274	an action specified by the department under Subsection 26-62-302(3) or Subsection
275	<u>26-62-304(3).</u>
276	(c) (i) A fee collected by the department under Subsection (3)(a) is a dedicated credit
277	for use by the department to implement this chapter.
278	(ii) A fee collected by the Insurance Department under Subsection (3)(b) is a dedicated
279	credit for use by the Insurance Department to perform the functions described in Subsection
280	<u>(3)(b).</u>
281	(d) The fees in Subsections (3)(a) and (b) may not exceed the amount necessary to
282	cover the cost the department incurs to implement this chapter.
283	(e) The department shall deposit in the General Fund the fees described in Subsection
284	(3)(a) as a dedicated credit to be used solely to pay for the cost of implementing this chapter.
285	(4) Before the conditions described in Subsection (1) are satisfied, the department:
286	(a) may, to the extent allowed under United State federal and state law:
287	(i) design the prescription drug importation program; and
288	(ii) negotiate with wholesalers in Canada and the United States regarding the potential
289	implementation of the prescription drug importation program; and
290	(b) may not:
291	(i) allow the importation of any prescription drugs under this chapter; or
292	(ii) implement any provisions of the prescription drug importation program that would
293	violate United States federal or state law.
294	Section 10. Section 26-62-401 is enacted to read:
295	26-62-401. Pharmaceutical manufacturer Prohibited conduct Penalties.
296	(1) A pharmaceutical manufacturer may not:
297	(a) take any action, by agreement, unilaterally, or otherwise, that has the effect of
298	fixing or otherwise controlling the price that a pharmaceutical supplier, distributor, or dispenser
299	charges or advertises for pharmaceuticals in the drug importation program; or
300	(b) discriminate against a pharmaceutical supplier, distributor, or dispenser based on
301	whether the supplier, distributor, or dispenser participates in the prescription drug importation
302	program.
303	Ĥ→ [(2) The attorney general may bring a civil action or seek an injunction against any
304	person who violates a provision of this section, and may seek any remedy available to the

305	H→ <u>attorney general for violations of 11tle /6, Chapter 10, Part 31, Utan Antitrust Act.</u>
306	Section 11. Section 26-62-402 is enacted to read:
307	26-62-402. Pharmaceutical manufacturer Report required.
308	(1) For each drug that has an annual wholesale acquisition cost of \$10,000 or more, a
309	pharmaceutical manufacturer shall submit a report to the department if a price increase for that
310	drug will result in an increase in the wholesale acquisition cost that is equal to:
311	(a) 7.5% or more over a period of 12 months; or
312	(b) 18% or more over a period of 36 months.
313	(2) The report described in Subsection (1) shall:
314	(a) be submitted to the department no later than 30 days before the day on which the
315	price increase takes effect; and
316	(b) include, for each drug for which a report is required under Subsection (1):
317	(i) the increase in the cost of the drug, expressed as a percentage increase based on the
318	price of the drug before the cost increase;
319	(ii) a justification for each price increase;
320	(iii) the date on which each price increase takes effect;
321	(iv) the total profit derived from sales of the drug, expressed in total dollars and as a
322	percentage of the pharmaceutical manufacturer's total profits for that calendar year;
323	(v) the total expenditures of the pharmaceutical manufacturer on materials and
324	manufacturing for the drug;
325	(vi) the total research and development costs paid by the pharmaceutical manufacturer
326	for the development and production of the drug;
327	(vii) the total administrative, marketing, and advertising costs for the drug; and
328	(viii) costs associated with direct-to-consumer coupons and patient assistance programs
329	for the drug.
330	(3) (a) The department shall publish information submitted to the department under
331	this section:
332	(i) at least once in every three month period; and
333	(ii) in a manner that allows the information to be identified separately for each drug.
334	(b) Notwithstanding Subsection (3)(a), the department may not disclose a trade secret,
335	as defined in Section 13-24-2, under this section. ←Ĥ

336	$\hat{H} \rightarrow \frac{(4)}{(4)}$ Information submitted to the department under this section is a private record for
337	the purpose of Title 63G, Chapter 2, Government Records Access and Management Act.] ←Ĥ
338	Section $\hat{\mathbf{H}} \rightarrow [12] \underline{11} \leftarrow \hat{\mathbf{H}}$. Section 63I-1-226 is amended to read:
339	63I-1-226. Repeal dates, Title 26.
340	(1) Section 26-1-40 is repealed July 1, 2019.
341	(2) Title 26, Chapter 9f, Utah Digital Health Service Commission Act, is repealed July
342	1, 2025.
343	(3) Section 26-10-11 is repealed July 1, 2020.
344	(4) Title 26, Chapter 33a, Utah Health Data Authority Act, is repealed July 1, 2024.
345	(5) Title 26, Chapter 36a, Hospital Provider Assessment Act, is repealed July 1, 2019.
346	(6) Title 26, Chapter 36b, Inpatient Hospital Assessment Act, is repealed July 1, 2021
347	[(7) Section 26-38-2.5 is repealed July 1, 2017.]
348	[(8) Section 26-38-2.6 is repealed July 1, 2017.]
349	[(9)] (7) Title 26, Chapter 56, Hemp Extract Registration Act, is repealed July 1, 2021
350	(8) Title 26, Chapter 62, Prescription Drug Affordability Act, is repealed July 1, 2028.
351	Section $\hat{\mathbf{H}} \rightarrow [13] \ \underline{12} \leftarrow \hat{\mathbf{H}}$. Section 63I-1-276 is amended to read:
352	63I-1-276. Repeal dates, Title 76.
353	(1) Subsection 76-10-526(15) is repealed July 1, 2018.
354	(2) Subsection 76-10-3104(3) is repealed July 1, 2028.
355	Section $\hat{H} \rightarrow [14] \underline{13} \leftarrow \hat{H}$. Section 76-10-3104 is amended to read:
356	76-10-3104. Illegal anticompetitive activities.
357	(1) Every contract, combination in the form of trust or otherwise, or conspiracy in
358	restraint of trade or commerce is declared to be illegal.
359	(2) It shall be unlawful for any person to monopolize, or attempt to monopolize, or
360	combine or conspire with any other person or persons to monopolize, any part of trade or
361	commerce.
362	(3) For purposes of the importation of prescription drugs under Title 26, Chapter 62,
363	Prescription Drug Affordability Act, in addition to the activities described in Subsections (1)
364	and (2), a unilateral act in the form of a trust or otherwise, in restraint of trade or commerce, is
365	unlawful.