

**As Passed by the House**

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**Am. Sub. H. B. No. 231**

**Representative Greenspan**

**Cosponsors: Representatives Jones, Miller, J., Patterson, Abrams, Carruthers, Crawley, Crossman, Ghanbari, Hicks-Hudson, Liston, Patton, Perales, Richardson, Robinson, Rogers, Russo, Scherer, Seitz, Sobecki, Sweeney, Sykes, Upchurch, West**

**A BILL**

To amend sections 3313.713, 3313.719, 4723.50, 1  
4729.01, 4729.51, 4729.513, 4729.541, 4729.60, 2  
and 4729.88 and to enact sections 3301.135, 3  
3313.7115, 3313.7116, 3314.147, 3326.60, 4  
3328.38, 4723.484, 4730.434, 4731.92, and 5  
5101.78 of the Revised Code to require the 6  
Department of Education to notify public and 7  
private schools of free and reduced cost 8  
epinephrine autoinjector programs, to enact the 9  
"Allison Rose Act" with regard to food allergy 10  
training for public schools, and to permit 11  
schools and camps to procure and use glucagon in 12  
certain circumstances. 13

**BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF OHIO:**

**Section 1.** That section 3313.719 be amended and section 14  
3301.135 of the Revised Code be enacted to read as follows: 15

**Sec. 3301.135.** The department of education annually shall 16  
compile a list of organizations and companies that offer free 17

and reduced cost epinephrine autoinjectors to qualifying school 18  
districts, other public schools, and chartered nonpublic 19  
schools. The department shall make this information readily 20  
available on their web site and send a copy of the list by mail 21  
or electronically to each school district, other public school, 22  
and chartered nonpublic school. 23

As used in this section, "other public school" has the 24  
same meaning as in section 3301.0711 of the Revised Code. 25

**Sec. 3313.719.** (A) The board of education of each city, 26  
local, exempted village, and joint vocational school district 27  
and the governing authority of each chartered nonpublic school 28  
shall establish a written policy with respect to protecting 29  
students with ~~peanut or other~~ food allergies. The policy shall 30  
be developed in consultation with parents, school nurses and 31  
other school employees, school volunteers, students, and 32  
community members. 33

(B) Each school district board may create training for all 34  
staff members and age-appropriate instruction for students in 35  
grades kindergarten through twelve on food allergies and ways in 36  
which to assist an individual experiencing an allergic reaction. 37

(C) Training completed under division (B) of this section 38  
may include instruction in food allergies, signs and symptoms of 39  
anaphylaxis, prevention of allergic reactions, management and 40  
administration of epinephrine, and follow-up and reporting 41  
procedures. 42

(D) Training completed under division (B) of this section 43  
shall qualify as a professional development activity for the 44  
renewal of educator licenses, in addition to activities approved 45  
by local professional development committees under division (F) 46

of section 3319.22 of the Revised Code. 47

(E) (1) The following are not liable in damages in a civil 48  
action for injury, death, or loss to person or property that 49  
allegedly arise from an act or omission associated with any 50  
training under divisions (B) and (C) of this section, unless the 51  
act or omission constitutes willful or wanton misconduct: 52

(a) A school or school district; 53

(b) A member of a district board of education; 54

(c) A district or school employee or contractor; 55

(d) A licensed health professional authorized to prescribe 56  
drugs who personally furnishes or prescribes epinephrine 57  
autoinjectors, consults with a superintendent, or issues a 58  
protocol pursuant to section 3313.7110 of the Revised Code; 59

(e) An anaphylaxis training organization and its personnel 60  
where leadership includes a physician authorized under Chapter 61  
4731. of the Revised Code to practice medicine and surgery or 62  
osteopathic medicine and surgery who is board-certified in 63  
allergy and immunology as that designation is issued by a 64  
medical specialty certifying board recognized by the American 65  
board of medical specialties or American osteopathic 66  
association. 67

(2) This section does not eliminate, limit, or reduce any 68  
other immunity or defense that a school or school district, 69  
member of a district board of education, district or school 70  
employee or contractor, or licensed health professional may be 71  
entitled to under Chapter 2744. or any other provision of the 72  
Revised Code or under the common law of this state. 73

**Section 2.** That existing section 3313.719 of the Revised 74

Code is hereby repealed. 75

**Section 3.** Sections 1 and 2 of this act shall be known as 76  
the "Allison Rose Act" in honor of Allison Rose Suhy. 77

**Section 4.** That sections 3313.713, 4723.50, 4729.01, 78  
4729.51, 4729.513, 4729.541, 4729.60, and 4729.88 be amended and 79  
sections 3313.7115, 3313.7116, 3314.147, 3326.60, 3328.38, 80  
4723.484, 4730.434, 4731.92, and 5101.78 of the Revised Code be 81  
enacted to read as follows: 82

**Sec. 3313.713.** (A) As used in this section: 83

(1) "Drug" means a drug, as defined in section 4729.01 of 84  
the Revised Code, that is to be administered pursuant to the 85  
instructions of the prescriber, whether or not required by law 86  
to be sold only upon a prescription. 87

(2) "Federal law" means the "Individuals with Disabilities 88  
Education Act of 1997," 111 Stat. 37, 20 U.S.C. 1400, as 89  
amended. 90

(3) "Prescriber" has the same meaning as in section 91  
4729.01 of the Revised Code. 92

(B) The board of education of each city, local, exempted 93  
village, and joint vocational school district shall adopt a 94  
policy on the authority of its employees, when acting in 95  
situations other than those governed by sections 2305.23, 96  
2305.231, 3313.712, 3313.7110, 3313.7112, ~~and~~ 3313.7113, and 97  
3313.7115 of the Revised Code, to administer drugs prescribed to 98  
students enrolled in the schools of the district. The policy 99  
shall provide either that: 100

(1) Except as otherwise required by federal law, no person 101  
employed by the board shall, in the course of such employment, 102

administer any drug prescribed to any student enrolled in the 103  
schools of the district. 104

(2) Designated persons employed by the board are 105  
authorized to administer to a student a drug prescribed for the 106  
student. Effective July 1, 2011, only employees of the board who 107  
are licensed health professionals, or who have completed a drug 108  
administration training program conducted by a licensed health 109  
professional and considered appropriate by the board, may 110  
administer to a student a drug prescribed for the student. 111  
Except as otherwise provided by federal law, the board's policy 112  
may provide that certain drugs or types of drugs shall not be 113  
administered or that no employee shall use certain procedures, 114  
such as injection, to administer a drug to a student. 115

(C) No drug prescribed for a student shall be administered 116  
pursuant to federal law or a policy adopted under division (B) 117  
of this section until the following occur: 118

(1) The board, or a person designated by the board, 119  
receives a written request, signed by the parent, guardian, or 120  
other person having care or charge of the student, that the drug 121  
be administered to the student. 122

(2) The board, or a person designated by the board, 123  
receives a statement, signed by the prescriber, that includes 124  
all of the following information: 125

(a) The name and address of the student; 126

(b) The school and class in which the student is enrolled; 127

(c) The name of the drug and the dosage to be 128  
administered; 129

(d) The times or intervals at which each dosage of the 130

drug is to be administered;	131
(e) The date the administration of the drug is to begin;	132
(f) The date the administration of the drug is to cease;	133
(g) Any severe adverse reactions that should be reported to the prescriber and one or more phone numbers at which the prescriber can be reached in an emergency;	134 135 136
(h) Special instructions for administration of the drug, including sterile conditions and storage.	137 138
(3) The parent, guardian, or other person having care or charge of the student agrees to submit a revised statement signed by the prescriber to the board or a person designated by the board if any of the information provided by the prescriber pursuant to division (C) (2) of this section changes.	139 140 141 142 143
(4) The person authorized by the board to administer the drug receives a copy of the statement required by division (C) (2) or (3) of this section.	144 145 146
(5) The drug is received by the person authorized to administer the drug to the student for whom the drug is prescribed in the container in which it was dispensed by the prescriber or a licensed pharmacist.	147 148 149 150
(6) Any other procedures required by the board are followed.	151 152
(D) If a drug is administered to a student, the board of education shall acquire and retain copies of the written requests required by division (C) (1) and the statements required by divisions (C) (2) and (3) of this section and shall ensure that by the next school day following the receipt of any such statement a copy is given to the person authorized to administer	153 154 155 156 157 158

drugs to the student for whom the statement has been received. 159  
The board, or a person designated by the board, shall establish 160  
a location in each school building for the storage of drugs to 161  
be administered under this section and federal law. All such 162  
drugs shall be stored in that location in a locked storage 163  
place, except that drugs that require refrigeration may be kept 164  
in a refrigerator in a place not commonly used by students. 165

