

General Assembly

Substitute Bill No. 6447

January Session, 2021



AN ACT CREATING THE COVERED CONNECTICUT PROGRAM TO EXPAND ACCESS TO AFFORDABLE HEALTH CARE.

Be it enacted by the Senate and House of Representatives in General Assembly convened:

- 1 Section 1. Subsections (a) and (b) of section 19a-754a of the general
- 2 statutes are repealed and the following is substituted in lieu thereof
- 3 (Effective July 1, 2021):
- 4 (a) There is established an Office of Health Strategy, which shall be
- 5 within the Department of Public Health for administrative purposes
- 6 only. The department head of said office shall be the executive director
- 7 of the Office of Health Strategy, who shall be appointed by the Governor
- 8 in accordance with the provisions of sections 4-5 to 4-8, inclusive, with
- 9 the powers and duties therein prescribed.
- 10 (b) The Office of Health Strategy shall be responsible for the 11 following:
- 12 (1) Developing and implementing a comprehensive and cohesive
- 13 health care vision for the state, including, but not limited to, a
- 14 coordinated state health care cost containment strategy;
- 15 (2) Promoting effective health planning and the provision of quality
- 16 health care in the state in a manner that ensures access for all state

- residents to cost-effective health care services, avoids the duplication of such services and improves the availability and financial stability of such services throughout the state;
- 20 (3) Directing and overseeing the State Innovation Model Initiative 21 and related successor initiatives;
- 22 (4) (A) Coordinating the state's health information technology 23 initiatives, (B) seeking funding for and overseeing the planning, 24 implementation and development of policies and procedures for the 25 administration of the all-payer claims database program established 26 under section 19a-775a, (C) establishing and maintaining a consumer 27 health information Internet web site under section 19a-755b, and (D) 28 designating an unclassified individual from the office to perform the 29 duties of a health information technology officer as set forth in sections 30 17b-59f and 17b-59g;
- 31 (5) Directing and overseeing the Health Systems Planning Unit 32 established under section 19a-612 and all of its duties and 33 responsibilities as set forth in chapter 368z; [and]
- (6) Convening forums and meetings with state government and
 external stakeholders, including, but not limited to, the Connecticut
 Health Insurance Exchange, to discuss health care issues designed to
 develop effective health care cost and quality strategies;
- 38 <u>(7) Administering the Covered Connecticut account established</u> 39 <u>under section 2 of this act;</u>
- 40 (8) Annually determining the amount described in, and reporting 41 such amount to the Insurance Commissioner pursuant to, subsection (b) 42 of section 4 of this act;
- 43 (9) Annually (A) developing a plan, pursuant to subsection (b) of 44 section 3 of this act, in consultation with the Connecticut Health 45 Insurance Exchange, Commissioner of Social Services and Insurance 46 Commissioner, and (B) submitting a report, pursuant to subsection (c)

- of section 3 of this act and in accordance with section 11-4a, to the joint
- 48 standing committee of the General Assembly having cognizance of
- 49 <u>matters relating to insurance; and</u>
- 50 (10) Not later than February 1, 2023, and annually thereafter,
- 51 providing to the Commissioner of Revenue Services a list of the drugs
- 52 that the Secretary of Health and Human Services determined, pursuant
- to 21 USC 356e, as amended from time to time, were in shortage in the
- 54 United States during the preceding calendar year.
- Sec. 2. (NEW) (Effective July 1, 2021) There is established an account
- to be known as the "Covered Connecticut account" which shall be a
- 57 separate, nonlapsing account within the General Fund. The account
- 58 shall be administered by the Office of Health Strategy, established under
- section 19a-754a of the general statutes, as amended by this act, and
- 60 contain any moneys required by law to be deposited in the account.
- 61 Subject to the approval required under subsection (d) of section 3 of this
- 62 act, moneys in the account shall be expended by the Connecticut Health
- 63 Insurance Exchange, established pursuant to section 38a-1081 of the
- 64 general statutes, and the Department of Social Services in accordance
- with the plan developed by the Office of Health Strategy pursuant to
- 66 subsection (b) of section 3 of this act.
- 67 Sec. 3. (NEW) (Effective July 1, 2021) (a) For the purposes of this
- 68 section:
- 69 (1) "Affordable Care Act" has the same meaning as provided in
- 70 section 38a-1080 of the general statutes;
- 71 (2) "Covered Connecticut account" means the Covered Connecticut
- 72 account established under section 2 of this act;
- 73 (3) "Exchange" has the same meaning as provided in section 38a-1080
- of the general statutes; and
- 75 (4) "Office of Health Strategy" means the Office of Health Strategy
- 76 established under section 19a-754a of the general statutes, as amended

