

As Introduced

133rd General Assembly

Regular Session

2019-2020

H. B. No. 700

Representatives Holmes, A., Crossman

A BILL

To amend sections 1751.91, 3719.063, 3923.89, 1
4723.52, 4729.283, 4729.45, 4729.75, 4729.80, 2
4729.84, 4730.56, 4731.83, 5119.363, and 3
5164.14; to amend, for the purpose of adopting a 4
new section number as indicated in parentheses, 5
section 3719.064 (3719.067); and to enact new 6
section 3719.064 and sections 3719.065, 3727.27, 7
3727.61, 4729.791, 4731.92, and 5119.441 of the 8
Revised Code regarding making addiction 9
treatment widely available. 10

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

Section 1. That sections 1751.91, 3719.063, 3923.89, 11
4723.52, 4729.283, 4729.45, 4729.75, 4729.80, 4729.84, 4730.56, 12
4731.83, 5119.363, and 5164.14 be amended; section 3719.064 13
(3719.067) be amended for the purpose of adopting a new section 14
number as indicated in parentheses; and new section 3719.064 and 15
sections 3719.065, 3727.27, 3727.61, 4729.791, 4731.92, and 16
5119.441 of the Revised Code be enacted to read as follows: 17

Sec. 1751.91. A health insuring corporation may provide 18
payment or reimbursement to a pharmacist for providing a health 19

care service to a patient if both of the following are the case: 20

(A) The pharmacist provided the health care service to the 21
patient in accordance with Chapter 4729. of the Revised Code, 22
including any of the following services: 23

(1) Managing drug therapy under a consult agreement with a 24
physician pursuant to section 4729.39 of the Revised Code; 25

(2) Administering immunizations in accordance with section 26
4729.41 of the Revised Code; 27

(3) Administering drugs in accordance with section 4729.45 28
or 4731.92 of the Revised Code. 29

(B) The patient's individual or group health insuring 30
corporation policy, contract, or agreement provides for payment 31
or reimbursement of the service. 32

Sec. 3719.063. In the absence of gross negligence or 33
intentional misconduct, a person who administers the drug 34
naltrexone by injection, the person's employer, and the facility 35
at which the drug is administered are not liable in any civil 36
action or subject to criminal prosecution or professional 37
discipline for any injury or damage caused by the injection or 38
drug if all of the following conditions are met: 39

(A) The individual to whom the drug is administered is 40
unable to have it administered as follows: 41

(1) By a person who routinely administers the drug to the 42
individual; 43

(2) At the facility at which the drug is routinely 44
administered to the individual; 45

(3) Under the direction of the drug's prescriber. 46

(B) The person who administers the drug under this section 47
is legally authorized to administer it by injection but is not 48
the prescriber of the drug or one who routinely administers it 49
to the individual. 50

(C) The drug is provided to the person who administers it 51
under this section in either of the following ways: 52

(1) By the individual to whom it is administered; 53

(2) By the pharmacy that has a record of a prescription 54
for the drug in the name of the individual to whom it is 55
administered. 56

(D) The person who administers the drug under this section 57
is authorized to do so by that person's employer or the facility 58
at which the drug is administered. 59

(E) This section does not apply in the case of an 60
individual who administers an injectable long-acting or 61
extended-release form of naltrexone in accordance with a 62
protocol as authorized by section 4731.92 of the Revised Code. 63

Sec. 3719.064. (A) As used in this section and in section 64
3719.065 of the Revised Code, "prescriber" means any of the 65
following: 66

(1) An advanced practice registered nurse who holds a 67
current, valid license issued under Chapter 4723. of the Revised 68
Code and is designated as a clinical nurse specialist, certified 69
nurse-midwife, or certified nurse practitioner; 70

(2) A physician authorized under Chapter 4731. of the 71
Revised Code to practice medicine and surgery or osteopathic 72
medicine and surgery; 73

(3) A physician assistant who is licensed under Chapter 74

4730. of the Revised Code, holds a valid prescriber number 75
issued by the state medical board, and has been granted 76
physician-delegated prescriptive authority. 77

(B) To the extent permitted by federal law, a prescriber 78
who prescribes opioid analgesics shall offer, during business 79
hours at the location where the prescriber practices, 80
administration of injectable long-acting or extended-release 81
forms of naltrexone. The administration may be delegated in 82
accordance with rules adopted under section 4723.48, 4730.203, 83
or 4731.053 of the Revised Code, as applicable. 84

A prescriber who delegates the administration of 85
injectable long-acting or extended-release forms of naltrexone 86
is not liable in damages to any person or government entity in a 87
civil action for injury, death, or loss to person or property 88
that allegedly arises from an act or omission of the delegate in 89
administering naltrexone, if the prescriber delegates in 90
accordance with this chapter and rules adopted under Chapter 91
4723., 4730., or 4731. of the Revised Code, as applicable. 92

Sec. 3719.065. (A) A prescriber who prescribes methadone 93
or noninjectable forms of buprenorphine shall taper the patient 94
off the drug within sixty days. If such tapering is not 95
possible, only daily doses of those drugs may be personally 96
furnished by the prescriber thereafter. 97

(B) Any prescriber who has obtained a waiver to treat 98
opioid addiction as provided under the federal Drug Addiction 99
Treatment Act of 2000 (DATA 2000), 21 U.S.C. 823(g), is required 100
to have completed training regarding injectable long-acting or 101
extended-release forms of naltrexone and burprenorphine. The 102
state board of pharmacy shall review training programs, 103
including training programs provided by organizations identified 104

in DATA 2000, and approve, for purposes of this section, those 105
it determines meet the requirement of providing the training as 106
specified in this division. 107