(E) No person who has been authorized by a board of 166  
education to administer a drug and has a copy of the most recent 167  
statement required by division (C) (2) or (3) of this section 168  
given to the person in accordance with division (D) of this 169  
section prior to administering the drug is liable in civil 170  
damages for administering or failing to administer the drug, 171  
unless such person acts in a manner that constitutes gross 172  
negligence or wanton or reckless misconduct. 173

(F) A board of education may designate a person or persons 174  
to perform any function or functions in connection with a drug 175  
policy adopted under this section either by name or by position, 176  
training, qualifications, or similar distinguishing factors. 177

(G) A policy adopted by a board of education pursuant to 178  
this section may be changed, modified, or revised by action of 179  
the board. 180

(H) Nothing in this section shall be construed to require 181  
a person employed by a board of education to administer a drug 182  
to a student unless the board's policy adopted in compliance 183  
with this section establishes such a requirement. A board shall 184  
not require an employee to administer a drug to a student if the 185  
employee objects, on the basis of religious convictions, to 186  
administering the drug. 187

Nothing in this section affects the application of section 188  
2305.23, 2305.231, 3313.712, 3313.7110, 3313.7112, ~~or~~ 3313.7113, 189  
or 3313.7115 of the Revised Code to the administration of 190  
emergency care or treatment to a student. 191

Nothing in this section affects the ability of a public or 192  
nonpublic school to participate in a school-based fluoride mouth 193  
rinse program established by the director of health pursuant to 194  
section 3701.136 of the Revised Code. Nothing in this section 195  
affects the ability of a person who is employed by, or who 196  
volunteers for, a school that participates in such a program to 197  
administer fluoride mouth rinse to a student in accordance with 198  
section 3701.136 of the Revised Code and any rules adopted by 199  
the director under that section. 200

(I) Nothing in this section shall be construed to require 201  
a school district to obtain written authorization or 202  
instructions from a health care provider to apply 203  
nonprescription topical ointments designed to prevent sunburn. 204  
Furthermore, nothing in this section shall be construed to 205  
prohibit a student to possess and self-apply nonprescription 206  
topical ointment designed to prevent sunburn while on school 207  
property or at a school-sponsored event without written 208  
authorization or instructions from a healthcare provider. The 209  
policy adopted by a school district pursuant to this section 210  
shall not require written authorization from a health care 211  
provider, but may require parental authorization, for the 212  
possession or application of such sunscreen. A designated person 213  
employed by the board of education of a school district shall 214  
apply sunscreen to a student in accordance with the school 215  
district's policy upon request. 216

Sec. 3313.7115. (A) As used in this section, "licensed 217

health professional authorized to prescribe drugs" and 218  
"prescriber" have the same meanings as in section 4729.01 of the 219  
Revised Code. 220

(B) The board of education of each city, local, exempted 221  
village, or joint vocational school district may procure 222  
injectable or nasally administered glucagon for each school 223  
operated by the district to have on the school premises for use 224  
in emergency situations identified under division (D) (5) of this 225  
section by doing one of the following: 226

(1) Having a licensed health professional authorized to 227  
prescribe drugs, acting in accordance with section 4723.484, 228  
4730.434, or 4731.92 of the Revised Code, personally furnish the 229  
injectable or nasally administered glucagon to the school or 230  
school district or issue a prescription for the drug in the name 231  
of the school or district; 232

(2) Having the district's superintendent obtain a 233  
prescriber-issued protocol that includes definitive orders for 234  
injectable or nasally administered glucagon and the dosages to 235  
be administered. 236

A district board that elects to procure injectable or 237  
nasally administered glucagon under this section is encouraged 238  
to maintain, at all times, at least two doses of the drug at 239  
each school operated by the district. 240

(C) A district board that elects to procure injectable or 241  
nasally administered glucagon under this section shall require 242  
the district's superintendent to adopt a policy governing 243  
maintenance and use of the drug. Before adopting the policy, the 244  
superintendent shall consult with a licensed health professional 245  
authorized to prescribe drugs. 246

<u>(D) The policy adopted under division (C) of this section</u>	247
<u>shall do all of the following:</u>	248
<u>(1) Identify the one or more locations in each school</u>	249
<u>operated by the district in which injectable or nasally</u>	250
<u>administered glucagon must be stored;</u>	251
<u>(2) Specify the conditions under which injectable or</u>	252
<u>nasally administered glucagon must be stored, replaced, and</u>	253
<u>disposed;</u>	254
<u>(3) Specify the individuals employed by or under contract</u>	255
<u>with the district board, in addition to a school nurse licensed</u>	256
<u>under section 3319.221 of the Revised Code or an athletic</u>	257
<u>trainer licensed under Chapter 4755. of the Revised Code, who</u>	258
<u>may access and use injectable or nasally administered glucagon</u>	259
<u>in an emergency situation identified under division (D) (5) of</u>	260
<u>this section;</u>	261
<u>(4) Specify any training that employees or contractors</u>	262
<u>specified under division (D) (3) of this section, other than a</u>	263
<u>school nurse or athletic trainer, must complete before being</u>	264
<u>authorized to access and use injectable or nasally administered</u>	265
<u>glucagon;</u>	266
<u>(5) Identify the emergency situations in which a school</u>	267
<u>nurse, athletic trainer, or other employees or contractors</u>	268
<u>specified under division (D) (3) of this section may access and</u>	269
<u>use injectable or nasally administered glucagon;</u>	270
<u>(6) Specify that assistance from an emergency medical</u>	271
<u>service provider must be requested immediately after a dose of</u>	272
<u>glucagon is administered;</u>	273
<u>(7) Specify the individuals, if any, in addition to</u>	274
<u>students, to whom a dose of glucagon may be administered in an</u>	275

<u>emergency situation specified under division (D) (5) of this</u>	276
<u>section.</u>	277
<u>(E) (1) The following are not liable in damages in a civil</u>	278
<u>action for injury, death, or loss to person or property that</u>	279
<u>allegedly arises from an act or omission associated with</u>	280
<u>procuring, maintaining, accessing, or using injectable or</u>	281
<u>nasally administered glucagon under this section, unless the act</u>	282
<u>or omission constitutes willful or wanton misconduct:</u>	283
<u>(a) A school or school district;</u>	284
<u>(b) A member of a district board of education;</u>	285
<u>(c) A district or school employee or contractor;</u>	286
<u>(d) A licensed health professional authorized to prescribe</u>	287
<u>drugs who personally furnishes or prescribes injectable or</u>	288
<u>nasally administered glucagon, consults with a superintendent,</u>	289
<u>or issues a protocol pursuant to this section.</u>	290
<u>(2) This section does not eliminate, limit, or reduce any</u>	291
<u>other immunity or defense that a school or school district,</u>	292
<u>member of a district board of education, district or school</u>	293
<u>employee or contractor, or licensed health professional may be</u>	294
<u>entitled to under Chapter 2744. or any other provision of the</u>	295
<u>Revised Code or under the common law of this state.</u>	296
<u>(F) A school district board of education may accept</u>	297
<u>donations of injectable or nasally administered glucagon from a</u>	298
<u>wholesale distributor of dangerous drugs or manufacturer of</u>	299
<u>dangerous drugs, as defined in section 4729.01 of the Revised</u>	300
<u>Code, and may accept donations of money from any person to</u>	301
<u>purchase the drug.</u>	302
<u>(G) A district board that elects to procure injectable or</u>	303

nasally administered glucagon under this section shall report to 304  
the department of education each procurement and each occurrence 305  
in which a dose of the drug is used from a school's supply. 306

**Sec. 3313.7116.** (A) With the approval of its governing 307  
authority, a chartered or nonchartered nonpublic school may 308  
procure injectable or nasally administered glucagon in the 309  
manner prescribed by section 3313.7115 of the Revised Code. A 310  
chartered or nonchartered nonpublic school that elects to do so 311  
shall comply with all provisions of that section as if it were a 312  
school district. 313

(B) (1) The following are not liable in damages in a civil 314  
action for injury, death, or loss to person or property that 315  
allegedly arises from an act or omission associated with 316  
procuring, maintaining, accessing, or using injectable or 317  
nasally administered glucagon under this section, unless the act 318  
or omission constitutes willful or wanton misconduct: 319

(a) A chartered or nonchartered nonpublic school; 320

(b) A member of a chartered or nonchartered nonpublic 321  
school governing authority; 322

(c) An employee or contractor of the school; 323

(d) A licensed health professional authorized to prescribe 324  
drugs who personally furnishes or prescribes injectable or 325  
nasally administered glucagon, provides a consultation, or 326  
issues a protocol pursuant to this section. 327

