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17 (b) "Controlled substance" means any substance listed
18 under Schedule II, Schedule III, Schedule IV, or Schedule V of
19 s. 893.03.

20 (c) "Dispenser" means a health care practitioner who,
21 within the scope of his or her practice act, is authorized to
22 dispense medicinal drugs and who does so under this act.

23 (d) "Free clinic" means a clinic that delivers only
24 medical diagnostic services or nonsurgical medical treatment
25 free of charge to low-income recipients.

26 (e) "Health care practitioner" or "practitioner" means a
27 practitioner licensed under this chapter, chapter 458, chapter
28 459, chapter 461, chapter 463, chapter 464, or chapter 466.

29 (f) "Indigent" means having a family income for the 12
30 months preceding the determination of income that is below 200
31 percent of the federal poverty level as defined by the most
32 recently revised poverty income guidelines published by the
33 United States Department of Health and Human Services.

34 (g) "Nonprofit health clinic" means a nonprofit legal
35 entity that provides medical care to patients who are indigent,
36 uninsured, or underinsured. The term includes, but is not
37 limited to, a federally qualified health center as defined in 42
38 U.S.C. s. 1396d(1)(2)(B) and a rural health clinic as defined in
39 42 U.S.C. s. 1396d(1)(1).

40 (h) "Nursing home facility" has the same meaning as in s.
41 400.021.

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42 (i) "Prescriber" means a health care practitioner who,
43 within the scope of his or her practice act, is authorized to
44 prescribe medicinal drugs.

45 (j) "Prescription drug" has the same meaning as the term
46 "medicinal drugs" or "drugs," as those terms are defined in s.
47 465.003(8), but does not include controlled substances, cancer
48 drugs donated under s. 499.029, or drugs with an approved
49 Federal Food and Drug Administration risk evaluation and
50 mitigation strategy that includes elements to assure safe use.

51 (k) "Program" means the Prescription Drug Donation
52 Repository Program created by this section.

53 (l) "Supply" means a material or an instrument used to
54 administer a prescription drug.

55 (m) "Tamper-evident packaging" means a package that has
56 one or more indicators or barriers to access which, if breached
57 or missing, can reasonably be expected to provide visible
58 evidence to consumers that tampering has occurred. The term
59 includes, but is not limited to, unopened unit-dose packaging,
60 multiple-dose packaging, and medications with a seal on their
61 immediate, outer, secondary, or tertiary packaging.

62 (n) "Underinsured" means having health care coverage or
63 prescription drug coverage, but having exhausted these benefits
64 or not having prescription drug coverage for the drug
65 prescribed.

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66 (o) "Uninsured" means having no health care coverage and
67 not being eligible for prescription drug coverage under a
68 program funded in whole or in part by the Federal Government.

69 (3) PRESCRIPTION DRUG DONATION REPOSITORY PROGRAM;
70 CREATION; PURPOSE.—The Prescription Drug Donation Repository
71 Program is created within the department to facilitate
72 prescription drugs and supplies donation to eligible patients.

73 (4) REPOSITORIES.—

74 (a) A repository may accept and dispense eligible
75 donations to eligible patients under the program. Repositories
76 must inspect, store, and dispense donations and report to the
77 department in accordance with this section.

78 (b) The following entities may participate as a
79 repository:

- 80 1. A health care practitioner's office.
- 81 2. A pharmacy.
- 82 3. A hospital with a closed drug delivery system.
- 83 4. A nursing home facility with a closed drug delivery
84 system.
- 85 5. A free clinic or nonprofit health clinic that is
86 licensed or permitted to dispense medicinal drugs in the state.

87 (c) An eligible entity must notify the department of its
88 intent to participate in the program as a repository before
89 accepting or dispensing any donations under the program. The
90 notification must be made on a physical or an electronic form

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91 prescribed by the department by rule and must, at a minimum,
92 include:

93 1. The name, street address, website, and telephone number
94 of the intended repository and any license or registration
95 number issued by the state to the intended repository, including
96 the name of the issuing agency.

97 2. The name and telephone number of the pharmacist
98 employed by or under contract with the intended repository who
99 is responsible for the inspection of donated prescription drugs
100 and supplies.

101 3. A signed and dated statement by the responsible
102 pharmacist affirming that the intended repository meets the
103 eligibility requirements of this section.

104 (b) A repository may withdraw from participation in the
105 program at any time by providing written notice to the
106 department or contractor, as appropriate, on a physical or an
107 electronic form prescribed by the department by rule. The
108 department shall adopt rules addressing the disposition of
109 prescription drugs and supplies in the possession of the
110 withdrawing repository.

111 (5) ELIGIBLE DONORS.—The following entities may donate
112 prescription drugs or supplies to a repository under the
113 program:

114 (a) Nursing home facilities with closed drug delivery
115 systems.

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116 (b) Hospices that have maintained control of a patient's
117 prescription drugs.

118 (c) Hospitals with closed drug delivery systems.

119 (d) Pharmacies.

120 (e) Drug manufacturers or wholesale distributors.

121 (f) Medical device manufacturers or suppliers.