Sec. ~~3719.064~~ 3719.067. (A) As used in this section: 108

(1) "Medication-assisted treatment" has the same meaning 109
as in section 340.01 of the Revised Code. 110

(2) "Prescriber" means any of the following: 111

(a) An advanced practice registered nurse who holds a 112
current, valid license issued under Chapter 4723. of the Revised 113
Code and is designated as a clinical nurse specialist, certified 114
nurse-midwife, or certified nurse practitioner; 115

(b) A physician authorized under Chapter 4731. of the 116
Revised Code to practice medicine and surgery or osteopathic 117
medicine and surgery; 118

(c) A physician assistant who is licensed under Chapter 119
4730. of the Revised Code, holds a valid prescriber number 120
issued by the state medical board, and has been granted 121
physician-delegated prescriptive authority. 122

(3) "Qualifying practitioner" has the same meaning as in 123
section 303(g) (2) (G) (iii) of the "Controlled Substances Act of 124
1970," 21 U.S.C. 823(g) (2) (G) (iii), as amended. 125

(B) Before initiating medication-assisted treatment, a 126
prescriber shall give the patient or the patient's 127
representative information about all drugs approved by the 128
United States food and drug administration for use in 129
medication-assisted treatment. The information must be provided 130
both orally and in writing. The prescriber or the prescriber's 131
delegate shall note in the patient's medical record when this 132

information was provided and make the record available to 133
employees of the board of nursing or state medical board on 134
their request. 135

If the prescriber is not a qualifying practitioner and the 136
patient's choice is opioid treatment and the prescriber 137
determines that such treatment is clinically appropriate and 138
meets generally accepted standards of medicine, the prescriber 139
shall refer the patient to an opioid treatment program licensed 140
under section 5119.37 of the Revised Code or a qualifying 141
practitioner. The prescriber or the prescriber's delegate shall 142
make a notation in the patient's medical record naming the 143
program or practitioner to whom the patient was referred and 144
specifying when the referral was made. 145

Sec. 3727.27. If a hospital fails to treat drug addiction 146
with at least eight inpatient beds and an outpatient program, as 147
determined by the director of health, any exemptions or 148
exclusions from taxation authorized by sections 140.08, 5709.08, 149
5709.12, 5709.121, division (B)(1) or (12) of section 5739.02, 150
and division (E)(8) of section 5751.01 of the Revised Code that 151
otherwise apply to the hospital shall cease to apply to that 152
hospital on and after the first day of January of the year 153
following the year in which the determination was made that the 154
hospital is no longer in compliance, notwithstanding anything to 155
the contrary in those sections. On and after that date, the real 156
property owned or held by the hospital shall become subject to 157
property taxation; purchases of tangible personal property or 158
services by the hospital shall be subject to sales and use taxes 159
levied under Chapter 5739. or 5741. of the Revised Code to the 160
extent otherwise applicable to such transactions; and the 161
hospital shall become a taxpayer for the purposes of the tax 162
levied under Chapter 5751. of the Revised Code. Such real 163

property shall continue to be taxable for each tax year until 164
the tax year preceding the tax year in which the determination 165
is made that tax-exempt status is restored; such purchases shall 166
continue to be subject to sales and use taxes levied under 167
Chapter 5739. or 5741. of the Revised Code until the first day 168
of the first month that begins after the date that determination 169
is made; and the hospital shall continue to be a taxpayer for 170
the purposes of the tax levied under Chapter 5751. of the 171
Revised Code until the first day of the tax period that begins 172
after the date that determination is made. 173

Nothing in this section affects the continued exemption 174
from taxation, under section 140.08 of the Revised Code, of 175
obligations issued under section 133.08, 140.06, or 339.15 of 176
the Revised Code or Section 3 of Article XVIII, Ohio 177
Constitution, to pay costs of hospital facilities or to refund 178
such obligations, the transfer of such obligations, the interest 179
and other income from such obligations, or any profit made on 180
their sale. 181

The director of health may adopt rules as the director 182
considers necessary to implement this section. 183

Sec. 3727.61. Each hospital shall perform, on demand and 184
regardless of ability to pay or health insurance coverage, a 185
laboratory test of liver function, the results of which may be 186
used by a person identified in division (B) of section 4731.92 187
of the Revised Code to determine whether it is appropriate to 188
administer to the person tested an injectable long-acting or 189
extended-release form of naltrexone for treatment of drug 190
addiction. 191

Sec. 3923.89. A sickness and accident insurer or public 192
employee benefit plan may provide payment or reimbursement to a 193

pharmacist for providing a health care service to a patient if	194
both of the following are the case:	195
(A) The pharmacist provided the health care service to the	196
patient in accordance with Chapter 4729. of the Revised Code,	197
including any of the following services:	198
(1) Managing drug therapy under a consult agreement with a	199
physician pursuant to section 4729.39 of the Revised Code;	200
(2) Administering immunizations in accordance with section	201
4729.41 of the Revised Code;	202
(3) Administering drugs in accordance with section 4729.45	203
<u>or 4731.92</u> of the Revised Code.	204
(B) The patient's individual or group policy of sickness	205
and accident insurance or public employee benefit plan provides	206
for payment or reimbursement of the service.	207
Sec. 4723.52. (A) As used in this section:	208
(1) "Community addiction services provider" has the same	209
meaning as in section 5119.01 of the Revised Code.	210
(2) "Medication-assisted treatment" has the same meaning	211
as in section 340.01 of the Revised Code.	212
(B) An advanced practice registered nurse shall comply	213
with section 3719.064 <u>3719.067</u> of the Revised Code and rules	214
adopted under section 4723.51 of the Revised Code when treating	215
a patient for addiction with medication-assisted treatment or	216
proposing to initiate such treatment.	217
(C) An advanced practice registered nurse who fails to	218
comply with this section shall treat not more than thirty	219
patients at any one time with medication-assisted treatment even	220

if the facility or location at which the treatment is provided 221
is either of the following: 222