(2) This division does not eliminate, limit, or reduce any 328  
other immunity or defense that a chartered or nonchartered 329  
nonpublic school or governing authority, member of a chartered 330  
or nonchartered nonpublic school governing authority, chartered 331  
or nonchartered nonpublic school employee or contractor, or 332

licensed health professional may be entitled to under any other 333  
provision of the Revised Code or the common law of this state. 334

(C) A chartered or nonchartered nonpublic school may 335  
accept donations of injectable or nasally administered glucagon 336  
from a wholesale distributor of dangerous drugs or manufacturer 337  
of dangerous drugs, as defined in section 4729.01 of the Revised 338  
Code, and may accept donations of money from any person to 339  
purchase the drug. 340

(D) A chartered or nonchartered nonpublic school that 341  
elects to procure injectable or nasally administered glucagon 342  
under this section shall report to the department of education 343  
each procurement and each occurrence in which a dose of the drug 344  
is used from the school's supply. 345

**Sec. 3314.147.** (A) With the approval of its governing 346  
authority, a community school established under this chapter may 347  
procure injectable or nasally administered glucagon in the 348  
manner prescribed by section 3313.7115 of the Revised Code. A 349  
community school that elects to do so shall comply with all 350  
provisions of that section as if it were a school district. 351

(B)(1) The following are not liable in damages in a civil 352  
action for injury, death, or loss to person or property that 353  
allegedly arises from an act or omission associated with 354  
procuring, maintaining, accessing, or using injectable or 355  
nasally administered glucagon under this section, unless the act 356  
or omission constitutes willful or wanton misconduct: 357

(a) A community school; 358

(b) A member of a community school governing authority; 359

(c) A community school employee or contractor; 360

(d) A licensed health professional authorized to prescribe 361  
drugs who personally furnishes or prescribes injectable or 362  
nasally administered glucagon, provides a consultation, or 363  
issues a protocol pursuant to this section. 364

(2) This division does not eliminate, limit, or reduce any 365  
other immunity or defense that a community school or governing 366  
authority, member of a community school governing authority, 367  
community school employee or contractor, or licensed health 368  
professional may be entitled to under Chapter 2744. or any other 369  
provision of the Revised Code or under the common law of this 370  
state. 371

(C) A community school may accept donations of injectable 372  
or nasally administered glucagon from a wholesale distributor of 373  
dangerous drugs or a manufacturer of dangerous drugs, as defined 374  
in section 4729.01 of the Revised Code, and may accept donations 375  
of money from any person to purchase the drug. 376

(D) A community school that elects to procure injectable 377  
or nasally administered glucagon under this section shall report 378  
to the department of education each procurement and each 379  
occurrence in which a dose of the drug is used from the school's 380  
supply. 381

**Sec. 3326.60.** (A) With the approval of its governing body, 382  
a STEM school established under this chapter may procure 383  
injectable or nasally administered glucagon in the manner 384  
prescribed by section 3313.7115 of the Revised Code. A STEM 385  
school that elects to do so shall comply with all provisions of 386  
that section as if it were a school district. 387

(B) (1) The following are not liable in damages in a civil 388  
action for injury, death, or loss to person or property that 389

allegedly arises from an act or omission associated with 390  
procuring, maintaining, accessing, or using injectable or 391  
nasally administered glucagon under this section, unless the act 392  
or omission constitutes willful or wanton misconduct: 393

(a) A STEM school; 394

(b) A member of a STEM school governing body; 395

(c) A STEM school employee or contractor; 396

(d) A licensed health professional authorized to prescribe 397  
drugs who personally furnishes or prescribes injectable or 398  
nasally administered glucagon, provides a consultation, or 399  
issues a protocol pursuant to this section. 400

(2) This division does not eliminate, limit, or reduce any 401  
other immunity or defense that a STEM school or governing body, 402  
member of a STEM school governing body, STEM school employee or 403  
contractor, or licensed health professional may be entitled to 404  
under Chapter 2744. or any other provision of the Revised Code 405  
or under the common law of this state. 406

(C) A STEM school may accept donations of injectable or 407  
nasally administered glucagon from a wholesale distributor of 408  
dangerous drugs or a manufacturer of dangerous drugs, as defined 409  
in section 4729.01 of the Revised Code, and may accept donations 410  
of money from any person to purchase the drug. 411

(D) A STEM school that elects to procure injectable or 412  
nasally administered glucagon under this section shall report to 413  
the department of education each procurement and each occurrence 414  
in which a dose of the drug is used from the school's supply. 415

**Sec. 3328.38.** (A) With the approval of its board of 416  
trustees, a college-preparatory boarding school established 417

under this chapter may procure injectable or nasally 418  
administered glucagon in the manner prescribed by section 419  
3313.7115 of the Revised Code. A college-preparatory boarding 420  
school that elects to do so shall comply with all provisions of 421  
that section as if it were a school district. 422

(B) (1) The following are not liable in damages in a civil 423  
action for injury, death, or loss to person or property that 424  
allegedly arises from an act or omission associated with 425  
procuring, maintaining, accessing, or using injectable or 426  
nasally administered glucagon under this section, unless the act 427  
or omission constitutes willful or wanton misconduct: 428

(a) A college-preparatory boarding school; 429

(b) A member of a college-preparatory boarding school 430  
board of trustees; 431

(c) A college-preparatory boarding school employee or 432  
contractor; 433

(d) A licensed health professional authorized to prescribe 434  
drugs who personally furnishes or prescribes injectable or 435  
nasally administered glucagon, provides a consultation, or 436  
issues a protocol pursuant to this section. 437

(2) This division does not eliminate, limit, or reduce any 438  
other immunity or defense that a college-preparatory boarding 439  
school or board of trustees, member of a college-preparatory 440  
boarding school board of trustees, college-preparatory boarding 441  
school employee or contractor, or licensed health professional 442  
may be entitled to under Chapter 2744. or any other provision of 443  
the Revised Code or under the common law of this state. 444

(C) A college-preparatory boarding school may accept 445  
donations of injectable or nasally administered glucagon from a 446

wholesale distributor of dangerous drugs or a manufacturer of 447  
dangerous drugs, as defined in section 4729.01 of the Revised 448  
Code, and may accept donations of money from any person to 449  
purchase the drug. 450

(D) A college-preparatory boarding school that elects to 451  
procure injectable or nasally administered glucagon under this 452  
section shall report to the department of education each 453  
procurement and each occurrence in which a dose of the drug is 454  
used from the school's supply. 455

**Sec. 4723.484.** (A) (1) Subject to division (A) (2) of this 456  
section, and notwithstanding any provision of this chapter or 457  
rule adopted by the board of nursing, a clinical nurse 458  
specialist, certified nurse-midwife, or certified nurse 459  
practitioner licensed as an advanced practice registered nurse 460  
under Chapter 4723. of the Revised Code may do either of the 461  
following without having examined an individual to whom glucagon 462  
may be administered: 463

(a) Personally furnish a supply of injectable or nasally 464  
administered glucagon for use in accordance with sections 465  
3313.7115, 3313.7116, 3314.147, 3326.60, 3328.38, and 5101.78 of 466  
the Revised Code; 467

(b) Issue a prescription for injectable or nasally 468  
administered glucagon for use in accordance with sections 469  
3313.7115, 3313.7116, 3314.147, 3326.60, 3328.38, and 5101.78 of 470  
the Revised Code. 471

(2) Injectable or nasally administered glucagon personally 472  
furnished or prescribed under division (A) (1) of this section 473  
must be furnished or prescribed in such a manner that it may be 474  
administered only in a manufactured dosage form. 475

(B) A nurse who acts in good faith in accordance with this 476  
section is not liable for or subject to any of the following for 477  
any action or omission of an entity to which injectable or 478  
nasally administered glucagon is furnished or a prescription is 479  
issued: damages in any civil action, prosecution in any criminal 480  
proceeding, or professional disciplinary action. 481

**Sec. 4723.50.** (A) As used in this section: 482

(1) "Controlled substance" has the same meaning as in 483  
section 3719.01 of the Revised Code. 484

(2) "Medication-assisted treatment" has the same meaning 485  
as in section 340.01 of the Revised Code. 486

(B) In accordance with Chapter 119. of the Revised Code, 487  
the board of nursing shall adopt rules as necessary to implement 488  
the provisions of this chapter pertaining to the authority of 489  
advanced practice registered nurses who are designated as 490  
clinical nurse specialists, certified nurse-midwives, and 491  
certified nurse practitioners to prescribe and furnish drugs and 492  
therapeutic devices. 493