- 77 by this act.
- 78 (b) The Office of Health Strategy shall, in consultation with the
- 79 exchange, Commissioner of Social Services and Insurance
- 80 Commissioner, annually develop a plan to, within the funds available
- 81 in the Covered Connecticut account and without recourse to any other
- 82 state funds, reduce this state's uninsured rate by, among other things,
- 83 reducing the burden that health care costs impose on insureds. Such
- 84 plan may, among other things, call for:
- 85 (1) The exchange to:
- 86 (A) Establish:
- 87 (i) A state subsidy program to provide premium subsidies, at defined
- 88 amounts and to individuals within defined income brackets, for
- 89 individuals with incomes not greater than six hundred per cent of the
- 90 federal poverty level; or
- 91 (ii) A reinsurance program; or
- 92 (B) Seek, in consultation with the Office of Health Strategy, and, if
- 93 issued, implement, a state innovation waiver pursuant to Section 1332
- 94 of the Affordable Care Act; or
- 95 (2) The Commissioner of Social Services to expand medical assistance
- 96 under chapter 319v of the general statutes to provide coverage to
- 97 additional individuals.
- 98 (c) Not later than January 1, 2022, and annually thereafter, the Office
- 99 of Health Strategy shall submit a report, in accordance with section 11-
- 4a of the general statutes, to the joint standing committee of the General
- 101 Assembly having cognizance of matters relating to insurance. Such
- report shall contain the plan developed pursuant to subsection (b) of
- this section.
- 104 (d) Not later than February 1, 2022, and annually thereafter, the joint

- standing committee of the General Assembly having cognizance of
- matters relating to insurance shall advise the Office of Health Strategy,
- 107 exchange, Commissioner of Social Services and Insurance
- 108 Commissioner of its approval or rejection of the plan contained in the
- report submitted by the Office of Health Strategy pursuant to subsection
- 110 (c) of this section. If the committee does not act on or before said date,
- said plan shall be deemed rejected.
- Sec. 4. (NEW) (Effective July 1, 2021) (a) For the purposes of this
- 113 section:
- 114 (1) "Covered Connecticut account" means the Covered Connecticut
- account established under section 2 of this act;
- 116 (2) "Exempt insurer" means an insurer that administers self-insured
- 117 health benefit plans and is exempt from third-party administrator
- licensure under subparagraph (C) of subdivision (11) of section 38a-720
- of the general statutes and section 38a-720a of the general statutes; and
- 120 (3) "Office of Health Strategy" means the Office of Health Strategy
- established under section 19a-754a of the general statutes, as amended
- by this act.
- (b) (1) Not later than July 1, 2022, and annually thereafter, the Office
- of Health Strategy shall:
- (A) Determine the difference between fifty million dollars and the
- amount of moneys deposited that year in the Covered Connecticut
- account pursuant to subsection (k) of section 7 of this act; and
- (B) Report the amount determined pursuant to subparagraph (A) of
- this subdivision to the Insurance Commissioner.
- 130 (2) The Office of Health Strategy shall, not later than July 1, 2021,
- 131 report to the Insurance Commissioner that the amount described in
- subparagraph (A) of subdivision (1) of this subsection is thirty million
- dollars for the year 2022.

- (c) (1) Each insurer and health care center doing health insurance business in this state, and each exempt insurer, shall annually pay to the Insurance Commissioner, for deposit in the Covered Connecticut account, a fee assessed by the commissioner pursuant to this section.
- (2) Not later than July 1, 2021, and annually thereafter, each insurer, health care center and exempt insurer described in subdivision (1) of this subsection shall report to the commissioner, on a form designated by the commissioner, the number of insured or enrolled lives in this state as of the May first immediately preceding for which such insurer, health care center or exempt insurer was providing health insurance coverage, or administering a self-insured health benefit plan providing coverage, of the types specified in subdivisions (1), (2), (4), (11) and (12) of section 38a-469 of the general statutes. Such number shall not include insured or enrolled lives covered under fully-insured group health insurance policies sold in the small group market, Medicare, any medical assistance program administered by the Department of Social Services, workers' compensation insurance or Medicare Part C plans.
- (3) Not later than August 1, 2021, and annually thereafter, the commissioner shall determine the fee to be assessed for that year against each insurer, health care center and exempt insurer described in subdivision (1) of this subsection. Such fee shall be determined by multiplying the number of insured or enrolled lives reported to the commissioner pursuant to subdivision (2) of this subsection by a factor, determined annually by the commissioner, to fully fund the amount reported by the Office of Health Strategy to the commissioner pursuant to subsection (b) of this section. The commissioner shall determine the factor by dividing the amount reported by the Office of Health Strategy to the commissioner pursuant to subsection (b) of this section by the total number of insured or enrolled lives reported to the commissioner pursuant to subdivision (2) of this subsection.
- (4) (A) Not later than August 1, 2021, and annually thereafter, the commissioner shall submit a statement to each insurer, health care center and exempt insurer described in subdivision (1) of this subsection