(1) Exempted by divisions (B) (2) (a) to (d) of section 223
4729.553 of the Revised Code from being required to possess a 224
category III terminal distributor of dangerous drugs license 225
with an office-based opioid treatment classification; 226

(2) A community addiction services provider that provides 227
alcohol and drug addiction services that are certified by the 228
department of mental health and addiction services under section 229
5119.36 of the Revised Code. 230

Sec. 4729.283. (A) A pharmacist may dispense naltrexone 231
without a written or oral prescription from a licensed health 232
professional authorized to prescribe drugs if all of the 233
following conditions are met: 234

(1) The pharmacist is able to verify a record of a 235
prescription for the injectable long-acting or extended-release 236
form of naltrexone in the name of the patient who is requesting 237
the drug, but the prescription does not provide for a refill or 238
the time permitted by rules adopted by the state board of 239
pharmacy for providing refills has elapsed. 240

(2) The pharmacist is unable to obtain authorization to 241
refill the prescription from the prescriber who issued it or 242
another prescriber responsible for the patient's care. 243

(3) In the exercise of the pharmacist's professional 244
judgment: 245

(a) The drug is necessary to continue the patient's 246
therapy for substance use disorder. 247

(b) Failure to dispense the drug to the patient could 248

result in harm to the health of the patient.	249
(B) Before dispensing naltrexone under this section, the pharmacist shall offer the patient the choice of receiving either the oral form or injectable long-acting or extended-release form, but only if both forms of the drug are available for dispensing at the time of the patient's request or within one day after the request.	250 251 252 253 254 255
(C) (1) With respect to naltrexone dispensed in an oral form under this section, the pharmacist shall not dispense an amount that exceeds a five-day supply.	256 257 258
(2) With respect to naltrexone dispensed in an injectable long-acting or extended-release form under this section, both of the following apply:	259 260 261
(a) The pharmacist shall exercise professional judgment in determining the amount of the drug dispensed.	262 263
(b) The pharmacist may administer the drug by injection to the patient but only in accordance with section 4729.45 <u>4731.92</u> of the Revised Code.	264 265 266
(D) A pharmacist who dispenses naltrexone under this section shall do all of the following:	267 268
(1) For one year after the date of dispensing, maintain a record in accordance with this chapter of the drug dispensed, including the amount and form dispensed, the original prescription number, the name and address of the patient and, if the individual receiving the drug is not the patient, the name and address of that individual;	269 270 271 272 273 274
(2) Notify the prescriber who issued the prescription described in division (A) (1) of this section or another	275 276

prescriber responsible for the patient's care not later than	277
five days after the drug is dispensed;	278
(3) If applicable, obtain authorization for additional	279
dispensing from one of the prescribers described in division (D)	280
(2) of this section.	281
(E) A pharmacist shall exercise professional judgment in	282
determining the number of times naltrexone may be dispensed	283
under this section to the same patient.	284
(F) This section does not limit the authority of a	285
pharmacist to dispense a dangerous drug under section 4729.281	286
of the Revised Code.	287
Sec. 4729.45. (A) As used in this section, "physician"	288
means an individual authorized under Chapter 4731. of the	289
Revised Code to practice medicine and surgery or osteopathic	290
medicine and surgery.	291
(B) (1) Subject to division (C) of this section, a	292
pharmacist licensed under this chapter may administer by	293
injection any of the following drugs as long as the drug that is	294
to be administered has been prescribed by a physician and the	295
individual to whom the drug was prescribed has an ongoing	296
physician-patient relationship with the physician:	297
(a) An opioid antagonist used for treatment of drug	298
addiction and administered in a long-acting or extended-release	299
form;	300
(b) An antipsychotic drug administered in a long-acting or	301
extended-release form;	302
(e) <u>(b)</u> Hydroxyprogesterone caproate;	303
(d) <u>(c)</u> Medroxyprogesterone acetate;	304

(e) <u>(d)</u> Cobalamin.	305
(2) As part of engaging in the administration of drugs by injection pursuant to this section, a pharmacist may administer epinephrine or diphenhydramine, or both, to an individual in an emergency situation resulting from an adverse reaction to a drug administered by the pharmacist.	306 307 308 309 310
(C) To be authorized to administer drugs pursuant to this section, a pharmacist must do all of the following:	311 312
(1) Successfully complete a course in the administration of drugs that satisfies the requirements established by the state board of pharmacy in rules adopted under division (H) <u>(G)</u> (1) (a) of this section;	313 314 315 316
(2) Receive and maintain certification to perform basic life-support procedures by successfully completing a basic life-support training course that is certified by the American red cross or American heart association or approved by the state board of pharmacy;	317 318 319 320 321
(3) Practice in accordance with a protocol that meets the requirements of division (F) <u>(E)</u> of this section.	322 323
(D) Each time a pharmacist administers a drug pursuant to this section, the pharmacist shall do all of the following:	324 325
(1) Obtain permission in accordance with the procedures specified in rules adopted under division (H) <u>(G)</u> of this section and comply with the following requirements:	326 327 328
(a) Except as provided in division (D) (1) (c) of this section, for each drug administered by a pharmacist to an individual who is eighteen years of age or older, the pharmacist shall obtain permission from the individual.	329 330 331 332

(b) For each drug administered by a pharmacist to an individual who is under eighteen years of age, the pharmacist shall obtain permission from the individual's parent or other person having care or charge of the individual.

(c) For each drug administered by a pharmacist to an individual who lacks the capacity to make informed health care decisions, the pharmacist shall obtain permission from the person authorized to make such decisions on the individual's behalf.