The board shall adopt rules that are consistent with a 494  
recommended exclusionary formulary the board receives from the 495  
committee on prescriptive governance pursuant to section 496  
4723.492 of the Revised Code. After reviewing a formulary 497  
submitted by the committee, the board may either adopt the 498  
formulary as a rule or ask the committee to reconsider and 499  
resubmit the formulary. The board shall not adopt any rule that 500  
does not conform to a formulary developed by the committee. 501

The exclusionary formulary shall permit, in a manner 502  
consistent with section 4723.481 of the Revised Code, the 503  
prescribing of controlled substances, including drugs that 504

contain buprenorphine used in medication-assisted treatment and 505  
both oral and long-acting opioid antagonists. The formulary 506  
shall not permit the prescribing or furnishing of any of the 507  
following: 508

(1) A drug or device to perform or induce an abortion; 509

(2) A drug or device prohibited by federal or state law. 510

(C) In addition to the rules described in division (B) of 511  
this section, the board shall adopt rules under this section 512  
that do the following: 513

(1) Establish standards for board approval of the course 514  
of study in advanced pharmacology and related topics required by 515  
section 4723.482 of the Revised Code; 516

(2) Establish requirements for board approval of the two- 517  
hour course of instruction in the laws of this state as required 518  
under division (C) (1) of section 4723.482 of the Revised Code 519  
~~and division (B) (2) of section 4723.484 of the Revised Code;~~ 520

(3) Establish criteria for the components of the standard 521  
care arrangements described in section 4723.431 of the Revised 522  
Code that apply to the authority to prescribe, including the 523  
components that apply to the authority to prescribe schedule II 524  
controlled substances. The rules shall be consistent with that 525  
section and include all of the following: 526

(a) Quality assurance standards; 527

(b) Standards for periodic review by a collaborating 528  
physician or podiatrist of the records of patients treated by 529  
the clinical nurse specialist, certified nurse-midwife, or 530  
certified nurse practitioner; 531

(c) Acceptable travel time between the location at which 532

the clinical nurse specialist, certified nurse-midwife, or 533  
certified nurse practitioner is engaging in the prescribing 534  
components of the nurse's practice and the location of the 535  
nurse's collaborating physician or podiatrist; 536

(d) Any other criteria recommended by the committee on 537  
prescriptive governance. 538

**Sec. 4729.01.** As used in this chapter: 539

(A) "Pharmacy," except when used in a context that refers 540  
to the practice of pharmacy, means any area, room, rooms, place 541  
of business, department, or portion of any of the foregoing 542  
where the practice of pharmacy is conducted. 543

(B) "Practice of pharmacy" means providing pharmacist care 544  
requiring specialized knowledge, judgment, and skill derived 545  
from the principles of biological, chemical, behavioral, social, 546  
pharmaceutical, and clinical sciences. As used in this division, 547  
"pharmacist care" includes the following: 548

(1) Interpreting prescriptions; 549

(2) Dispensing drugs and drug therapy related devices; 550

(3) Compounding drugs; 551

(4) Counseling individuals with regard to their drug 552  
therapy, recommending drug therapy related devices, and 553  
assisting in the selection of drugs and appliances for treatment 554  
of common diseases and injuries and providing instruction in the 555  
proper use of the drugs and appliances; 556

(5) Performing drug regimen reviews with individuals by 557  
discussing all of the drugs that the individual is taking and 558  
explaining the interactions of the drugs; 559

(6) Performing drug utilization reviews with licensed health professionals authorized to prescribe drugs when the pharmacist determines that an individual with a prescription has a drug regimen that warrants additional discussion with the prescriber;

(7) Advising an individual and the health care professionals treating an individual with regard to the individual's drug therapy;

(8) Acting pursuant to a consult agreement with one or more physicians authorized under Chapter 4731. of the Revised Code to practice medicine and surgery or osteopathic medicine and surgery, if an agreement has been established;

(9) Engaging in the administration of immunizations to the extent authorized by section 4729.41 of the Revised Code;

(10) Engaging in the administration of drugs to the extent authorized by section 4729.45 of the Revised Code.

(C) "Compounding" means the preparation, mixing, assembling, packaging, and labeling of one or more drugs in any of the following circumstances:

(1) Pursuant to a prescription issued by a licensed health professional authorized to prescribe drugs;

(2) Pursuant to the modification of a prescription made in accordance with a consult agreement;

(3) As an incident to research, teaching activities, or chemical analysis;

(4) In anticipation of orders for drugs pursuant to prescriptions, based on routine, regularly observed dispensing patterns;

(5) Pursuant to a request made by a licensed health professional authorized to prescribe drugs for a drug that is to be used by the professional for the purpose of direct administration to patients in the course of the professional's practice, if all of the following apply:

(a) At the time the request is made, the drug is not commercially available regardless of the reason that the drug is not available, including the absence of a manufacturer for the drug or the lack of a readily available supply of the drug from a manufacturer.

(b) A limited quantity of the drug is compounded and provided to the professional.

(c) The drug is compounded and provided to the professional as an occasional exception to the normal practice of dispensing drugs pursuant to patient-specific prescriptions.

(D) "Consult agreement" means an agreement that has been entered into under section 4729.39 of the Revised Code.

(E) "Drug" means:

(1) Any article recognized in the United States pharmacopoeia and national formulary, or any supplement to them, intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in humans or animals;

(2) Any other article intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in humans or animals;

(3) Any article, other than food, intended to affect the structure or any function of the body of humans or animals;

(4) Any article intended for use as a component of any

article specified in division (E) (1), (2), or (3) of this 616  
section; but does not include devices or their components, 617  
parts, or accessories. 618

(F) "Dangerous drug" means any of the following: 619

(1) Any drug to which either of the following applies: 620

(a) Under the "Federal Food, Drug, and Cosmetic Act," 52 621  
Stat. 1040 (1938), 21 U.S.C.A. 301, as amended, the drug is 622  
required to bear a label containing the legend "Caution: Federal 623  
law prohibits dispensing without prescription" or "Caution: 624  
Federal law restricts this drug to use by or on the order of a 625  
licensed veterinarian" or any similar restrictive statement, or 626  
the drug may be dispensed only upon a prescription; 627

(b) Under Chapter 3715. or 3719. of the Revised Code, the 628  
drug may be dispensed only upon a prescription. 629

(2) Any drug that contains a schedule V controlled 630  
substance and that is exempt from Chapter 3719. of the Revised 631  
Code or to which that chapter does not apply; 632

(3) Any drug intended for administration by injection into 633  
the human body other than through a natural orifice of the human 634  
body; 635

(4) Any drug that is a biological product, as defined in 636  
section 3715.01 of the Revised Code. 637

(G) "Federal drug abuse control laws" has the same meaning 638  
as in section 3719.01 of the Revised Code. 639

(H) "Prescription" means all of the following: 640

(1) A written, electronic, or oral order for drugs or 641  
combinations or mixtures of drugs to be used by a particular 642

individual or for treating a particular animal, issued by a 643  
licensed health professional authorized to prescribe drugs; 644

(2) For purposes of sections 2925.61, 4723.488, 4730.431, 645  
and 4731.94 of the Revised Code, a written, electronic, or oral 646  
order for naloxone issued to and in the name of a family member, 647  
friend, or other individual in a position to assist an 648  
individual who there is reason to believe is at risk of 649  
experiencing an opioid-related overdose. 650

(3) For purposes of section 4729.44 of the Revised Code, a 651  
written, electronic, or oral order for naloxone issued to and in 652  
the name of either of the following: 653

(a) An individual who there is reason to believe is at 654  
risk of experiencing an opioid-related overdose; 655

(b) A family member, friend, or other individual in a 656  
position to assist an individual who there is reason to believe 657  
is at risk of experiencing an opioid-related overdose. 658

(4) For purposes of sections 4723.4810, 4729.282, 659  
4730.432, and 4731.93 of the Revised Code, a written, 660  
electronic, or oral order for a drug to treat chlamydia, 661  
gonorrhoea, or trichomoniasis issued to and in the name of a 662  
patient who is not the intended user of the drug but is the 663  
sexual partner of the intended user; 664

(5) For purposes of sections 3313.7110, 3313.7111, 665  
3314.143, 3326.28, 3328.29, 4723.483, 4729.88, 4730.433, 666  
4731.96, and 5101.76 of the Revised Code, a written, electronic, 667  
or oral order for an epinephrine autoinjector issued to and in 668  
the name of a school, school district, or camp; 669