- that includes the proposed fee imposed under this section for such insurer, health care center or exempt insurer determined in accordance with this subsection. Each such insurer, health care center and exempt insurer shall pay such fee to the commissioner not later than November first of that year.
- (B) Any insurer, health care center or exempt insurer described in subdivision (1) of this subsection that is aggrieved by an assessment levied under this subsection may appeal therefrom in the same manner as provided for appeals under section 38a-52 of the general statutes.
- (5) Any insurer, health care center or exempt insurer that fails to file the report required under subdivision (2) of this subsection, or pay the fee assessed under subdivision (1) of this subsection, shall pay a late filing or payment fee, as applicable, of one hundred dollars per day for each day from the date such report or payment was due. The commissioner shall deposit all late fees paid pursuant to this subdivision in the Covered Connecticut account. The commissioner may require an insurer, health care center or exempt insurer subject to this subsection to produce any records in its possession, and may require any other person to produce any records in such other person's possession, that were used to prepare such report for examination by the commissioner or the commissioner's designee. If the commissioner determines there exists anything other than a good faith discrepancy between the actual number of insured or enrolled lives that should have been reported to the commissioner pursuant to subdivision (2) of this subsection and the number actually reported, such insurer, health care center or exempt insurer shall be liable to this state for a civil penalty of not more than fifteen thousand dollars for each report filed for which the commissioner determines there is such a discrepancy.
- (6) (A) The commissioner shall apply any overpayment of the fee imposed under this section by an insurer, health care center or exempt insurer for a given year as a credit against the fee due from such insurer, health care center or exempt insurer under this section for the succeeding year if:

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200201	(i) The amount of the overpayment exceeds five thousand dollars; and		
202	(ii) On or before April first of the year following the overpayment, the		
203	insurer, health care center or exempt insurer:		
204	(I) Notifies the commissioner of the amount of the overpayment; and		
205	(II) Provides the commissioner with evidence sufficient to prove the		
206	amount of the overpayment.		
207	(B) Not later than ninety days after the commissioner receives the		
208	notice and supporting evidence under subparagraph (A)(ii) of this		
209	subdivision, the commissioner shall:		
210	(i) Determine whether the insurer, health care center or exempt		
211	insurer made an overpayment; and		
212	(ii) Notify the insurer, health care center or exempt insurer of the		
213	commissioner's determination under subparagraph (B)(i) of this		
214	subdivision.		
215	(C) Failure of an insurer, health care center or exempt insurer to		
216	notify the commissioner of the amount of an overpayment within the		
217	time prescribed in subparagraph (A)(ii) of this subdivision constitutes a		
218	waiver of any demand of the insurer, health care center or exempt		
219	insurer against this state on account of such overpayment.		
220	(D) Nothing in this subdivision shall be construed to prohibit or limit		
221	the right of an insurer, health care center or exempt insurer to appear		
222	pursuant to subparagraph (B) of subdivision (4) of this subsection.		
223	(d) If another state, territory or district of the United States, or a		
224	foreign country, imposes on a Connecticut domiciled insurer, fraternal		
225	benefit society, hospital service corporation, medical service		
226	corporation, health care center or other domestic entity a retaliatory		
227	charge for the fee imposed under this section, such domestic entity may,		

- 228 not later than sixty days after receipt of notice of the imposition of the 229 retaliatory charge for such fee, appeal to the Insurance Commissioner 230 for a verification that the fee imposed under this section is subject to 231 retaliation by another state, territory or district of the United States, or a 232 foreign country. If the commissioner verifies, upon appeal to and 233 certification by the commissioner, that the fee imposed under this 234 section is the subject of a retaliatory tax, fee, assessment or other 235 obligation by another state, territory or district of the United States, or a 236 foreign country, such fee shall not be assessed against nondomestic 237 insurers and nondomestic exempt insurers pursuant to this section. Any 238 such domestic insurer, fraternal benefit society, hospital service 239 corporation, medical service corporation, health care center or other 240 entity aggrieved by the commissioner's decision issued under this 241 subsection may appeal therefrom in the same manner as provided 242 under section 38a-52 of the general statutes.
- 243 (e) The Insurance Commissioner may adopt regulations, in 244 accordance with chapter 54 of the general statutes, to implement the 245 provisions of this section.
- Sec. 5. Section 38a-1084 of the general statutes is repealed and the following is substituted in lieu thereof (*Effective July 1, 2021*):
- 248 The exchange shall:
- 249 (1) Administer the exchange for both qualified individuals and qualified employers;
- 251 (2) Commission surveys of individuals, small employers and health care providers on issues related to health care and health care coverage;
- 253 (3) Implement procedures for the certification, recertification and 254 decertification, consistent with guidelines developed by the Secretary 255 under Section 1311(c) of the Affordable Care Act, and section 38a-1086, 256 of health benefit plans as qualified health plans;
- 257 (4) Provide for the operation of a toll-free telephone hotline to