~~(2) In the case of an opioid antagonist described in division (B) of this section, obtain in accordance with division (E) of this section test results indicating that it is appropriate to administer the drug to the individual if either of the following is to be administered:~~

~~(a) The initial dose of the drug;~~

~~(b) Any subsequent dose, if the administration occurs more than thirty days after the previous dose of the drug was administered.~~

~~(3) Observe the individual to whom the drug is administered to determine whether the individual has an adverse reaction to the drug;~~

~~(4) (3) Notify the physician who prescribed the drug that the drug has been administered to the individual.~~

~~(E) A pharmacist may obtain the test results described in division (D) (2) of this section in either of the following ways:~~

~~(1) From the physician;~~

~~(2) By ordering blood and urine tests for the individual to whom the opioid antagonist is to be administered.~~

~~If a pharmacist orders blood and urine tests, the pharmacist shall evaluate the results of the tests to determine whether they indicate that it is appropriate to administer the opioid antagonist. A pharmacist's authority to evaluate test results under this division does not authorize the pharmacist to make a diagnosis.~~

~~(F)~~All of the following apply with respect to the protocol required by division (C) (3) of this section:

(1) The protocol must be established by a physician who has a scope of practice that includes treatment of the condition for which the individual has been prescribed the drug to be administered.

(2) The protocol must satisfy the requirements established in rules adopted under division ~~(H)~~(G) (1) (b) of this section.

(3) The protocol must do all of the following:

(a) Specify a definitive set of treatment guidelines;

(b) Specify the locations at which a pharmacist may engage in the administration of drugs pursuant to this section;

(c) Include provisions for implementing the requirements of division (D) of this section, including for purposes of division (D) ~~(3)~~(2) of this section provisions specifying the length of time and location at which a pharmacist must observe an individual who receives a drug to determine whether the individual has an adverse reaction to the drug;

(d) Specify procedures to be followed by a pharmacist when administering epinephrine, diphenhydramine, or both, to an individual who has an adverse reaction to a drug administered by the pharmacist.

(G) <u>(F)</u> A pharmacist shall not do either of the following:	389
(1) Engage in the administration of drugs pursuant to this section unless the requirements of division (C) of this section have been met;	390 391 392
(2) Delegate to any person the pharmacist's authority to engage in the administration of drugs pursuant to this section.	393 394
(H) <u>(G)</u> (1) The state board of pharmacy shall adopt rules to implement this section. The rules shall be adopted in accordance with Chapter 119. of the Revised Code and include all of the following:	395 396 397 398
(a) Requirements for courses in administration of drugs;	399
(b) Requirements for protocols to be followed by pharmacists in administering drugs pursuant to this section;	400 401
(c) Procedures to be followed by a pharmacist in obtaining permission to administer a drug to an individual.	402 403
(2) The board shall consult with the state medical board before adopting rules regarding requirements for protocols under this section.	404 405 406
Sec. 4729.75. The state board of pharmacy may establish and maintain a drug database. The board shall use the drug database to monitor the misuse and diversion of the following: controlled substances, as defined in section 3719.01 of the Revised Code; medical marijuana, as authorized under Chapter 3796. of the Revised Code; and other dangerous drugs the board includes in the database pursuant to rules adopted under section 4729.84 of the Revised Code.	407 408 409 410 411 412 413 414
The board also shall use the drug database to monitor naltrexone, <u>including the administration of injectable long-</u>	415 416

acting or extended-release forms of naltrexone as authorized 417
under section 4731.92 of the Revised Code. 418

In establishing and maintaining the database, the board 419
shall electronically collect information pursuant to sections 420
4729.77, 4729.771, 4729.772, 4729.78, ~~and 4729.79~~, and 4729.791 421
of the Revised Code and shall disseminate information as 422
authorized or required by sections 4729.80 and 4729.81 of the 423
Revised Code. The board's collection and dissemination of 424
information shall be conducted in accordance with rules adopted 425
under section 4729.84 of the Revised Code. 426

Sec. 4729.791. (A) (1) If the state board of pharmacy 427
establishes and maintains a drug database pursuant to section 428
4729.75 of the Revised Code, the following individuals who 429
administer injectable long-acting or extended-release forms of 430
naltrexone shall submit to the board the information identified 431
in division (A) (2) of this section: 432

(a) A licensed health professional authorized to prescribe 433
drugs; 434

(b) An individual identified in division (B) of section 435
4731.92. 436

(2) Each individual identified in division (A) of this 437
section shall submit the following information to the board: 438

(a) The individual's name and licensing board; 439

(b) The name of the individual receiving the drug by 440
injection; 441

(c) The date the drug was administered; 442

(d) The name, strength, and national drug code of the drug 443
furnished; 444

(e) Any other information specified by the board in rules 445
adopted under section 4729.84 of the Revised Code. 446

(B) If the state board of pharmacy establishes and 447
maintains a drug database pursuant to section 4729.75 of the 448
Revised Code, each licensed health professional who receives 449
test results indicating whether or not it is appropriate to 450
administer to an individual an injectable long-acting or 451
extended-release form of naltrexone shall submit to the board 452
the following: 453

(1) Health professional identification; 454

(2) Patient identification; 455

(3) Date and results of the test; 456

(4) Any other information specified by the board in rules 457
adopted under section 4729.84 of the Revised Code. 458

(C) Information required by this section shall be 459
transmitted as specified by the board in rules adopted under 460
section 4729.84 of the Revised Code. 461

The information shall be submitted electronically in the 462
format specified by the board, except that the board may grant a 463
waiver allowing the individual to submit the information in 464
another format. 465

The information shall be submitted in accordance with any 466
time limits specified by the board, except that the board may 467
grant an extension if either of the following occurs: 468