(6) For purposes of Chapter 3728. and sections 4723.483, 670  
4729.88, 4730.433, and 4731.96 of the Revised Code, a written, 671

electronic, or oral order for an epinephrine autoinjector issued 672  
to and in the name of a qualified entity, as defined in section 673  
3728.01 of the Revised Code; 674

(7) For purposes of sections 3313.7115, 3313.7116, 675  
3314.147, 3326.60, 3328.38, 4723.484, 4730.434, 4731.92, and 676  
5101.78 of the Revised Code, a written, electronic, or oral 677  
order for injectable or nasally administered glucagon in the 678  
name of a school, school district, or camp. 679

(I) "Licensed health professional authorized to prescribe 680  
drugs" or "prescriber" means an individual who is authorized by 681  
law to prescribe drugs or dangerous drugs or drug therapy 682  
related devices in the course of the individual's professional 683  
practice, including only the following: 684

(1) A dentist licensed under Chapter 4715. of the Revised 685  
Code; 686

(2) A clinical nurse specialist, certified nurse-midwife, 687  
or certified nurse practitioner who holds a current, valid 688  
license to practice nursing as an advanced practice registered 689  
nurse issued under Chapter 4723. of the Revised Code; 690

(3) An optometrist licensed under Chapter 4725. of the 691  
Revised Code to practice optometry under a therapeutic 692  
pharmaceutical agents certificate; 693

(4) A physician authorized under Chapter 4731. of the 694  
Revised Code to practice medicine and surgery, osteopathic 695  
medicine and surgery, or podiatric medicine and surgery; 696

(5) A physician assistant who holds a license to practice 697  
as a physician assistant issued under Chapter 4730. of the 698  
Revised Code, holds a valid prescriber number issued by the 699  
state medical board, and has been granted physician-delegated 700

prescriptive authority; 701

(6) A veterinarian licensed under Chapter 4741. of the 702  
Revised Code. 703

(J) "Sale" or "sell" includes any transaction made by any 704  
person, whether as principal proprietor, agent, or employee, to 705  
do or offer to do any of the following: deliver, distribute, 706  
broker, exchange, gift or otherwise give away, or transfer, 707  
whether the transfer is by passage of title, physical movement, 708  
or both. 709

(K) "Wholesale sale" and "sale at wholesale" mean any sale 710  
in which the purpose of the purchaser is to resell the article 711  
purchased or received by the purchaser. 712

(L) "Retail sale" and "sale at retail" mean any sale other 713  
than a wholesale sale or sale at wholesale. 714

(M) "Retail seller" means any person that sells any 715  
dangerous drug to consumers without assuming control over and 716  
responsibility for its administration. Mere advice or 717  
instructions regarding administration do not constitute control 718  
or establish responsibility. 719

(N) "Price information" means the price charged for a 720  
prescription for a particular drug product and, in an easily 721  
understandable manner, all of the following: 722

(1) The proprietary name of the drug product; 723

(2) The established (generic) name of the drug product; 724

(3) The strength of the drug product if the product 725  
contains a single active ingredient or if the drug product 726  
contains more than one active ingredient and a relevant strength 727  
can be associated with the product without indicating each 728

active ingredient. The established name and quantity of each 729  
active ingredient are required if such a relevant strength 730  
cannot be so associated with a drug product containing more than 731  
one ingredient. 732

(4) The dosage form; 733

(5) The price charged for a specific quantity of the drug 734  
product. The stated price shall include all charges to the 735  
consumer, including, but not limited to, the cost of the drug 736  
product, professional fees, handling fees, if any, and a 737  
statement identifying professional services routinely furnished 738  
by the pharmacy. Any mailing fees and delivery fees may be 739  
stated separately without repetition. The information shall not 740  
be false or misleading. 741

(O) "Wholesale distributor of dangerous drugs" or 742  
"wholesale distributor" means a person engaged in the sale of 743  
dangerous drugs at wholesale and includes any agent or employee 744  
of such a person authorized by the person to engage in the sale 745  
of dangerous drugs at wholesale. 746

(P) "Manufacturer of dangerous drugs" or "manufacturer" 747  
means a person, other than a pharmacist or prescriber, who 748  
manufactures dangerous drugs and who is engaged in the sale of 749  
those dangerous drugs. 750

(Q) "Terminal distributor of dangerous drugs" or "terminal 751  
distributor" means a person who is engaged in the sale of 752  
dangerous drugs at retail, or any person, other than a 753  
manufacturer, repackager, outsourcing facility, third-party 754  
logistics provider, wholesale distributor, or pharmacist, who 755  
has possession, custody, or control of dangerous drugs for any 756  
purpose other than for that person's own use and consumption. 757

"Terminal distributor" includes pharmacies, hospitals, nursing homes, and laboratories and all other persons who procure dangerous drugs for sale or other distribution by or under the supervision of a pharmacist, licensed health professional authorized to prescribe drugs, or other person authorized by the state board of pharmacy.

(R) "Promote to the public" means disseminating a representation to the public in any manner or by any means, other than by labeling, for the purpose of inducing, or that is likely to induce, directly or indirectly, the purchase of a dangerous drug at retail.

(S) "Person" includes any individual, partnership, association, limited liability company, or corporation, the state, any political subdivision of the state, and any district, department, or agency of the state or its political subdivisions.

(T) "Animal shelter" means a facility operated by a humane society or any society organized under Chapter 1717. of the Revised Code or a dog pound operated pursuant to Chapter 955. of the Revised Code.

(U) "Food" has the same meaning as in section 3715.01 of the Revised Code.

(V) "Pain management clinic" has the same meaning as in section 4731.054 of the Revised Code.

(W) "Investigational drug or product" means a drug or product that has successfully completed phase one of the United States food and drug administration clinical trials and remains under clinical trial, but has not been approved for general use by the United States food and drug administration.

"Investigational drug or product" does not include controlled 787  
substances in schedule I, as defined in section 3719.01 of the 788  
Revised Code. 789

(X) "Product," when used in reference to an 790  
investigational drug or product, means a biological product, 791  
other than a drug, that is made from a natural human, animal, or 792  
microorganism source and is intended to treat a disease or 793  
medical condition. 794

(Y) "Third-party logistics provider" means a person that 795  
provides or coordinates warehousing or other logistics services 796  
pertaining to dangerous drugs including distribution, on behalf 797  
of a manufacturer, wholesale distributor, or terminal 798  
distributor of dangerous drugs, but does not take ownership of 799  
the drugs or have responsibility to direct the sale or 800  
disposition of the drugs. 801

(Z) "Repackager of dangerous drugs" or "repackager" means 802  
a person that repacks and relabels dangerous drugs for sale or 803  
distribution. 804

(AA) "Outsourcing facility" means a facility that is 805  
engaged in the compounding and sale of sterile drugs and is 806  
registered as an outsourcing facility with the United States 807  
food and drug administration. 808

(BB) "Laboratory" means a laboratory licensed under this 809  
chapter as a terminal distributor of dangerous drugs and 810  
entrusted to have custody of any of the following drugs and to 811  
use the drugs for scientific and clinical purposes and for 812  
purposes of instruction: dangerous drugs that are not controlled 813  
substances, as defined in section 3719.01 of the Revised Code; 814  
dangerous drugs that are controlled substances, as defined in 815

that section; and controlled substances in schedule I, as 816  
defined in that section. 817

**Sec. 4729.51.** (A) No person other than a licensed 818  
manufacturer of dangerous drugs, outsourcing facility, third- 819  
party logistics provider, repackager of dangerous drugs, or 820  
wholesale distributor of dangerous drugs shall possess for sale, 821  
sell, distribute, or deliver, at wholesale, dangerous drugs or 822  
investigational drugs or products, except as follows: 823

(1) A licensed terminal distributor of dangerous drugs 824  
that is a pharmacy may make occasional sales of dangerous drugs 825  
or investigational drugs or products at wholesale. 826

(2) A licensed terminal distributor of dangerous drugs 827  
having more than one licensed location may transfer or deliver 828  
dangerous drugs from one licensed location to another licensed 829  
location owned by the terminal distributor if the license issued 830  
for each location is in effect at the time of the transfer or 831  
delivery. 832

(3) A licensed terminal distributor of dangerous drugs 833  
that is not a pharmacy may make occasional sales of naloxone at 834  
wholesale. 835

(4) A licensed terminal distributor of dangerous drugs 836  
that is not a pharmacy may make occasional sales of dangerous 837  
drugs at wholesale if the drugs being sold are in shortage, as 838  
defined in rules adopted by the state board of pharmacy under 839  
section 4729.26 of the Revised Code. 840