258 respond to requests for assistance;

- 259 (5) Provide for enrollment periods, as provided under Section 260 1311(c)(6) of the Affordable Care Act;
 - (6) Maintain an Internet web site through which enrollees and prospective enrollees of qualified health plans may obtain standardized comparative information on such plans including, but not limited to, the enrollee satisfaction survey information under Section 1311(c)(4) of the Affordable Care Act and any other information or tools to assist enrollees and prospective enrollees evaluate qualified health plans offered through the exchange;
 - (7) Publish the average costs of licensing, regulatory fees and any other payments required by the exchange and the administrative costs of the exchange, including information on moneys lost to waste, fraud and abuse, on an Internet web site to educate individuals on such costs;
 - (8) On or before the open enrollment period for plan year 2017, assign a rating to each qualified health plan offered through the exchange in accordance with the criteria developed by the Secretary under Section 1311(c)(3) of the Affordable Care Act, and determine each qualified health plan's level of coverage in accordance with regulations issued by the Secretary under Section 1302(d)(2)(A) of the Affordable Care Act;
 - (9) Use a standardized format for presenting health benefit options in the exchange, including the use of the uniform outline of coverage established under Section 2715 of the Public Health Service Act, 42 USC 300gg-15, as amended from time to time;
 - (10) Inform individuals, in accordance with Section 1413 of the Affordable Care Act, of eligibility requirements for the Medicaid program under Title XIX of the Social Security Act, as amended from time to time, the Children's Health Insurance Program (CHIP) under Title XXI of the Social Security Act, as amended from time to time, or any applicable state or local public program, and enroll an individual in such program if the exchange determines, through screening of the

- application by the exchange, that such individual is eligible for any suchprogram;
- (11) Collaborate with the Department of Social Services, to the extent possible, to allow an enrollee who loses premium tax credit eligibility under Section 36B of the Internal Revenue Code and is eligible for HUSKY A or any other state or local public program, to remain enrolled in a qualified health plan;
 - (12) Establish and make available by electronic means a calculator to determine the actual cost of coverage after application of any premium tax credit under Section 36B of the Internal Revenue Code and any cost-sharing reduction under Section 1402 of the Affordable Care Act;
- (13) Establish a program for small employers through which qualified employers may access coverage for their employees and that shall enable any qualified employer to specify a level of coverage so that any of its employees may enroll in any qualified health plan offered through the exchange at the specified level of coverage;
 - (14) Offer enrollees and small employers the option of having the exchange collect and administer premiums, including through allocation of premiums among the various insurers and qualified health plans chosen by individual employers;
- (15) Grant a certification, subject to Section 1411 of the Affordable Care Act, attesting that, for purposes of the individual responsibility penalty under Section 5000A of the Internal Revenue Code, an individual is exempt from the individual responsibility requirement or from the penalty imposed by said Section 5000A because:
- (A) There is no affordable qualified health plan available through the exchange, or the individual's employer, covering the individual; or
- 316 (B) The individual meets the requirements for any other such 317 exemption from the individual responsibility requirement or penalty;

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318	(16) Provide to the Secretary of the Treasury of the United States the		
319	following:		
320	(A) A list of the individuals granted a certification under subdivision		
321	(15) of this section, including the name and taxpayer identification		
322	number of each individual;		
323	(B) The name and taxpayer identification number of each individual		
324	who was an employee of an employer but who was determined to b		
325	eligible for the premium tax credit under Section 36B of the Internal		
326	Revenue Code because:		
327	(i) The employer did not provide minimum essential health benefi		
328	coverage; or		
329	(ii) The employer provided the minimum essential coverage but it		
330	was determined under Section 36B(c)(2)(C) of the Internal Revenue		
331	Code to be unaffordable to the employee or not provide the required		
332	minimum actuarial value; and		
333	(C) The name and taxpayer identification number of:		
334	(i) Each individual who notifies the exchange under Section		
335	1411(b)(4) of the Affordable Care Act that such individual has changed		
336	employers; and		
337	(ii) Each individual who ceases coverage under a qualified health		
338	plan during a plan year and the effective date of that cessation;		
339	(17) Provide to each employer the name of each employee, as		
340	described in subparagraph (B) of subdivision (16) of this section, of th		
341	employer who ceases coverage under a qualified health plan during a		
342	plan year and the effective date of the cessation;		
343	(18) Perform duties required of, or delegated to, the exchange by the		
344	Secretary or the Secretary of the Treasury of the United States related to		
345	determining eligibility for premium tax credits, reduced cost-sharing or		