(1) The individual's transmission system suffers a 469
mechanical or electronic failure or the individual cannot meet 470
the deadline for other reasons beyond the individual's control. 471

<u>(2) The board is unable to receive electronic submissions.</u>	472
<u>(D) If the board becomes aware of an individual's failure</u>	473
<u>to comply with this section, the board shall notify the</u>	474
<u>government entity responsible for licensing the individual.</u>	475
Sec. 4729.80. (A) If the state board of pharmacy	476
establishes and maintains a drug database pursuant to section	477
4729.75 of the Revised Code, the board is authorized or required	478
to provide information from the database only as follows:	479
(1) On receipt of a request from a designated	480
representative of a government entity responsible for the	481
licensure, regulation, or discipline of health care	482
professionals with authority to prescribe, administer, or	483
dispense drugs, the board may provide to the representative	484
information from the database relating to the professional who	485
is the subject of an active investigation being conducted by the	486
government entity or relating to a professional who is acting as	487
an expert witness for the government entity in such an	488
investigation.	489
(2) On receipt of a request from a federal officer, or a	490
state or local officer of this or any other state, whose duties	491
include enforcing laws relating to drugs, the board shall	492
provide to the officer information from the database relating to	493
the person who is the subject of an active investigation of a	494
drug abuse offense, as defined in section 2925.01 of the Revised	495
Code, being conducted by the officer's employing government	496
entity.	497
(3) Pursuant to a subpoena issued by a grand jury, the	498
board shall provide to the grand jury information from the	499
database relating to the person who is the subject of an	500

investigation being conducted by the grand jury. 501

(4) Pursuant to a subpoena, search warrant, or court order 502
in connection with the investigation or prosecution of a 503
possible or alleged criminal offense, the board shall provide 504
information from the database as necessary to comply with the 505
subpoena, search warrant, or court order. 506

(5) On receipt of a request from a prescriber or the 507
prescriber's delegate approved by the board, the board shall 508
provide to the prescriber a report of information from the 509
database relating to a patient who is either a current patient 510
of the prescriber or a potential patient of the prescriber based 511
on a referral of the patient to the prescriber, if all of the 512
following conditions are met: 513

(a) The prescriber certifies in a form specified by the 514
board that it is for the purpose of providing medical treatment 515
to the patient who is the subject of the request; 516

(b) The prescriber has not been denied access to the 517
database by the board. 518

(6) On receipt of a request from a pharmacist or the 519
pharmacist's delegate approved by the board, the board shall 520
provide to the pharmacist information from the database relating 521
to a current patient of the pharmacist, if the pharmacist 522
certifies in a form specified by the board that it is for the 523
purpose of the pharmacist's practice of pharmacy involving the 524
patient who is the subject of the request and the pharmacist has 525
not been denied access to the database by the board. 526

(7) On receipt of a request from an individual seeking the 527
individual's own database information in accordance with the 528
procedure established in rules adopted under section 4729.84 of 529

the Revised Code, the board may provide to the individual the 530
individual's own prescription history. 531

(8) On receipt of a request from a medical director or a 532
pharmacy director of a managed care organization that has 533
entered into a contract with the department of medicaid under 534
section 5167.10 of the Revised Code and a data security 535
agreement with the board required by section 5167.14 of the 536
Revised Code, the board shall provide to the medical director or 537
the pharmacy director information from the database relating to 538
a medicaid recipient enrolled in the managed care organization, 539
including information in the database related to prescriptions 540
for the recipient that were not covered or reimbursed under a 541
program administered by the department of medicaid. 542

(9) On receipt of a request from the medicaid director, 543
the board shall provide to the director information from the 544
database relating to a recipient of a program administered by 545
the department of medicaid, including information in the 546
database related to prescriptions for the recipient that were 547
not covered or paid by a program administered by the department. 548

(10) On receipt of a request from a medical director of a 549
managed care organization that has entered into a contract with 550
the administrator of workers' compensation under division (B) (4) 551
of section 4121.44 of the Revised Code and a data security 552
agreement with the board required by section 4121.447 of the 553
Revised Code, the board shall provide to the medical director 554
information from the database relating to a claimant under 555
Chapter 4121., 4123., 4127., or 4131. of the Revised Code 556
assigned to the managed care organization, including information 557
in the database related to prescriptions for the claimant that 558
were not covered or reimbursed under Chapter 4121., 4123., 559

4127., or 4131. of the Revised Code, if the administrator of workers' compensation confirms, upon request from the board, that the claimant is assigned to the managed care organization.

(11) On receipt of a request from the administrator of workers' compensation, the board shall provide to the administrator information from the database relating to a claimant under Chapter 4121., 4123., 4127., or 4131. of the Revised Code, including information in the database related to prescriptions for the claimant that were not covered or reimbursed under Chapter 4121., 4123., 4127., or 4131. of the Revised Code.

(12) On receipt of a request from a prescriber or the prescriber's delegate approved by the board, the board shall provide to the prescriber information from the database relating to a patient's mother, if the prescriber certifies in a form specified by the board that it is for the purpose of providing medical treatment to a newborn or infant patient diagnosed as opioid dependent and the prescriber has not been denied access to the database by the board.

(13) On receipt of a request from the director of health, the board shall provide to the director information from the database relating to the duties of the director or the department of health in implementing the Ohio violent death reporting system established under section 3701.93 of the Revised Code.