(B) No licensed manufacturer, outsourcing facility, third- 841  
party logistics provider, repackager, or wholesale distributor 842  
shall possess for sale, sell, or distribute, at wholesale, 843  
dangerous drugs or investigational drugs or products to any 844

person other than the following: 845

(1) Subject to division (D) of this section, a licensed 846  
terminal distributor of dangerous drugs; 847

(2) Subject to division (C) of this section, any person 848  
exempt from licensure as a terminal distributor of dangerous 849  
drugs under section 4729.541 of the Revised Code; 850

(3) A licensed manufacturer, outsourcing facility, third- 851  
party logistics provider, repackager, or wholesale distributor; 852

(4) A terminal distributor, manufacturer, outsourcing 853  
facility, third-party logistics provider, repackager, or 854  
wholesale distributor that is located in another state, is not 855  
engaged in the sale of dangerous drugs within this state, and is 856  
actively licensed to engage in the sale of dangerous drugs by 857  
the state in which the distributor conducts business. 858

(C) No licensed manufacturer, outsourcing facility, third- 859  
party logistics provider, repackager, or wholesale distributor 860  
shall possess for sale, sell, or distribute, at wholesale, 861  
dangerous drugs or investigational drugs or products to either 862  
of the following: 863

(1) A prescriber who is employed by either of the 864  
following: 865

(a) A pain management clinic that is not licensed as a 866  
terminal distributor of dangerous drugs with a pain management 867  
clinic classification issued under section 4729.552 of the 868  
Revised Code; 869

(b) A facility, clinic, or other location that provides 870  
office-based opioid treatment but is not licensed as a terminal 871  
distributor of dangerous drugs with an office-based opioid 872

treatment classification issued under section 4729.553 of the Revised Code if such a license is required by that section.

(2) A business entity described in division (A)(2) or (3) of section 4729.541 of the Revised Code that is, or is operating, either of the following:

(a) A pain management clinic without a license as a terminal distributor of dangerous drugs with a pain management clinic classification issued under section 4729.552 of the Revised Code;

(b) A facility, clinic, or other location that provides office-based opioid treatment without a license as a terminal distributor of dangerous drugs with an office-based opioid treatment classification issued under section 4729.553 of the Revised Code if such a license is required by that section.

(D) No licensed manufacturer, outsourcing facility, third-party logistics provider, repackager, or wholesale distributor shall possess dangerous drugs or investigational drugs or products for sale at wholesale, or sell or distribute such drugs at wholesale, to a licensed terminal distributor of dangerous drugs, except as follows:

(1) In the case of a terminal distributor with a category II license, only dangerous drugs in category II, as defined in division (A)(1) of section 4729.54 of the Revised Code;

(2) In the case of a terminal distributor with a category III license, dangerous drugs in category II and category III, as defined in divisions (A)(1) and (2) of section 4729.54 of the Revised Code;

(3) In the case of a terminal distributor with a limited category II or III license, only the dangerous drugs specified

in the license.	902
(E) (1) Except as provided in division (E) (2) of this section, no person shall do any of the following:	903 904
(a) Sell or distribute, at retail, dangerous drugs;	905
(b) Possess for sale, at retail, dangerous drugs;	906
(c) Possess dangerous drugs.	907
(2) (a) Divisions (E) (1) (a), (b), and (c) of this section do not apply to any of the following:	908 909
(i) A licensed terminal distributor of dangerous drugs;	910
(ii) A person who possesses, or possesses for sale or sells, at retail, a dangerous drug in accordance with Chapters 3719., 4715., 4723., 4725., 4729., 4730., 4731., and 4741. of the Revised Code;	911 912 913 914
(iii) Any of the persons identified in divisions (A) (1) to (5) and (13) of section 4729.541 of the Revised Code, but only to the extent specified in that section.	915 916 917
(b) Division (E) (1) (c) of this section does not apply to any of the following:	918 919
(i) A licensed manufacturer, outsourcing facility, third-party logistics provider, repackager, or wholesale distributor;	920 921
(ii) Any of the persons identified in divisions (A) (6) to (12) of section 4729.541 of the Revised Code, but only to the extent specified in that section.	922 923 924
(F) No licensed terminal distributor of dangerous drugs or person that is exempt from licensure under section 4729.541 of the Revised Code shall purchase dangerous drugs or investigational drugs or products from any person other than a	925 926 927 928

licensed manufacturer, outsourcing facility, third-party 929  
logistics provider, repackager, or wholesale distributor, except 930  
as follows: 931

(1) A licensed terminal distributor of dangerous drugs or 932  
person that is exempt from licensure under section 4729.541 of 933  
the Revised Code may make occasional purchases of dangerous 934  
drugs or investigational drugs or products that are sold in 935  
accordance with division (A) (1) or (3) of this section. 936

(2) A licensed terminal distributor of dangerous drugs 937  
having more than one licensed location may transfer or deliver 938  
dangerous drugs or investigational drugs or products from one 939  
licensed location to another licensed location if the license 940  
issued for each location is in effect at the time of the 941  
transfer or delivery. 942

(G) No licensed terminal distributor of dangerous drugs 943  
shall engage in the retail sale or other distribution of 944  
dangerous drugs or investigational drugs or products or maintain 945  
possession, custody, or control of dangerous drugs or 946  
investigational drugs or products for any purpose other than the 947  
distributor's personal use or consumption, at any establishment 948  
or place other than that or those described in the license 949  
issued by the board to such terminal distributor. 950

(H) Nothing in this section shall be construed to 951  
interfere with the performance of official duties by any law 952  
enforcement official authorized by municipal, county, state, or 953  
federal law to collect samples of any drug, regardless of its 954  
nature or in whose possession it may be. 955

(I) Notwithstanding anything to the contrary in this 956  
section, the board of education of a city, local, exempted 957

village, or joint vocational school district may distribute 958  
epinephrine autoinjectors for use in accordance with section 959  
3313.7110 of the Revised Code ~~and,~~ may distribute inhalers for 960  
use in accordance with section 3313.7113 of the Revised Code, 961  
and may distribute injectable or nasally administered glucagon 962  
for use in accordance with section 3313.7115 of the Revised 963  
Code. 964

**Sec. 4729.513.** A manufacturer of dangerous drugs may 965  
donate inhalers, as defined in section 3313.7113 of the Revised 966  
Code, ~~and epinephrine autoinjectors,~~ or injectable or nasally 967  
administered glucagon to any of the following: 968

(A) The board of education of a city, local, exempted 969  
village, or joint vocational school district; 970

(B) A community school established under Chapter 3314. of 971  
the Revised Code; 972

(C) A STEM school established under Chapter 3326. of the 973  
Revised Code; 974

(D) A college-preparatory boarding school established 975  
under Chapter 3328. of the Revised Code; 976

(E) A chartered or nonchartered nonpublic school; 977

(F) A residential camp, as defined in section 2151.011 of 978  
the Revised Code; 979

(G) A child day camp, as defined in section 5104.01 of the 980  
Revised Code; 981

(H) A child day camp operated by any county, township, 982  
municipal corporation, township park district created under 983  
section 511.18 of the Revised Code, park district created under 984  
section 1545.04 of the Revised Code, or joint recreation 985

<u>district established under section 755.14 of the Revised Code.</u>	986
<b>Sec. 4729.541.</b> (A) Except as provided in divisions (B) to	987
(D) of this section, all of the following are exempt from	988
licensure as a terminal distributor of dangerous drugs:	989
(1) A licensed health professional authorized to prescribe	990
drugs;	991
(2) A business entity that is a corporation formed under	992
division (B) of section 1701.03 of the Revised Code, a limited	993
liability company formed under Chapter 1705. of the Revised	994
Code, or a professional association formed under Chapter 1785.	995
of the Revised Code if the entity has a sole shareholder who is	996
a prescriber and is authorized to provide the professional	997
services being offered by the entity;	998
(3) A business entity that is a corporation formed under	999
division (B) of section 1701.03 of the Revised Code, a limited	1000
liability company formed under Chapter 1705. of the Revised	1001
Code, a partnership or a limited liability partnership formed	1002
under Chapter 1775. of the Revised Code, or a professional	1003
association formed under Chapter 1785. of the Revised Code, if,	1004
to be a shareholder, member, or partner, an individual is	1005
required to be licensed, certified, or otherwise legally	1006
authorized under Title XLVII of the Revised Code to perform the	1007
professional service provided by the entity and each such	1008
individual is a prescriber;	1009
(4) An individual who holds a current license,	1010
certificate, or registration issued under Title XLVII of the	1011
Revised Code and has been certified to conduct diabetes	1012
education by a national certifying body specified in rules	1013
adopted by the state board of pharmacy under section 4729.68 of	1014

the Revised Code, but only with respect to insulin that will be 1015  
used for the purpose of diabetes education and only if diabetes 1016  
education is within the individual's scope of practice under 1017  
statutes and rules regulating the individual's profession; 1018

