecting the examination report with directions to examiners to reopen the  
535 examination for the purposes of obtaining additional data, documentation or information and re-  
536 filing pursuant to the above provisions; or

537 (iii) calling for an investigatory hearing with no less than 20 days' notice to the pharmacy  
538 benefit manager for purposes of obtaining additional documentation, data, information and  
539 testimony.

540 (e) Notwithstanding any general or special law to the contrary, including clause 26 of  
541 section 7 of chapter 4 and chapter 66, the records of any such audit, examination or other

542 inspection and the information contained in the records, reports or books of any pharmacy  
543 benefit manager examined pursuant to this section shall be confidential and open only to the  
544 inspection of the commissioner, or the examiners and assistants. Access to such confidential  
545 material may be granted by the commissioner to law enforcement officials of the commonwealth  
546 or any other state or agency of the federal government at any time, so long as the agency or  
547 office receiving the information agrees in writing to keep such material confidential. Nothing  
548 herein shall be construed to prohibit the required production of such records, and information  
549 contained in the reports of such company or organization before any court of the commonwealth  
550 or any master or auditor appointed by any such court, in any criminal or civil proceeding,  
551 affecting such pharmacy benefit manager, its officers, partners, directors or employees. The final  
552 report of any such audit, examination or any other inspection by or on behalf of the division of  
553 insurance shall be a public record.

554 Section 13.

555 A pharmacy benefit manager shall be required to submit to periodic audits by a licensed  
556 carrier if the pharmacy benefit manager has entered into a contract with the carrier to provide  
557 pharmacy benefits to the carrier or its members. The commissioner shall direct or provide  
558 specifications for such audits

559 Section 14.

560 (a) A contract between a pharmacy benefit manager and a participating pharmacy or  
561 pharmacist or contracting agent shall not include any provision that prohibits, restricts, or limits a  
562 pharmacist or contracting agent or pharmacy's right to provide an insured with information on  
563 the amount of the insured's cost share for such insured's prescription drug and the clinical

564 efficacy of a more affordable alternative drug if one is available. Neither a pharmacy nor a  
565 pharmacist shall be penalized by a pharmacy benefit manager for disclosing such information to  
566 an insured or for selling to an insured a more affordable alternative if one is available.

567 (b) A pharmacy benefit manager shall not charge a pharmacist or pharmacy a fee related  
568 to the adjudication of a claim, including, without limitation, a fee for: (i) the receipt and  
569 processing of a pharmacy claim; (ii) the development or management of claims processing  
570 services in a pharmacy benefit manager network; or (iii) participation in a pharmacy benefit  
571 manager network, unless such fee is set out in a contract between the pharmacy benefit manager  
572 and the pharmacist or contracting agent or pharmacy.

573 (c) A contract between a pharmacy benefit manager and a participating pharmacy or  
574 pharmacist or contracting agent shall not include any provision that prohibits, restricts, or limits  
575 disclosure of information to the division deemed necessary by the division to ensure a pharmacy  
576 benefit manager's compliance with the requirements under this section or section 21C of chapter  
577 94C.

578 SECTION 24. Sections 1 to 22 shall take effect 6 months after the effective date of this  
579 act.

580 SECTION 25. The commissioner of insurance shall promulgate regulations to implement  
581 chapter 176X of the General Laws, as inserted by section 23, not later than 1 year after the  
582 effective date of this act.