(14) On receipt of a request from a requestor described in division (A)(1), (2), (5), or (6) of this section who is from or participating with another state's prescription monitoring program, the board may provide to the requestor information from the database, but only if there is a written agreement under

which the information is to be used and disseminated according 590
to the laws of this state. 591

(15) On receipt of a request from a delegate of a retail 592
dispensary licensed under Chapter 3796. of the Revised Code who 593
is approved by the board to serve as the dispensary's delegate, 594
the board shall provide to the delegate a report of information 595
from the database pertaining only to a patient's use of medical 596
marijuana, if both of the following conditions are met: 597

(a) The delegate certifies in a form specified by the 598
board that it is for the purpose of dispensing medical marijuana 599
for use in accordance with Chapter 3796. of the Revised Code. 600

(b) The retail dispensary or delegate has not been denied 601
access to the database by the board. 602

(16) On receipt of a request from a judge of a program 603
certified by the Ohio supreme court as a specialized docket 604
program for drugs, the board shall provide to the judge, or an 605
employee of the program who is designated by the judge to 606
receive the information, information from the database that 607
relates specifically to a current or prospective program 608
participant. 609

(17) On receipt of a request from a coroner, deputy 610
coroner, or coroner's delegate approved by the board, the board 611
shall provide to the requestor information from the database 612
relating to a deceased person about whom the coroner is 613
conducting or has conducted an autopsy or investigation. 614

(18) On receipt of a request from a prescriber, the board 615
may provide to the prescriber a summary of the prescriber's 616
prescribing record if such a record is created by the board. 617
Information in the summary is subject to the confidentiality 618

requirements of this chapter. 619

(19) (a) On receipt of a request from a pharmacy's 620
responsible person, the board may provide to the responsible 621
person a summary of the pharmacy's dispensing record if such a 622
record is created by the board. Information in the summary is 623
subject to the confidentiality requirements of this chapter. 624

(b) As used in division (A) (19) (a) of this section, 625
"responsible person" has the same meaning as in rules adopted by 626
the board under section 4729.26 of the Revised Code. 627

(20) The board may provide information from the database 628
without request to a prescriber or pharmacist who is authorized 629
to use the database pursuant to this chapter. 630

(21) (a) On receipt of a request from a prescriber or 631
pharmacist, or the prescriber's or pharmacist's delegate, who is 632
a designated representative of a peer review committee, the 633
board shall provide to the committee information from the 634
database relating to a prescriber who is subject to the 635
committee's evaluation, supervision, or discipline if the 636
information is to be used for one of those purposes. The board 637
shall provide only information that it determines, in accordance 638
with rules adopted under section 4729.84 of the Revised Code, is 639
appropriate to be provided to the committee. 640

(b) As used in division (A) (21) (a) of this section, "peer 641
review committee" has the same meaning as in section 2305.25 of 642
the Revised Code, except that it includes only a peer review 643
committee of a hospital or a peer review committee of a 644
nonprofit health care corporation that is a member of the 645
hospital or of which the hospital is a member. 646

(22) Any personal health information submitted to the 647

board pursuant to section 4729.772 of the Revised Code may be 648
provided by the board only as authorized by the submitter of the 649
information and in accordance with rules adopted under section 650
4729.84 of the Revised Code. 651

(23) On receipt of a request from an individual identified 652
in division (B) of section 4731.92 of the Revised Code, the 653
board shall provide to the individual a report of information 654
from the database pertaining only to a patient's treatment for 655
drug addiction. 656

(B) The state board of pharmacy shall maintain a record of 657
each individual or entity that requests information from the 658
database pursuant to this section. In accordance with rules 659
adopted under section 4729.84 of the Revised Code, the board may 660
use the records to document and report statistics and law 661
enforcement outcomes. 662

The board may provide records of an individual's requests 663
for database information only to the following: 664

(1) A designated representative of a government entity 665
that is responsible for the licensure, regulation, or discipline 666
of health care professionals with authority to prescribe, 667
administer, or dispense drugs who is involved in an active 668
criminal or disciplinary investigation being conducted by the 669
government entity of the individual who submitted the requests 670
for database information; 671

(2) A federal officer, or a state or local officer of this 672
or any other state, whose duties include enforcing laws relating 673
to drugs and who is involved in an active investigation being 674
conducted by the officer's employing government entity of the 675
individual who submitted the requests for database information; 676

(3) A designated representative of the department of 677
medicaid regarding a prescriber who is treating or has treated a 678
recipient of a program administered by the department and who 679
submitted the requests for database information. 680

(C) Information contained in the database and any 681
information obtained from it is confidential and is not a public 682
record. Information contained in the records of requests for 683
information from the database is confidential and is not a 684
public record. Information contained in the database that does 685
not identify a person, including any licensee or registrant of 686
the board or other entity, may be released in summary, 687
statistical, or aggregate form. 688

(D) A pharmacist or prescriber shall not be held liable in 689
damages to any person in any civil action for injury, death, or 690
loss to person or property on the basis that the pharmacist or 691
prescriber did or did not seek or obtain information from the 692
database. 693

Sec. 4729.84. For purposes of establishing and maintaining 694
a drug database pursuant to section 4729.75 of the Revised Code, 695
the state board of pharmacy shall adopt rules in accordance with 696
Chapter 119. of the Revised Code to carry out and enforce 697
sections 4729.75 to 4729.83 of the Revised Code. The rules shall 698
specify all of the following: 699

(A) A means of identifying each patient, each terminal 700
distributor of dangerous drugs, each purchase at wholesale of 701
dangerous drugs, and each retail dispensary licensed under 702
Chapter 3796. of the Revised Code about which information is 703
entered into the drug database; 704

(B) Requirements for the transmission of information from 705

terminal distributors of dangerous drugs, manufacturers of 706
dangerous drugs, outsourcing facilities, repackagers of 707
dangerous drugs, wholesale distributors of dangerous drugs, 708
prescribers, ~~and~~ retail dispensaries, and other individuals 709
required to transmit information to the board; 710