346	individual responsibility requirement exemptions;	
347 348 349	(19) Select entities qualified to serve as Navigators in accordance with Section 1311(i) of the Affordable Care Act and award grants to enable Navigators to:	
350 351	(A) Conduct public education activities to raise awareness of the availability of qualified health plans;	
352 353 354 355	(B) Distribute fair and impartial information concerning enrollment in qualified health plans and the availability of premium tax credits under Section 36B of the Internal Revenue Code and cost-sharing reductions under Section 1402 of the Affordable Care Act;	
356	(C) Facilitate enrollment in qualified health plans;	
357 358 359 360 361 362 363	(D) Provide referrals to the Office of the Healthcare Advocate or health insurance ombudsman established under Section 2793 of the Public Health Service Act, 42 USC 300gg-93, as amended from time to time, or any other appropriate state agency or agencies, for any enrollee with a grievance, complaint or question regarding the enrollee's health benefit plan, coverage or a determination under that plan or coverage; and	
364 365 366	(E) Provide information in a manner that is culturally and linguistically appropriate to the needs of the population being served by the exchange;	
367 368 369 370	(20) Review the rate of premium growth within and outside the exchange and consider such information in developing recommendations on whether to continue limiting qualified employer status to small employers;	
371 372 373	(21) Credit the amount, in accordance with Section 10108 of the Affordable Care Act, of any free choice voucher to the monthly premium of the plan in which a qualified employee is enrolled and	

collect the amount credited from the offering employer;

375376377	(22) Consult with stakeholders relevant to carrying out the activities required under sections 38a-1080 to 38a-1090, inclusive, including, but not limited to:	
378 379 380 381	have background or experience in making informed decisions regarding health, medical and scientific matters and are enrollees in qualified	
382 383	(B) Individuals and entities with experience in facilitating enrollment in qualified health plans;	
384 385	(C) Representatives of small employers and self-employed individuals;	
386	(D) The Department of Social Services; and	
387	(E) Advocates for enrolling hard-to-reach populations;	
388	(23) Meet the following financial integrity requirements:	
389 390 391 392	(A) Keep an accurate accounting of all activities, receipts and expenditures and annually submit to the Secretary, the Governor, the Insurance Commissioner and the General Assembly a report concerning such accountings;	
393 394 395 396	(B) Fully cooperate with any investigation conducted by the Secretary pursuant to the Secretary's authority under the Affordable Care Act and allow the Secretary, in coordination with the Inspector General of the United States Department of Health and Human Services, to:	
397	(i) Investigate the affairs of the exchange;	
398	(ii) Examine the properties and records of the exchange; and	
399 400	(iii) Require periodic reports in relation to the activities undertaken by the exchange; and	

- (C) Not use any funds in carrying out its activities under sections 38a-1080 to 38a-1089, inclusive, that are intended for the administrative and operational expenses of the exchange, for staff retreats, promotional giveaways, excessive executive compensation or promotion of federal or state legislative and regulatory modifications;
- (24) (A) Seek to include the most comprehensive health benefit plans that offer high quality benefits at the most affordable price in the exchange, (B) encourage health carriers to offer tiered health care provider network plans that have different cost-sharing rates for different health care provider tiers and reward enrollees for choosing low-cost, high-quality health care providers by offering lower copayments, deductibles or other out-of-pocket expenses, and (C) offer any such tiered health care provider network plans through the exchange; [and]
- (25) Report at least annually to the General Assembly on the effect of adverse selection on the operations of the exchange and make legislative recommendations, if necessary, to reduce the negative impact from any such adverse selection on the sustainability of the exchange, including recommendations to ensure that regulation of insurers and health benefit plans are similar for qualified health plans offered through the exchange and health benefit plans offered outside the exchange. The exchange shall evaluate whether adverse selection is occurring with respect to health benefit plans that are grandfathered under the Affordable Care Act, self-insured plans, plans sold through the exchange and plans sold outside the exchange; and
- (26) Annually consult with the Office of Health Strategy, established under section 19a-754a, as amended by this act, Commissioner of Social Services and Insurance Commissioner to develop the annual plan required under subsection (b) of section 3 of this act and, subject to the terms of such plan, the approval required under subsection (d) of section 3 of this act and within the funds available in the Covered Connecticut account established under section 2 of this act:

