Representative Gage Froerer proposes the following substitute bill:

1	CHARITABLE PRESCRIPTION DRUG RECYCLING
2	PROGRAM
3	2016 GENERAL SESSION
4	STATE OF UTAH
5	Chief Sponsor: Gage Froerer
6	Senate Sponsor:
7	
8	LONG TITLE
9	General Description:
10	This bill creates a program that allows certain pharmacies to accept and dispense
11	donated unused prescription medications to certain individuals.
12	Highlighted Provisions:
13	This bill:
14	 amends the Pharmacy Practice Act;
15	defines terms;
16	 directs the Division of Occupational and Professional Licensing (DOPL) to make
17	rules, in consultation with the Utah State Board of Pharmacy, to create a charitable
18	prescription drug recycling program;
19	 establishes criteria for prescription drugs eligible for the program;
20	establishes requirements for donors and pharmacies;
21	 limits the liability of program participants and drug manufacturers;
22	 directs DOPL to make rules establishing certain requirements, standards,
23	procedures, and processes; and
24	makes technical changes.
25	Money Appropriated in this Bill:





26	None
27	Other Special Clauses:
28	None
29	Utah Code Sections Affected:
30	AMENDS:
31	58-17b-502, as last amended by Laws of Utah 2015, Chapter 336
32	58-17b-503, as last amended by Laws of Utah 2011, Chapter 366
33	ENACTS:
34	58-17b-901, Utah Code Annotated 1953
35	58-17b-902, Utah Code Annotated 1953
36	58-17b-903, Utah Code Annotated 1953
37	58-17b-904, Utah Code Annotated 1953
38	58-17b-905, Utah Code Annotated 1953
39	58-17b-906, Utah Code Annotated 1953
40	58-17b-907, Utah Code Annotated 1953
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41	Be it enacted by the Legislature of the state of Utah:
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42 43	Section 1. Section 58-17b-502 is amended to read:
42 43 44	Section 1. Section 58-17b-502 is amended to read: 58-17b-502 . Unprofessional conduct.
42 43 44 45	Section 1. Section 58-17b-502 is amended to read: 58-17b-502 . Unprofessional conduct . "Unprofessional conduct" includes:
42 43 44 45 46	Section 1. Section 58-17b-502 is amended to read: 58-17b-502. Unprofessional conduct. "Unprofessional conduct" includes: (1) willfully deceiving or attempting to deceive the division, the board, or their agents
42 43 44 45 46 47	Section 1. Section 58-17b-502 is amended to read: 58-17b-502. Unprofessional conduct. "Unprofessional conduct" includes: (1) willfully deceiving or attempting to deceive the division, the board, or their agents as to any relevant matter regarding compliance under this chapter;
42 43 44 45 46 47 48	Section 1. Section 58-17b-502 is amended to read: 58-17b-502. Unprofessional conduct. "Unprofessional conduct" includes: (1) willfully deceiving or attempting to deceive the division, the board, or their agents as to any relevant matter regarding compliance under this chapter; (2) (a) except as provided in Subsection (2)(b):
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57	(iii) providing compensation for services to a veterinarian.
58	(3) misbranding or adulteration of any drug or device or the sale, distribution, or
59	dispensing of any outdated, misbranded, or adulterated drug or device;
60	(4) engaging in the sale or purchase of drugs or devices that are samples or packages
61	bearing the inscription "sample" or "not for resale" or similar words or phrases;
62	(5) except as provided in Section 58-17b-503 or Part 9, Charitable Prescription Drug
63	Recycling Act, accepting back and redistributing [of] any unused drug, or a part of it, after it
64	has left the premises of any pharmacy, unless the drug is in a unit pack, as defined in Section
65	58-17b-503, or the manufacturer's sealed container, as defined in rule;
66	(6) an act in violation of this chapter committed by a person for any form of
67	compensation if the act is incidental to the person's professional activities, including the
68	activities of a pharmacist, pharmacy intern, or pharmacy technician;
69	(7) violating Federal Title II, P.L. 91, Controlled Substances Act, Title 58, Chapter 37
70	Utah Controlled Substances Act, or rules or regulations adopted under either act;
71	(8) requiring or permitting pharmacy interns or technicians to engage in activities
72	outside the scope of practice for their respective license classifications, as defined in this
73	chapter and division rules made in collaboration with the board, or beyond their scope of
74	training and ability;
75	(9) administering:
76	(a) without appropriate training, as defined by rule;
77	(b) without a physician's order, when one is required by law; and
78	(c) in conflict with a practitioner's written guidelines or written protocol for
79	administering;
80	(10) disclosing confidential patient information in violation of the provisions of the
81	Health Insurance Portability and Accountability Act of 1996 or other applicable law;
82	(11) engaging in the practice of pharmacy without a licensed pharmacist designated as
83	the pharmacist-in-charge;
84	(12) failing to report to the division any adverse action taken by another licensing
85	jurisdiction, government agency, law enforcement agency, or court for conduct that in