(C) An electronic format for the submission of information 711
from persons identified in division (B) of this section; 712

(D) A procedure whereby a person unable to submit 713
information electronically may obtain a waiver to submit 714
information in another format; 715

(E) A procedure whereby the board may grant a request from 716
a law enforcement agency or a government entity responsible for 717
the licensure, regulation, or discipline of licensed health 718
professionals authorized to prescribe drugs that information 719
that has been stored for three years be retained when the 720
information pertains to an open investigation being conducted by 721
the agency or entity; 722

(F) A procedure whereby a person identified in division 723
(B) of this section may apply for an extension to the time by 724
which information must be transmitted to the board; 725

(G) A procedure whereby a person or government entity to 726
which the board is authorized to provide information may submit 727
a request to the board for the information and the board may 728
verify the identity of the requestor; 729

(H) Standards for determining what information is 730
appropriate to be provided under division (A) (21) of section 731
4729.80 of the Revised Code; 732

(I) A procedure whereby the board can use the database 733
request records required by division (B) of section 4729.80 of 734

the Revised Code to document and report statistics and law 735
enforcement outcomes; 736

(J) A procedure whereby an individual may request the 737
individual's own database information and the board may verify 738
the identity of the requestor; 739

(K) A reasonable fee that the board may charge under 740
section 4729.83 of the Revised Code for providing an individual 741
with the individual's own database information pursuant to 742
section 4729.80 of the Revised Code; 743

(L) The other specific dangerous drugs that, in addition 744
to controlled substances, must be included in the database; 745

(M) The types of pharmacies licensed as terminal 746
distributors of dangerous drugs that are required to submit 747
prescription information to the board pursuant to section 748
4729.77 of the Revised Code; 749

(N) Additional data fields, recognized by the American 750
society for automation in pharmacy, that licensed terminal 751
distributors of dangerous drugs must submit to the board 752
pursuant to section 4729.77 of the Revised Code; 753

(O) The information regarding medical marijuana dispensed 754
to a patient that a retail dispensary is required to submit to 755
the board pursuant to section 4729.771 of the Revised Code; 756

(P) Requirements for the transmission of information 757
pursuant to section 4729.772 of the Revised Code and 758
requirements for the release of such information by the board; 759

(Q) Any additional information that must be submitted to 760
the board pursuant to section 4729.791 of the Revised Code. 761

Sec. 4730.56. (A) As used in this section: 762

(1) "Community addiction services provider" has the same meaning as in section 5119.01 of the Revised Code.	763 764
(2) "Medication-assisted treatment" has the same meaning as in section 340.01 of the Revised Code.	765 766
(B) A physician assistant shall comply with section 3719.064 <u>3719.067</u> of the Revised Code and rules adopted under section 4730.55 of the Revised Code when treating a patient with medication-assisted treatment or proposing to initiate such treatment.	767 768 769 770 771
(C) A physician assistant who fails to comply with this section shall treat not more than thirty patients at any one time with medication-assisted treatment even if the facility or location at which the treatment is provided is either of the following:	772 773 774 775 776
(1) Exempted by divisions (B) (2) (a) to (d) of section 4729.553 of the Revised Code from being required to possess a category III terminal distributor of dangerous drugs license with an office-based opioid treatment classification;	777 778 779 780
(2) A community addiction services provider that provides alcohol and drug addiction services that are certified by the department of mental health and addiction services under section 5119.36 of the Revised Code.	781 782 783 784
Sec. 4731.83. (A) As used in this section:	785
(1) "Medication-assisted treatment" has the same meaning as in section 340.01 of the Revised Code.	786 787
(2) "Physician" means an individual authorized by this chapter to practice medicine and surgery or osteopathic medicine and surgery.	788 789 790

(B) A physician shall comply with section ~~3719.064~~ 791
3719.067 of the Revised Code and rules adopted under section 792
4731.056 of the Revised Code when treating a patient with 793
medication-assisted treatment or proposing to initiate such 794
treatment. 795

(C) A physician who fails to comply with this section 796
shall treat not more than thirty patients at any one time with 797
medication-assisted treatment even if the facility or location 798
at which the treatment is provided is either of the following: 799

(1) Exempted by divisions (B) (2) (a) to (d) of section 800
4729.553 of the Revised Code from being required to possess a 801
category III terminal distributor of dangerous drugs license 802
with an office-based opioid treatment classification; 803

(2) A community addiction services provider that provides 804
alcohol and drug addiction services that are certified by the 805
department of mental health and addiction services under section 806
5119.36 of the Revised Code. 807

Sec. 4731.92. (A) As used in this section, "physician" 808
means an individual authorized to practice medicine and surgery 809
or osteopathic medicine and surgery. 810

(B) Notwithstanding any conflicting provision of the 811
Revised Code or rule adopted under it, any of the following 812
individuals who comply with division (C) of this section may 813
administer by injection, in accordance with a protocol that 814
meets the requirements of division (F) of this section, long- 815
acting or extended-release forms of naltrexone for treatment of 816
drug addiction: 817

(1) A pharmacist licensed or otherwise authorized to 818
practice by the state board of pharmacy under Chapter 4729. of 819