- 461 (8) "Pharmaceutical manufacturer" means a person that 462 manufactures a prescription drug and sells, directly or through another 463 person, the prescription drug for distribution in this state;
- (9) "Prescription drug" means a legend drug approved by the federal Food and Drug Administration, or any successor agency, and prescribed by a health care provider to an individual in this state;
- (10) "Reference price" means the wholesale acquisition cost of a drug
 (A) on January 1, 2021, or (B) if the drug is first commercially marketed
 in the United States after January 1, 2021, on the date such drug is first
 commercially marketed in the United States; and
- 471 (11) "Wholesale acquisition cost" has the same meaning as provided 472 in 42 USC 1395w-3a, as amended from time to time.
- Sec. 7. (NEW) (*Effective July 1, 2021*) (a) (1) Notwithstanding any provision of the general statutes and except as provided in subdivision (2) of this subsection, no pharmaceutical manufacturer shall, on or after January 1, 2022, sell a prescription drug in this state at a price that exceeds the sum of:
- 478 (A) The reference price for the prescription drug, adjusted for any 479 increase or decrease in the consumer price index; and
- (B) Two per cent of the reference price for the prescription drug for each twelve-month period that has elapsed since the date on which the reference price for such prescription drug was determined, compounded annually on the anniversary of such date.
 - (2) A pharmaceutical manufacturer may sell a prescription drug in this state at a price that exceeds the amount determined for the prescription drug under subdivision (1) of this subsection if the Secretary of Health and Human Services determines, pursuant to 21 USC 356e, as amended from time to time, that such prescription drug is in shortage in the United States.

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- (b) (1) Except as provided in subdivision (2) of this subsection, any pharmaceutical manufacturer that violates the provisions of subsection (a) of this section shall be liable to this state for a civil penalty. Such civil penalty shall be determined and collected on a calendar year basis, and the amount of such civil penalty for a calendar year shall be equal to eighty per cent of the difference between:
 - (A) The revenue that the pharmaceutical manufacturer earned from all sales of the identified prescription drug in this state during the calendar year; and
 - (B) The revenue that the pharmaceutical manufacturer would have earned from all sales of the identified prescription drug in this state during the calendar year if the pharmaceutical manufacturer had sold such identified prescription drug at a price that did not exceed the amount described in subsection (a) of this section.
- (2) No pharmaceutical manufacturer of an identified prescription drug shall be liable to this state for the civil penalty imposed under subdivision (1) of this subsection unless the pharmaceutical manufacturer made at least two hundred fifty thousand dollars in total annual sales in this state for the calendar year for which such civil penalty would otherwise be imposed.
 - (c) (1) (A) Not later than March 1, 2023, and annually thereafter, each pharmaceutical manufacturer that violated subsection (a) of this section during the preceding calendar year shall:
- 513 (i) Pay to the commissioner the civil penalty imposed under 514 subsection (b) of this section for such calendar year; and
- 515 (ii) File with the commissioner a statement for such calendar year in 516 a form and manner, and containing all information, prescribed by the 517 commissioner.
- 518 (B) A pharmaceutical manufacturer that is required to file a statement 519 and pay a civil penalty pursuant to subparagraph (A) of this subdivision

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- 520 shall electronically file such statement and make such payment by electronic funds transfer in the manner provided by chapter 228g of the general statutes, irrespective of whether the pharmaceutical 523 manufacturer would have otherwise been required to electronically file such statement or make such payment by electronic funds transfer 525 under chapter 228g of the general statutes.
 - (2) If no statement is filed pursuant to subdivision (1) of this subsection, the commissioner may make such statement at any time thereafter, according to the best obtainable information and the prescribed form.
 - (d) The commissioner may examine such records of a pharmaceutical manufacturer that is subject to the civil penalty imposed under subsection (b) of this section as the commissioner deems necessary. If the commissioner determines from such examination that the pharmaceutical manufacturer failed to pay the full amount of such civil penalty, the commissioner shall bill such pharmaceutical manufacturer for the full amount of such civil penalty.
 - (e) (1) The commissioner may require all pharmaceutical manufacturers subject to a civil penalty imposed under this section to keep such records as the commissioner may prescribe, and may require the production of books, papers, documents and other data, to provide or secure information pertinent to the determination of the civil penalty and the enforcement and collection thereof.
 - (2) The commissioner, or any person authorized by the commissioner, may examine the books, papers, records and equipment of any person liable under the provisions of this section and may investigate the character of the business of such person to verify the accuracy of any statement made or, if no statement is made by the person, to ascertain and determine the amount required to be paid.
 - (f) Any pharmaceutical manufacturer that is subject to a civil penalty imposed under this section and aggrieved by the action of the