(13) as a pharmacist or pharmacy intern, compounding a prescription drug in a dosage

substance would be considered unprofessional conduct under this section; and

88	form which is regularly and commonly available from a manufacturer in quantities and
89	strengths prescribed by a practitioner.
90	Section 2. Section 58-17b-503 is amended to read:
91	58-17b-503. Exception to unprofessional conduct.
92	(1) For purposes of this section:
93	(a) "Licensed intermediate care facility for people with an intellectual disability" means
94	an intermediate care facility for people with an intellectual disability that is licensed as a
95	nursing care facility or a small health care facility under Title 26, Chapter 21, Health Care
96	Facility Licensing and Inspection Act.
97	(b) "Nursing care facility" [has the same definition as] means the same as that term is
98	defined in Section 26-21-2.
99	(c) "Unit pack" means a tamper-resistant nonreusable single-dose single-drug package
100	with identification that indicates the lot number and expiration date for the drug.
101	(2) [Notwithstanding the provisions of Subsection 58-17b-502(5), a] A pharmacist
102	may <u>:</u>
103	(a) accept and redistribute an unused drug under Part 9, Charitable Prescription Drug
104	Recycling Act; or
105	(b) accept back and redistribute any unused drug, or a part of it, after it has left the
106	premises of the pharmacy if:
107	[(a)] (i) the drug was prescribed to a patient in a nursing care facility, $[a]$ licensed
108	intermediate care facility for people with an intellectual disability, or state prison facility,
109	county jail, or state hospital;
110	[(b)] (ii) the drug was stored under the supervision of a licensed health care provider
111	according to manufacturer recommendations;
112	[(c)] (iii) the drug is in a unit pack or in the manufacturer's sealed container;
113	[(d)] (iv) the drug was returned to the original dispensing pharmacy;
114	[(e)] (v) the drug was initially dispensed by a licensed pharmacist or licensed pharmacy
115	intern; and
116	[(f)] (vi) accepting back and [redistribution] redistributing of the drug complies with
117	federal Food and Drug Administration and Drug Enforcement Administration regulations.
118	Section 3. Section 58-17b-901 is enacted to read:

119	Part 9. Charitable Prescription Drug Recycling Act
120	<u>58-17b-901.</u> Title.
121	This part is known as the "Charitable Prescription Drug Recycling Act."
122	Section 4. Section 58-17b-902 is enacted to read:
123	58-17b-902. Definitions.
124	As used in this part:
125	(1) "Assisted living facility" means the same as that term is defined in Section 26-21-2.
126	(2) "Cancer drug" means a drug that controls or kills neoplastic cells and includes a
127	drug used in chemotherapy to destroy cancer cells.
128	(3) "Charitable clinic" means a charitable nonprofit corporation that:
129	(a) holds a valid exemption from federal income taxation issued under Section 501(a),
130	Internal Revenue Code;
131	(b) is exempt from federal income taxation under Section 501(c)(3), Internal Revenue
132	Code;
133	(c) provides, on an outpatient basis, for a period of less than 24 consecutive hours, to
134	an individual not residing or confined at a facility owned or operated by the charitable
135	nonprofit corporation:
136	(i) advice;
137	(ii) counseling;
138	(iii) diagnosis;
139	(iv) treatment;
140	(v) surgery; or
141	(vi) care or services relating to the preservation or maintenance of health; and
142	(d) has a licensed outpatient pharmacy.
143	(4) "Charitable pharmacy" means an eligible pharmacy that is operated by a charitable
144	<u>clinic.</u>
145	(5) "County health department" means the same as that term is defined in Section
146	<u>26A-1-102.</u>
147	(6) "Donated prescription drug" means a prescription drug that an eligible donor
148	donates to an eligible pharmacy under the program.
149	(7) "Eligible donor" means a donor that donates a prescription drug from within the