<u>the Revised Code;</u>	820
<u>(2) A psychologist licensed or otherwise authorized to</u>	821
<u>practice by the state board of psychology under Chapter 4732. of</u>	822
<u>the Revised Code;</u>	823
<u>(3) An individual licensed or otherwise authorized to</u>	824
<u>practice by the chemical dependency professionals board under</u>	825
<u>Chapter 4758. of the Revised Code;</u>	826
<u>(4) An individual licensed or otherwise authorized to</u>	827
<u>practice by the counselor, social worker, and marriage and</u>	828
<u>family therapist board under Chapter 4757. of the Revised Code;</u>	829
<u>(5) An individual licensed or otherwise authorized to</u>	830
<u>practice by the state board of emergency medical, fire, and</u>	831
<u>transportation services under Chapter 4765. of the Revised Code;</u>	832
<u>(6) A police officer;</u>	833
<u>(7) A licensed health care professional not otherwise</u>	834
<u>listed in this section that is specifically identified in a</u>	835
<u>protocol that meets the requirements of division (F) of this</u>	836
<u>section.</u>	837
<u>(C) To be authorized to administer injectable long-acting</u>	838
<u>or extended-release forms of naltrexone pursuant to this</u>	839
<u>section, an individual identified in division (B) of this</u>	840
<u>section must do all of the following:</u>	841
<u>(1) Successfully complete an online course in the</u>	842
<u>administration of drugs that satisfies the requirements</u>	843
<u>established by the state medical board in rules adopted under</u>	844
<u>division (I) of this section;</u>	845
<u>(2) Receive and maintain certification to perform basic</u>	846
<u>life-support procedures by successfully completing a basic life-</u>	847

support training course certified by the American red cross or 848
American heart association; 849

(3) Practice in accordance with a protocol that meets the 850
requirements of division (F) of this section. 851

(D) Each time an individual administers a drug pursuant to 852
this section, the individual shall do both of the following: 853

(1) Except as provided in division (E) (2) of this section, 854
obtain in accordance with division (E) of this section test 855
results indicating that it is appropriate to administer the 856
drug; 857

(2) Submit to the state board of pharmacy the information 858
identified in section 4729.791 of the Revised Code. 859

(E) (1) An individual identified in division (B) of this 860
section may obtain the test results described in division (D) (1) 861
of this section in any of the following ways: 862

(a) From a physician; 863

(b) From the drug database established under section 864
4729.75 of the Revised Code; 865

(c) From a hospital; 866

(d) From the person on whom the test described in division 867
(D) (1) of this section was performed. 868

(2) If the individual seeking to administer a drug in 869
accordance with this section is unable to obtain test results 870
indicating that it is appropriate to administer the drug and the 871
recipient of the drug declares that the recipient is unable to 872
get the test, the individual may administer the drug to the 873
recipient for not more than sixty days. 874

(F) The protocol required by division (C) (3) of this section must do both of the following: 875
876

(1) Be established by a physician whose regular practice includes treatment of the condition for which the recipient is receiving the drug to be administered; 877
878
879

(2) Satisfy the requirements established in rules adopted under division (I) of this section. 880
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(G) An individual identified in division (B) of this section is not liable for damages in any civil action allegedly arising from, or subject to prosecution in any criminal proceeding or professional disciplinary action for, any act or omission associated with administering injectable long-acting or extended-release forms of naltrexone under this section, unless the act or omission constitutes willful or wanton misconduct. 882
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(H) Nothing in this section requires an individual identified in division (B) of this section to administer a drug by injection. 889
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(I) The state medical board shall adopt rules to implement this section. The rules shall be adopted in accordance with Chapter 119. of the Revised Code and include at least the following: 892
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(1) Requirements for online courses in the administration of drugs; 896
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(2) Requirements for protocols established under this section. 898
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Sec. 5119.363. The director of mental health and addiction services shall adopt rules governing the duties of boards of alcohol, drug addiction, and mental health services under 900
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section 340.20 of the Revised Code and the duties of community 903
addiction services providers under section 5119.362 of the 904
Revised Code. The rules shall be adopted in accordance with 905
Chapter 119. of the Revised Code. 906

The director shall adopt rules under this section that 907
authorize the department of mental health and addiction services 908
to determine an advanced practice registered nurse's, physician 909
assistant's, or physician's compliance with section ~~3719.064-~~ 910
3719.067 of the Revised Code if such practitioner works for a 911
community addiction services provider. 912

Sec. 5119.441. (A) The department of mental health and 913
addiction services shall procure injectable long-acting or 914
extended-release forms of naltrexone and buprenorphine directly 915
from drug manufacturers and coordinate with state, county, and 916
municipal agencies to distribute the drugs as needed to treat 917
drug-addicted individuals in this state, including distribution 918
to individuals identified in division (B) of section 4731.92 of 919
the Revised Code. The department shall require monitoring and 920
monthly administration of the drugs by boards of health, boards 921
of alcohol, drug addiction, and mental health services, courts, 922
and parole and probation officers. 923

(B) The department shall contract with a licensed terminal 924
distributor of dangerous drugs to serve as a central pharmacy 925
that is responsible for obtaining statewide contract pricing and 926
from which political subdivisions can make direct purchases of 927
injectable long-acting or extended-release forms of naltrexone 928
and buprenorphine. 929

(C) In procuring injectable long-acting or extended- 930
release forms of naltrexone and buprenorphine pursuant to this 931
section, the department may use rebates to further discount the 932

<u>drug's price.</u>	933
Sec. 5164.14. The medicaid program may cover a health care	934
service that a pharmacist provides to a medicaid recipient in	935
accordance with Chapter 4729. of the Revised Code, including any	936
of the following services:	937
(A) Managing drug therapy under a consult agreement with a	938
physician pursuant to section 4729.39 of the Revised Code;	939
(B) Administering immunizations in accordance with section	940
4729.41 of the Revised Code;	941
(C) Administering drugs in accordance with section 4729.45	942
<u>or 4731.92</u> of the Revised Code.	943
Section 2. That existing sections 1751.91, 3719.063,	944
3719.064, 3923.89, 4723.52, 4729.283, 4729.45, 4729.75, 4729.80,	945
4729.84, 4730.56, 4731.83, 5119.363, and 5164.14 of the Revised	946
Code are hereby repealed.	947