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commissioner under subdivision (2) of subsection (c) of this section or subsection (d) of this section may apply to the commissioner, in writing and not later than sixty days after the notice of such action is delivered or mailed to such pharmaceutical manufacturer, for a hearing, setting forth the reasons why such hearing should be granted and the amount by which the civil penalty should be reduced. The commissioner shall promptly consider each such application and may grant or deny the hearing requested. If the hearing request is denied, the commissioner shall immediately notify the pharmaceutical manufacturer. If the hearing request is granted, the commissioner shall notify the pharmaceutical manufacturer of the date, time and place for such hearing. After such hearing, the commissioner may make such order as appears just and lawful to the commissioner and shall furnish a copy of such order to the pharmaceutical manufacturer. The commissioner may, by notice in writing, order a hearing on the commissioner's own initiative and require a pharmaceutical manufacturer, or any other person who the commissioner believes to be in possession of relevant information concerning such pharmaceutical manufacturer, to appear before the commissioner or the commissioner's authorized agent with any specified books of account, papers or other documents for examination under oath.

(g) Any pharmaceutical manufacturer that is aggrieved by any order, decision, determination or disallowance of the commissioner made under subsection (f) of this section may, not later than thirty days after service of notice of such order, decision, determination or disallowance, take an appeal therefrom to the superior court for the judicial district of New Britain, which appeal shall be accompanied by a citation to the commissioner to appear before said court. Such citation shall be signed by the same authority and such appeal shall be returnable at the same time and served and returned in the same manner as is required in case of a summons in a civil action. The authority issuing the citation shall take from the appealant a bond or recognizance to this state, with surety, to prosecute the appeal to effect and to comply with the orders and decrees of the court in the premises. Such appeals shall be preferred

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cases, to be heard, unless cause appears to the contrary, at the first session, by the court or by a committee appointed by the court. Said court may grant such relief as may be equitable and, if the civil penalty was paid prior to the granting of such relief, may order the Treasurer to pay the amount of such relief. If the appeal was taken without probable cause, the court may tax double or triple costs, as the case demands and, upon all such appeals that are denied, costs may be taxed against such pharmaceutical manufacturer at the discretion of the court but no costs shall be taxed against this state.

(h) The commissioner, and any agent of the commissioner duly authorized to conduct any inquiry, investigation or hearing pursuant to this section, shall have power to administer oaths and take testimony under oath relative to the matter of inquiry or investigation. At any hearing ordered by the commissioner, the commissioner, or the commissioner's agent authorized to conduct such hearing and having authority by law to issue such process, may subpoena witnesses and require the production of books, papers and documents pertinent to such inquiry or investigation. No witness under subpoena authorized to be issued under the provisions of this section shall be excused from testifying or from producing books, papers or documentary evidence on the ground that such testimony or the production of such books, papers or documentary evidence would tend to incriminate such witness, but such books, papers or documentary evidence so produced shall not be used in any criminal proceeding against such witness. If any person disobeys such process or, having appeared in obedience thereto, refuses to answer any pertinent question put to such person by the commissioner, or the commissioner's authorized agent, or to produce any books, papers or other documentary evidence pursuant thereto, the commissioner, or such agent, may apply to the Superior Court of the judicial district wherein the pharmaceutical manufacturer resides or wherein the business was conducted, or to any judge of such court if the same is not in session, setting forth such disobedience to process or refusal to answer, and such court or such judge shall cite such person to appear before such court or such judge to answer such question or to

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produce such books, papers or other documentary evidence and, upon such person's refusal so to do, shall commit such person to a community correctional center until such person testifies, but not for a period longer than sixty days. Notwithstanding the serving of the term of such commitment by any person, the commissioner may proceed in all respects with such inquiry and examination as if the witness had not previously been called upon to testify. Officers who serve subpoenas issued by the commissioner or under the commissioner's authority and witnesses attending hearings conducted by the commissioner pursuant to this section shall receive fees and compensation at the same rates as officers and witnesses in the courts of this state, to be paid on vouchers of the commissioner on order of the Comptroller from the proper appropriation for the administration of this section.