122 (g) Prescribers who receive prescription drugs or supplies
123 directly from a drug manufacturer, wholesale distributor, or
124 pharmacy.

125 (6) ELIGIBLE DONATIONS; DONATION REQUIREMENTS; PROHIBITED
126 DONATIONS.—

127 (a) An eligible donor may only donate a prescription drug
128 to a repository if:

129 1. The drug is approved for medical use in the United
130 States.

131 2. The drug is in unopened tamper-evident packaging.

132 3. The drug requires storage at normal room temperature
133 per the manufacturer or federal requirements.

134 4. The drug has been stored according to manufacturer or
135 federal storage requirements.

136 5. The drug does not have any physical signs of tampering
137 or adulteration and there is no reason to believe that the drug
138 is adulterated.

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139 6. The packaging does not have any physical signs of
140 tampering, misbranding, deterioration, compromised integrity, or
141 adulteration.

142 7. The packaging indicates the expiration date of the
143 drug. If the lot number is not retrievable, all specified
144 medications must be destroyed in the event of a recall.

145 8. The drug has an expiration date that is more than 3
146 months after the date that the drug was donated.

147 (b) An eligible donor may donate a prescription drug
148 supply to a repository only if it is in unopened tamper-evident
149 packaging.

150 (c) Donations must be made on the premises of a repository
151 to a person designated by the repository. A drop box may not be
152 used to accept donations.

153 (d) A prescription drug or supply may not be donated to a
154 specific patient.

155 (7) INSPECTION AND STORAGE.—

156 (a) Upon receipt of a proposed donation, a licensed
157 pharmacist employed by or under contract with a repository shall
158 inspect it to determine whether it meets the requirements of
159 subsections (5) and (6). The repository shall quarantine
160 donations until such inspection is complete and the donation is
161 approved for dispensing.

162 (b) The inspecting pharmacist must sign an inspection
163 record on a physical or an electronic form prescribed by the

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164 department by rule which verifies that the prescription drug or
165 supply meets the criteria of subsections (5) and (6) and must
166 attach the record to the inventory required by paragraph (d). A
167 repository that receives drugs and supplies from another
168 repository is not required to reinspect them.

169 (c) Repositories shall store donations in a secure storage
170 area under the environmental conditions specified by the
171 manufacturer or federal storage requirements. Donations under
172 the program may not be stored with other inventory.

173 (d) Repositories shall maintain an inventory of the name,
174 strength, available quantity, and expiration date of donations
175 and the transaction date and name, address, and phone number of
176 the donor. Repositories shall record such inventory on a
177 physical or an electronic form prescribed by the department by
178 rule.

179 (e) By the fifth day of each month, a repository shall
180 submit to the department its inventory records of donations in
181 the previous month.

182 (f) The department may facilitate redistribution of
183 donations between repositories. A repository that receives
184 donations may, after notifying the department, distribute the
185 donations to another local repository.

186 (8) ELIGIBLE PATIENTS; DISPENSING REQUIREMENTS; PATIENT
187 NOTICE; PROHIBITIONS.-

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188 (a) A repository may dispense an eligible donation to a
189 Florida resident who is indigent, uninsured, or underinsured,
190 and who has a valid prescription for such prescription drug or
191 supply, as applicable.

192 (b) Each new eligible patient must submit an intake
193 collection form to a repository to receive a donation under the
194 program, using a physical or an electronic form prescribed by
195 department in rule. Such form shall include, at a minimum:

196 1. The name, street address, and telephone number of the
197 eligible patient.

198 2. The basis for eligibility, which must specify that the
199 patient is indigent, uninsured, or underinsured.

200 3. A statement physically or electronically signed and
201 dated by the patient affirming that the patient meets the
202 eligibility requirements of this section and will inform the
203 repository if the patient's eligibility changes.

204 4. Notice that the prescription drug was donated to the
205 program, the donors and participants in the program are immune
206 from civil or criminal liability or disciplinary action, and the
207 eligible patient is not required to pay for the prescription
208 drug.

209 5. A statement physically or electronically signed and
210 dated by the eligible patient acknowledging receipt of the
211 information required by subparagraph 4. of this paragraph.

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212 (c) By the fifth day of each month, the repository shall
213 submit to the department a summary of each intake collection
214 form it collected in the previous month.

215 (d) A dispenser may dispense donations, if available, only
216 to an eligible patient who has submitted a completed intake
217 collection form.

218 (e) A dispenser may provide dispensing and consulting
219 services to an eligible patient.

220 (f) Donations under the program may not be sold or resold.

221 (g) A dispenser may not submit a claim or otherwise seek
222 reimbursement from any public or private third-party payor for
223 donations dispensed under the program.

224 (9) RECALLED PRESCRIPTION DRUGS AND SUPPLIES.—

225 (a) Each repository shall establish and follow a protocol
226 for notifying recipients in the event a prescription drug
227 donated under the program is recalled.

228 (b) Repositories shall destroy all donated prescription
229 drugs that are recalled, expired, or not suitable for
230 dispensing. Repositories must complete a destruction form for
231 all such drugs, using a physical or an electronic form
232 prescribed by department in rule.