150	state and is:
151	(a) a nursing care facility;
152	(b) an assisted living facility;
153	(c) a licensed intermediate care facility for people with an intellectual disability;
154	(d) a manufacturer;
155	(e) a pharmaceutical wholesale distributor;
156	(f) an eligible pharmacy; or
157	(g) a physician's office.
158	(8) "Eligible pharmacy" means a pharmacy that:
159	(a) is registered by the division as eligible to participate in the program; and
160	(b) is operated by:
161	(i) a county;
162	(ii) a county health department;
163	(iii) a pharmacy under contract with a county health department;
164	(iv) the Department of Health, created in Section 26-1-4;
165	(v) the Division of Substance Abuse and Mental Health, created in Section
166	<u>62A-15-103; or</u>
167	(vi) a charitable clinic.
168	(9) "Eligible prescription drug" means a prescription drug, described in Section
169	58-17b-904, that is not a controlled substance.
170	(10) "Licensed intermediate care facility for people with an intellectual disability"
171	means the same as that term is defined in Section 58-17b-503.
172	(11) "Medically indigent individual" means an individual who does not have health
173	insurance and lacks reasonable means to purchase prescribed medications.
174	(12) "Nursing care facility" means the same as that term is defined in Section
175	<u>26-18-501.</u>
176	(13) "Physician's office" means a fixed medical facility that:
177	(a) is staffed by a physician, physician's assistant, nurse practitioner, or registered
178	nurse, licensed under Title 58, Occupations and Professions; and
179	(b) treats an individual who presents at, or is transported to, the facility.
180	(14) "Program" means the Charitable Prescription Drug Recycling Program created in

181	Section 58-17b-903.
182	(15) "Unit pack" means the same as that term is defined in Section 58-17b-503.
183	(16) "Unlawful conduct" means the same as that term is defined in Sections 58-1-501
184	and 58-17b-501.
185	(17) "Unprofessional conduct" means the same as that term is defined in Sections
186	<u>58-1-501</u> and <u>58-17b-502</u> .
187	Section 5. Section 58-17b-903 is enacted to read:
188	58-17b-903. Charitable Prescription Drug Recycling Program Creation
189	Requirements.
190	(1) There is created the Charitable Prescription Drug Recycling Program.
191	(2) The division, in consultation with the board, shall:
192	(a) implement the program, on a statewide basis, to permit an eligible donor to transfer
193	an eligible prescription drug to an eligible pharmacy for dispensing to a medically indigent
194	individual;
195	(b) in accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act,
196	make rules necessary to implement the program; and
197	(c) provide technical assistance to entities that desire to participate in the program.
198	Section 6. Section 58-17b-904 is enacted to read:
199	58-17b-904. Criteria for eligible prescription drugs.
200	An eligible pharmacy may not accept or dispense an unused prescription drug under the
201	program unless the unused prescription drug:
202	(1) (a) is in the original sealed unit pack; or
203	(b) is an injectable medication;
204	(2) (a) is unopened; or
205	(b) is a cancer drug packaged in an unopened single-unit dose that has been removed
206	from a multi-dose package;
207	(3) is accepted and dispensed by the eligible pharmacy before:
208	(a) a beyond use date that appears on the label;
209	(b) the expiration date recommended by the manufacturer; or
210	(c) a date, established by division rule for a specific prescription drug, in accordance
211	with Title 63G. Chapter 3. Utah Administrative Rulemaking Act, that is later than the date in