(i) The amount of any civil penalty unpaid under the provisions this section may be collected under the provisions of section 12-35 of the general statutes. The warrant provided under section 12-35 of the general statutes shall be signed by the commissioner or the commissioner's authorized agent. The amount of any such civil penalty shall be a lien on the real property of the pharmaceutical manufacturer from the last day of the month next preceding the due date of such civil penalty until such civil penalty is paid. The commissioner may record such lien in the records of any town in which the real property of such pharmaceutical manufacturer is situated, but no such lien shall be enforceable against a bona fide purchaser or qualified encumbrancer of such real property. When any civil penalty with respect to which a lien was recorded under the provisions of this subsection is satisfied, the commissioner shall, upon request of any interested party, issue a certificate discharging such lien, which certificate shall be recorded in the same office in which such lien was recorded. Any action for the foreclosure of such lien shall be brought by the Attorney General in the name of this state in the Superior Court for the judicial district in which the real property subject to such lien is situated, or, if such property is located in two or more judicial districts, in the Superior Court for any one such judicial district, and the court may limit the time for

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redemption or order the sale of such real property or make such other or further decree as it judges equitable. The provisions of section 12-39g of the general statutes shall apply to all civil penalties imposed under this section.

- (j) (1) Any officer or employee of a pharmaceutical manufacturer who owes a duty to the pharmaceutical manufacturer to pay a civil penalty imposed under this section on behalf of such pharmaceutical manufacturer, file a statement with the commissioner pursuant to subsection (c) of this section on behalf of such pharmaceutical manufacturer, keep records or supply information to the commissioner on behalf of such pharmaceutical manufacturer pursuant to this section and wilfully fails, at the time required under this section, to pay such civil penalty, file such statement, keep such records or supply such information on behalf of such pharmaceutical manufacturer shall, in addition to any other penalty provided by law, be fined not more than one thousand dollars or imprisoned not more than one year, or both. Notwithstanding the provisions of section 54-193 of the general statutes, no such officer or employee shall be prosecuted for a violation of the provisions of this subdivision committed on or after July 1, 2021, except within three years next after such violation is committed.
- (2) Any officer or employee of a pharmaceutical manufacturer who owes a duty to the pharmaceutical manufacturer to deliver or disclose to the commissioner, or the commissioner's authorized agent, any list, statement, return, account statement or other document on behalf of such pharmaceutical manufacturer and wilfully delivers or discloses to the commissioner, or the commissioner's authorized agent, any such list, statement, return, account statement or other document that such officer or employee knows to be fraudulent or false in any material matter shall, in addition to any other penalty provided by law, be guilty of a class D felony.
- (3) No officer or employee of a pharmaceutical manufacturer shall be charged with an offense under subdivisions (1) and (2) of this subsection in relation to the same civil penalty, but such officer or employee may

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- be charged and prosecuted for both such offenses upon the same information.
- 688 (k) The proceeds from all civil penalties imposed under this section 689 shall be deposited in the Covered Connecticut account. Each civil 690 penalty imposed under this section shall be deemed to constitute a civil 691 fine or penalty within the meaning of 42 USC 1396b(w), as amended 692 from time. No portion of any civil penalty imposed under this section shall be waived under section 12-3a of the general statutes or any other 693 694 applicable law. No tax credit shall be allowable against any civil penalty 695 imposed under this section.
- (l) Not later than July 1, 2023, and annually thereafter, the commissioner shall prepare a list containing the name of each pharmaceutical manufacturer that violated subsection (a) of this section during the preceding calendar year. The commissioner shall make each list publicly available.
- 701 (m) The commissioner may adopt regulations, in accordance with the 702 provisions of chapter 54 of the general statutes, to implement the 703 provisions of this section.
 - Sec. 8. (NEW) (*Effective July 1, 2021*) (a) No pharmaceutical manufacturer of an identified prescription drug shall withdraw the identified prescription drug from sale in this state for the purpose of avoiding the civil penalty imposed under subsection (b) of section 7 of this act.
 - (b) Any pharmaceutical manufacturer that intends to withdraw an identified prescription drug from sale in this state shall send advance written notice to the Office of Health Strategy disclosing such pharmaceutical manufacturer's intention at least one hundred eighty days before such withdrawal.
 - (c) Any pharmaceutical manufacturer that violates any provision of subsection (a) or (b) of this section shall be liable to this state for a civil penalty in the amount of five hundred thousand dollars.

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This act shall take effect as follows and shall amend the following						
sections:						
Section 1	July 1, 2021	19a-754a(a) and (b)				
Sec. 2	July 1, 2021	New section				
Sec. 3	July 1, 2021	New section				
Sec. 4	July 1, 2021	New section				
Sec. 5	July 1, 2021	38a-1084				
Sec. 6	July 1, 2021	New section				
Sec. 7	July 1, 2021	New section				
Sec. 8	July 1, 2021	New section				

Statement of Legislative Commissioners:

In Section 3(a), Subdiv. (5) was deleted to eliminate an unnecessary definition.

INS Joint Favorable Subst.