233 (10) RECORDKEEPING.—

234 (a) Repositories shall maintain records of prescription
235 drugs and supplies that are accepted, donated, dispensed,

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236 distributed, or destroyed under the program in physical or
237 electronic forms prescribed by the department in rule.

238 (b) All required records must be maintained in accordance
239 with any applicable practice act. Repositories shall submit
240 these records monthly to the department for data collection.

241 (11) REGISTRIES; PUBLICATION OF FORMS.—

242 (a) The department shall establish and maintain registries
243 of all repositories and of prescription drugs and supplies
244 available under the program. The registry of repositories must
245 include each repository's name, address, website, and telephone
246 number. The registry of available prescription drugs and
247 supplies must include the name, strength, available quantity,
248 and expiration date of the prescription drug or supplies and the
249 name and contact information of each repository where such drugs
250 or supplies are available. The department shall publish the
251 registries on its website.

252 (b) The department shall publish all forms required by
253 this section on its website.

254 (12) IMMUNITY FROM LIABILITY, DISCIPLINARY ACTION.—

255 (a) Any donor of prescription drugs or supplies and any
256 participant in the program who exercises reasonable care in
257 donating, accepting, distributing, or dispensing prescription
258 drugs or supplies under the program is immune from civil or
259 criminal liability and from professional disciplinary action by

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260 the state for any injury, death, or loss to person or property
261 relating to such activities.

262 (b) A pharmaceutical manufacturer who exercises reasonable
263 care is not liable for any claim or injury arising from the
264 donation of any prescription drug or supply under this section,
265 including, but not limited to, liability for failure to transfer
266 or communicate product or consumer information regarding the
267 donated prescription drug, including its expiration date.

268 (13) RULEMAKING.—The department shall adopt rules
269 necessary to administer this section.

270 Section 2. Paragraph (o) is added to subsection (5) of
271 section 252.36, Florida Statutes, to read:

272 252.36 Emergency management powers of the Governor.—

273 (5) In addition to any other powers conferred upon the
274 Governor by law, she or he may:

275 (o) Waive the patient eligibility requirements of s.
276 465.1902.

277 Section 3. For the 2019-2020 fiscal year, two full-time
278 equivalent positions with associated salary rate of 150,449 are
279 authorized and the sums of \$325,423 in recurring funds and
280 \$78,233 in nonrecurring funds from the Grants and Donations
281 Trust Fund are appropriated to the Department of Health for the
282 purpose of implementing the requirements of this act.

283 Section 4. This act shall take effect July 1, 2019.
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T I T L E A M E N D M E N T

Remove everything before the enacting clause and insert:
An act relating to the Prescription Drug Donation
Repository Program; creating s. 465.1902, F.S.; providing a
short title; defining terms; creating the Prescription Drug
Donation Repository Program within the Department of
Health; specifying the purpose of the program; specifying
entities that may participate as repositories; requiring a
repository to notify the department of its intent to
participate in the program; providing notification
requirements; providing a procedure for a repository to
withdraw from participation in the program; requiring the
department to adopt rules regarding the disposition of
prescription drugs and supplies of a withdrawing
repository; specifying entities that are eligible donors;
providing criteria and procedures for eligible donations;
prohibiting donations to specific patients; providing
inspection, inventory, and storage requirements for
repositories; requiring inspection of donated prescription
drugs and supplies by a licensed pharmacist; requiring a
repository to notify the department within a specified
timeframe after receiving a donation of prescription drugs
or supplies; authorizing the department to redistribute

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310 prescription drugs or supplies; authorizing a repository to
311 transfer prescription drugs or supplies to another
312 repository after notifying the department; specifying
313 patients eligible to receive donated prescription drugs and
314 supplies; specifying conditions for dispensing donated
315 prescription drugs and supplies to eligible patients;
316 providing intake collection form requirements; requiring
317 the intake form to provide certain notice to patients;
318 prohibiting the sale of donated prescription drugs and
319 supplies under the program; requiring repositories to
320 establish a protocol for notifying recipients of a
321 prescription drug recall; providing for destruction of
322 donated prescription drugs under certain circumstances;
323 providing recordkeeping requirements; requiring the
324 department to establish, maintain, and publish a registry
325 of participating repositories and available donated
326 prescription drugs and supplies; requiring the department
327 to publish certain information and forms on its website;
328 providing immunity from civil and criminal liability and
329 from professional disciplinary action for participants
330 under certain circumstances; providing immunity to
331 pharmaceutical manufacturers, under certain circumstances,
332 from any claim or injury arising from the donation of any
333 prescription drug or supply under the program; requiring
334 the department to adopt rules; amending s. 252.36, F.S.;

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COMMITTEE/SUBCOMMITTEE AMENDMENT

Bill No. CS/CS/HB 59 (2019)

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335 | authorizing the Governor to waive program patient
336 | eligibility requirements during a declared state of
337 | emergency; authorizing positions and providing
338 | appropriations; providing an effective date.