212	Subsection $(3)(a)$ or $(3)(b)$;
213	(4) (a) is not adulterated or mislabeled; and
214	(b) the pharmacist or licensed pharmacist technician accepting or dispensing the
215	prescription drug does not have reason to believe that the prescription drug is adulterated or
216	mislabeled.
217	Section 7. Section 58-17b-905 is enacted to read:
218	58-17b-905. Participation in program Requirements Fees.
219	(1) An eligible donor or an eligible pharmacy may participate in the program.
220	(2) An eligible pharmacy:
221	(a) shall comply with all applicable federal and state laws related to the storage and
222	distribution of a prescription drug;
223	(b) shall comply with all applicable federal and state laws related to the acceptance and
224	transfer of a prescription drug, including 21 U.S.C. Chapter 9, Subchapter V, Part H,
225	Pharmaceutical Distribution Supply Chain;
226	(c) shall, before accepting or dispensing a prescription drug under the program, inspect
227	each prescription drug to determine whether the prescription drug is an eligible prescription
228	<u>drug;</u>
229	(d) may dispense an eligible prescription drug to a medically indigent individual who:
230	(i) is a resident of the state; and
231	(ii) has a prescription issued by a practitioner;
232	(e) may charge a handling fee, adopted by the division under Section 63J-1-504; and
233	(f) may not accept, transfer, or dispense a prescription drug in violation of the federal
234	Food, Drug, and Cosmetic Act, 21 U.S.C. Sec. 301 et seq.
235	Section 8. Section 58-17b-906 is enacted to read:
236	58-17b-906. Liability of participating organizations and manufacturers.
237	In the absence of bad faith or gross negligence, a person is not criminally or civilly
238	liable for injury, death, or loss of property based solely on the fact that the person
239	manufactured, provided, donated, accepted, or dispensed an eligible prescription drug under
240	this part.
241	Section 9. Section 58-17b-907 is enacted to read:
242	58-17h-907 Rules made by the division

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243	The rules made by the division under Subsection 58-17b-903(2)(b) shall include:
244	(1) registration requirements to establish the eligibility of a pharmacy to participate in
245	the program;
246	(2) a formulary that includes all eligible prescription drugs approved by the federal
247	Food and Drug Administration;
248	(3) standards and procedures for:
249	(a) verifying whether a pharmacy or pharmacist participating in the program is licensed
250	and in good standing with the board;
251	(b) handling of a donated eligible prescription drug, including:
252	(i) acceptance;
253	(ii) identification, including redundant criteria for verification;
254	(iii) documentation, under 21 U.S.C. Sec. 360eee-1, of transaction information, history,
255	and statements;
256	(iv) safe storage;
257	(v) security;
258	(vi) inspection;
259	(vii) transfer; and
260	(viii) dispensing;
261	(c) a pharmacist or licensed pharmacy technician working in or consulting with a
262	participating eligible donor;
263	(d) disposition of a donated prescription drug that is a controlled substance;
264	(e) record keeping regarding:
265	(i) the eligible donor that donated each prescription drug;
266	(ii) the identification and evaluation of a donated prescription drug by a pharmacist or
267	licensed pharmacy technician; and
268	(iii) the dispensing or disposition of a prescription drug;
269	(f) determining the status of a medically indigent individual;
270	(g) labeling requirements to:
271	(i) ensure compliance with patient privacy laws relating to:
272	(A) an individual who receives an eligible prescription drug; and
273	(B) patient information that may appear on a donated prescription drug;

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274	(ii) clearly identify an eligible prescription drug dispensed under the program; and
275	(iii) communicate necessary information regarding the manufacturer's recommended
276	expiration date or the beyond use date; and
277	(h) ensuring compliance with the requirements of this part;
278	(4) a process for seeking input from:
279	(a) the Department of Health, created in Section 26-1-4, to establish program standards
280	and procedures for assisted living facilities and nursing care facilities; and
281	(b) the Division of Substance Abuse and Mental Health, created in Section
282	62A-15-103, to establish program standards and procedures for mental health and substance
283	abuse clients; and
284	(5) the creation of a special training program that a pharmacist and a licensed pharmacy
285	technician at an eligible pharmacy must complete before participating in the program.