(5) An individual who holds a valid certificate issued by 1019  
a nationally recognized S.C.U.B.A. diving certifying 1020  
organization approved by the state board of pharmacy under rules 1021  
adopted by the board, but only with respect to medical oxygen 1022  
that will be used for the purpose of emergency care or treatment 1023  
at the scene of a diving emergency; 1024

(6) With respect to epinephrine autoinjectors that may be 1025  
possessed under section 3313.7110, 3313.7111, 3314.143, 3326.28, 1026  
or 3328.29 of the Revised Code, any of the following: the board 1027  
of education of a city, local, exempted village, or joint 1028  
vocational school district; a chartered or nonchartered 1029  
nonpublic school; a community school established under Chapter 1030  
3314. of the Revised Code; a STEM school established under 1031  
Chapter 3326. of the Revised Code; or a college-preparatory 1032  
boarding school established under Chapter 3328. of the Revised 1033  
Code; 1034

(7) With respect to epinephrine autoinjectors that may be 1035  
possessed under section 5101.76 of the Revised Code, any of the 1036  
following: a residential camp, as defined in section 2151.011 of 1037  
the Revised Code; a child day camp, as defined in section 1038  
5104.01 of the Revised Code; or a child day camp operated by any 1039  
county, township, municipal corporation, township park district 1040  
created under section 511.18 of the Revised Code, park district 1041  
created under section 1545.04 of the Revised Code, or joint 1042  
recreation district established under section 755.14 of the 1043  
Revised Code; 1044

(8) With respect to epinephrine autoinjectors that may be 1045  
possessed under Chapter 3728. of the Revised Code, a qualified 1046  
entity, as defined in section 3728.01 of the Revised Code; 1047

(9) With respect to inhalers that may be possessed under 1048  
section 3313.7113, 3313.7114, 3314.144, 3326.30, or 3328.30 of 1049  
the Revised Code, any of the following: the board of education 1050  
of a city, local, exempted village, or joint vocational school 1051  
district; a chartered or nonchartered nonpublic school; a 1052  
community school established under Chapter 3314. of the Revised 1053  
Code; a STEM school established under Chapter 3326. of the 1054  
Revised Code; or a college-preparatory boarding school 1055  
established under Chapter 3328. of the Revised Code; 1056

(10) With respect to inhalers that may be possessed under 1057  
section 5101.77 of the Revised Code, any of the following: a 1058  
residential camp, as defined in section 2151.011 of the Revised 1059  
Code; a child day camp, as defined in section 5104.01 of the 1060  
Revised Code; or a child day camp operated by any county, 1061  
township, municipal corporation, township park district created 1062  
under section 511.18 of the Revised Code, park district created 1063  
under section 1545.04 of the Revised Code, or joint recreation 1064  
district established under section 755.14 of the Revised Code; 1065

(11) With respect to naloxone that may be possessed under 1066  
section 2925.61 of the Revised Code, a law enforcement agency 1067  
and its peace officers; 1068

(12) With respect to naloxone that may be possessed under 1069  
section 4729.514 of the Revised Code, a service entity, as 1070  
defined in that section; 1071

(13) A facility that is owned and operated by the United 1072  
States department of defense, the United States department of 1073

veterans affairs, or any other federal agency; 1074

(14) With respect to injectable or nasally administered 1075  
glucagon that may be possessed under sections 3313.7115, 1076  
3313.7116, 3314.147, 3326.60, and 3328.38 of the Revised Code, 1077  
any of the following: the board of education of a city, local, 1078  
exempted village, or joint vocational school district; a 1079  
chartered or nonchartered nonpublic school; a community school 1080  
established under Chapter 3314. of the Revised Code; a STEM 1081  
school established under Chapter 3326. of the Revised Code; or a 1082  
college-preparatory boarding school established under Chapter 1083  
3328. of the Revised Code; 1084

(15) With respect to injectable or nasally administered 1085  
glucagon that may be possessed under section 5101.78 of the 1086  
Revised Code, any of the following: a residential camp, as 1087  
defined in section 2151.011 of the Revised Code; a child day 1088  
camp, as defined in section 5104.01 of the Revised Code; or a 1089  
child day camp operated by any county, township, municipal 1090  
corporation, township park district created under section 511.18 1091  
of the Revised Code, park district created under section 1545.04 1092  
of the Revised Code, or joint recreation district established 1093  
under section 755.14 of the Revised Code. 1094

(B) If a person described in division (A) of this section 1095  
is a pain management clinic or is operating a pain management 1096  
clinic, the person shall hold a license as a terminal 1097  
distributor of dangerous drugs with a pain management clinic 1098  
classification issued under section 4729.552 of the Revised 1099  
Code. 1100

(C) If a person described in division (A) of this section 1101  
is operating a facility, clinic, or other location described in 1102  
division (B) of section 4729.553 of the Revised Code that must 1103

hold a category III terminal distributor of dangerous drugs 1104  
license with an office-based opioid treatment classification, 1105  
the person shall hold a license with that classification. 1106

(D) Any of the persons described in divisions (A) (1) to 1107  
(12) of this section shall hold a license as a terminal 1108  
distributor of dangerous drugs in order to possess, have custody 1109  
or control of, and distribute any of the following: 1110

(1) Dangerous drugs that are compounded or used for the 1111  
purpose of compounding; 1112

(2) A schedule I, II, III, IV, or V controlled substance, 1113  
as defined in section 3719.01 of the Revised Code. 1114

**Sec. 4729.60.** (A) (1) Before a licensee identified in 1115  
division (B) (1) (a) of section 4729.52 of the Revised Code may 1116  
sell or distribute dangerous drugs at wholesale to any person, 1117  
except as provided in division (A) (2) of this section, the 1118  
licensee shall query the roster established pursuant to section 1119  
4729.59 of the Revised Code to determine whether the purchaser 1120  
is a licensed terminal distributor of dangerous drugs. 1121

If no documented query is conducted before a sale is made, 1122  
it shall be presumed that the sale of dangerous drugs by the 1123  
licensee is in violation of division (B) of section 4729.51 of 1124  
the Revised Code and the purchase of dangerous drugs by the 1125  
purchaser is in violation of division (E) of section 4729.51 of 1126  
the Revised Code. If a licensee conducts a documented query and 1127  
relies on the results of the query in selling or distributing 1128  
dangerous drugs at wholesale to the terminal distributor of 1129  
dangerous drugs, the licensee shall be deemed not to have 1130  
violated division (B) of section 4729.51 of the Revised Code in 1131  
making the sale. 1132

(2) Division (A) (1) of this section does not apply when a  
licensee identified in division (B) (1) (a) of section 4729.52 of  
the Revised Code sells or distributes dangerous drugs at  
wholesale to any of the following:

(a) A person specified in division (B) (4) of section  
4729.51 of the Revised Code;

(b) Any of the persons described in divisions (A) (1) to  
~~(13)~~ (15) of section 4729.541 of the Revised Code, but only if  
the purchaser is not required to obtain licensure as provided in  
divisions (B) to (D) of that section.

(B) Before a licensed terminal distributor of dangerous  
drugs may purchase dangerous drugs at wholesale, the terminal  
distributor shall query the roster established pursuant to  
section 4729.59 of the Revised Code to confirm the seller is  
licensed to engage in the sale or distribution of dangerous  
drugs at wholesale.

If no documented query is conducted before a purchase is  
made, it shall be presumed that the purchase of dangerous drugs  
by the terminal distributor is in violation of division (F) of  
section 4729.51 of the Revised Code and the sale of dangerous  
drugs by the seller is in violation of division (A) of section  
4729.51 of the Revised Code. If a licensed terminal distributor  
of dangerous drugs conducts a documented query at least annually  
and relies on the results of the query in purchasing dangerous  
drugs at wholesale, the terminal distributor shall be deemed not  
to have violated division (F) of section 4729.51 of the Revised  
Code in making the purchase.

**Sec. 4729.88.** (A) Notwithstanding any provision of this  
chapter or rule adopted by the state board of pharmacy, a

pharmacist may dispense epinephrine autoinjectors pursuant to a 1162  
prescription issued under section 4723.483, 4730.433, or 4731.96 1163  
of the Revised Code. 1164

A pharmacist who in good faith dispenses epinephrine 1165  
autoinjectors under this ~~section~~ division is not liable for or 1166  
subject to any of the following for any action or omission of an 1167  
entity to which an epinephrine autoinjector is dispensed: 1168  
damages in any civil action, prosecution in any criminal 1169  
proceeding, or professional disciplinary action. 1170

(B) Notwithstanding any provision of this chapter or rule 1171  
adopted by the state board of pharmacy, a pharmacist may 1172  
dispense injectable or nasally administered glucagon pursuant to 1173  
a prescription issued under section 4723.484, 4730.434, or 1174  
4731.92 of the Revised Code. 1175

A pharmacist who in good faith dispenses injectable or 1176  
nasally administered glucagon under this division is not liable 1177  
for or subject to any of the following for any action or 1178  
omission of an entity to which the drug is dispensed: damages in 1179  
any civil action, prosecution in any criminal proceeding, or 1180  
professional disciplinary action. 1181

**Sec. 4730.434.** (A) (1) Subject to division (A) (2) of this 1182  
section and notwithstanding any provision of this chapter or 1183  
rule adopted by the state medical board, a physician assistant 1184  
who holds a valid prescriber number issued by the board and has 1185  
been granted physician-delegated prescriptive authority may do 1186  
either of the following without having examined an individual to 1187  
whom glucagon may be administered: 1188

(a) Personally furnish a supply of injectable or nasally 1189  
administered glucagon for use in accordance with section 1190

3313.7115, 3313.7116, 3314.147, 3326.60, 3328.38, or 5101.78 of 1191  
the Revised Code; 1192

(b) Issue a prescription for injectable or nasally 1193  
administered glucagon in accordance with section 3313.7115, 1194  
3313.7116, 3314.147, 3326.60, 3328.38, or 5101.78 of the Revised 1195  
Code. 1196

(2) Injectable or nasally administered glucagon personally 1197  
furnished or prescribed under division (A)(1) of this section 1198  
must be furnished or prescribed in such a manner that it may be 1199  
administered only in a manufactured dosage form. 1200

(B) A physician assistant who acts in good faith in 1201  
accordance with this section is not liable for or subject to any 1202  
of the following for any action or omission of an entity to 1203  
which injectable or nasally administered glucagon is furnished 1204  
or a prescription is issued: damages in any civil action, 1205  
prosecution in any criminal proceeding, or professional 1206  
disciplinary action. 1207

**Sec. 4731.92.** (A) As used in this section, "physician" 1208  
means an individual authorized under this chapter to practice 1209  
medicine and surgery, osteopathic medicine and surgery, or 1210  
podiatric medicine and surgery. 1211

(B) (1) Subject to division (B) (2) of this section, and 1212  
notwithstanding any provision of this chapter or rule adopted by 1213  
the state medical board, a physician may do either of the 1214  
following without having examined an individual to whom glucagon 1215  
may be administered: 1216

(a) Personally furnish a supply of injectable or nasally 1217  
administered glucagon for use in accordance with section 1218  
3313.7115, 3313.7116, 3314.147, 3326.60, 3328.38, or 5101.78 of 1219

Revised Code; 1220

(b) Issue a prescription for injectable or nasally administered glucagon for use in accordance with section 3313.7115, 3313.7116, 3314.147, 3326.60, 3328.38, or 5101.78 of the Revised Code. 1221  
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(2) Injectable or nasally administered glucagon personally furnished or prescribed under division (B)(1) of this section must be furnished or prescribed in such a manner that it may be administered only in a manufactured dosage form. 1225  
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(C) A physician who acts in good faith in accordance with this section is not liable for or subject to any of the following for any action or omission of an entity to which injectable or nasally administered glucagon is furnished or a prescription is issued: damages in any civil action, prosecution in any criminal proceeding, or professional disciplinary action. 1229  
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**Sec. 5101.78.** (A) As used in this section, "licensed health professional authorized to prescribe drugs" and "prescriber" have the same meanings as in section 4729.01 of the Revised Code. 1235  
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(B) A residential camp, as defined in section 2151.011 of the Revised Code; a child day camp, as defined in section 5104.01 of the Revised Code; or a child day camp operated by any county, township, municipal corporation, township park district created under section 511.18 of the Revised Code, park district created under section 1545.04 of the Revised Code, or joint recreation district established under section 755.14 of the Revised Code may procure injectable or nasally administered glucagon for use in emergency situations identified under division (D)(5) of this section by doing one of the following: 1239  
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(1) Having a licensed health professional authorized to 1249  
prescribe drugs, acting in accordance with section 4723.484, 1250  
4730.434, or 4731.92 of the Revised Code, personally furnish the 1251  
injectable or nasally administered glucagon to the camp or issue 1252  
a prescription for the drug in the name of the camp; 1253

(2) Obtaining a prescriber-issued protocol that includes 1254  
definitive orders for injectable or nasally administered 1255  
glucagon and the dosages to be administered; 1256

A camp that elects to procure injectable or nasally 1257  
administered glucagon under this section is encouraged to 1258  
maintain at least two doses of the drug at all times. 1259

(C) A camp that elects to procure injectable or nasally 1260  
administered glucagon under this section shall adopt a policy 1261  
governing maintenance and use of the drug. Before adopting the 1262  
policy, the camp shall consult with a licensed health 1263  
professional authorized to prescribe drugs. 1264

(D) The policy adopted under division (C) of this section 1265  
shall do all of the following: 1266

(1) Identify the one or more locations at the camp in 1267  
which injectable or nasally administered glucagon must be 1268  
stored; 1269

(2) Specify the conditions under which injectable or 1270  
nasally administered glucagon must be stored, replaced, or 1271  
disposed; 1272

(3) Specify the individuals employed by or under contract 1273  
with the camp, or who volunteer at the camp, who may access and 1274  
use injectable or nasally administered glucagon in an emergency 1275  
situation identified under division (D) (5) of this section; 1276

(4) Specify any training that employees, contractors, or 1277  
volunteers specified under division (D) (3) of this section must 1278  
complete before being authorized to access and use injectable or 1279  
nasally administered glucagon; 1280

(5) Identify the emergency situations, including when an 1281  
individual exhibits signs and symptoms of severe hypoglycemia, 1282  
in which employees, contractors, or volunteers specified under 1283  
division (D) (3) of this section may access and use injectable or 1284  
nasally administered glucagon; 1285

(6) Specify that assistance from an emergency medical 1286  
service provider must be requested immediately after a dose of 1287  
glucagon is administered; 1288

(7) Specify the individuals to whom a dose of glucagon may 1289  
be administered in an emergency situation specified under 1290  
division (D) (5) of this section. 1291

(E) (1) The following are not liable in damages in a civil 1292  
action for injury, death, or loss to person or property that 1293  
allegedly arises from an act or omission associated with 1294  
procuring, maintaining, accessing, or using injectable or 1295  
nasally administered glucagon under this section, unless the act 1296  
or omission constitutes willful or wanton misconduct: 1297

(a) A camp; 1298

(b) A camp employee, contractor, or volunteer; 1299

(c) A licensed health professional authorized to prescribe 1300  
drugs who personally furnishes or prescribes injectable or 1301  
nasally administered glucagon, provides a consultation, or 1302  
issues a protocol pursuant to this section; 1303

(2) This section does not eliminate, limit, or reduce any 1304

other immunity or defense that a camp; camp employee, 1305  
contractor, or volunteer; or licensed health professional may be 1306  
entitled to under Chapter 2744. or any other provision of the 1307  
Revised Code or under the common law of this state. 1308

(F) A camp may accept donations of injectable or nasally 1309  
administered glucagon from a wholesale distributor of dangerous 1310  
drugs or manufacturer of dangerous drugs, as defined in section 1311  
4729.01 of the Revised Code, and may accept donations of money 1312  
from any person to purchase the drug. 1313

(G) A camp that elects to procure injectable or nasally 1314  
administered glucagon under this section shall report to the 1315  
department of job and family services each procurement and each 1316  
occurrence in which a dose of the drug is used from the camp's 1317  
supply. 1318

**Section 5.** That existing sections 3313.713, 4723.50, 1319  
4729.01, 4729.51, 4729.513, 4729.541, 4729.60, and 4729.88 of 1320  
the Revised Code are hereby repealed. 1321

**Section 6.** Section 4729.01 of the Revised Code is 1322  
presented in this act as a composite of the section as amended 1323  
by both Sub. S.B. 119 and Sub. S.B. 229 of the 132nd General 1324  
Assembly. The General Assembly, applying the principle stated in 1325  
division (B) of section 1.52 of the Revised Code that amendments 1326  
are to be harmonized if reasonably capable of simultaneous 1327  
operation, finds that the composite is the resulting version of 1328  
the section in effect prior to the effective date of the section 1329  
as presented in this act. 1330