As Introduced

131st General Assembly Regular Session 2015-2016

S. B. No. 357

Senator Hite

A BILL

To amend sections 1739.05 and 5167.12 and to enact	1
sections 1751.691, 3923.851, and 5164.091 of the	2
Revised Code regarding health insurance,	3
Medicaid, and abuse-deterrent opioid analgesic	4
drug products.	5

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

Section 1. That sections 1739.05 and 5167.12 be amended	6
and sections 1751.691, 3923.851, and 5164.091 of the Revised	7
Code be enacted to read as follows:	8
Sec. 1739.05. (A) A multiple employer welfare arrangement	9
that is created pursuant to sections 1739.01 to 1739.22 of the	10
Revised Code and that operates a group self-insurance program	11
may be established only if any of the following applies:	12
(1) The arrangement has and maintains a minimum enrollment	13
(1) The arrangement has and marntains a minimum enroriment	тэ
of three hundred employees of two or more employers.	14
(2) The arrangement has and maintains a minimum enrollment	15
of three hundred self-employed individuals.	16
(2) The experiment has and maintains a minimum envellment	17
(3) The arrangement has and maintains a minimum enrollment	т /
of three hundred employees or self-employed individuals in any	18

combination of divisions (A)(1) and (2) of this section. 19 (B) A multiple employer welfare arrangement that is 20 created pursuant to sections 1739.01 to 1739.22 of the Revised 21 Code and that operates a group self-insurance program shall 22 comply with all laws applicable to self-funded programs in this 23 state, including sections 3901.04, 3901.041, 3901.19 to 3901.26, 24 3901.38, 3901.381 to 3901.3814, 3901.40, 3901.45, 3901.46, 25 3901.491, 3902.01 to 3902.14, 3923.24, 3923.282, 3923.30, 26 3923.301, 3923.38, 3923.581, 3923.63, 3923.80, 3923.85, 27 <u>3923.851,</u> 3924.031, 3924.032, and 3924.27 of the Revised Code. 28 (C) A multiple employer welfare arrangement created 29 30

pursuant to sections 1739.01 to 1739.22 of the Revised Code30shall solicit enrollments only through agents or solicitors31licensed pursuant to Chapter 3905. of the Revised Code to sell32or solicit sickness and accident insurance.33

(D) A multiple employer welfare arrangement created 34 pursuant to sections 1739.01 to 1739.22 of the Revised Code 35 shall provide benefits only to individuals who are members, 36 employees of members, or the dependents of members or employees, 37 or are eligible for continuation of coverage under section 38 1751.53 or 3923.38 of the Revised Code or under Title X of the 39 "Consolidated Omnibus Budget Reconciliation Act of 1985," 100 40 Stat. 227, 29 U.S.C.A. 1161, as amended. 41

(E) A multiple employer welfare arrangement created
pursuant to sections 1739.01 to 1739.22 of the Revised Code is
subject to, and shall comply with, sections 3903.81 to 3903.93
of the Revised Code in the same manner as other life or health
insurers, as defined in section 3903.81 of the Revised Code.

Sec. 1751.691. (A) As used in this section:

Page 2

47

(1) "Abuse-deterrent opioid analgesic drug product" means	48
a brand or generic opioid analgesic drug product approved by the	49
United States food and drug administration with abuse-deterrence	50
labeling claims indicating its abuse-deterrent properties are	51
expected to deter or reduce its abuse.	52
(2) "Opioid analgesic" has the same meaning as in section	53
<u>3719.01 of the Revised Code.</u>	54
<u>5715.01 01 the Revised code.</u>	51
(3) "Prescriber" has the same meaning as in section	55
4729.01 of the Revised Code.	56
(B) Notwithstanding section 3901.71 of the Revised Code,	57
an individual or group health insuring corporation policy,	58
contract, or agreement that is delivered, issued for delivery,	59
or renewed in this state and covers opioid analgesic drug	60
products as part of providing any coverage of prescription drugs	61
shall provide access to abuse-deterrent opioid analgesic drug	62
products in the drug formulary or other list of covered drugs	63
that applies under the policy, contract, or agreement.	64
(C) Doth of the following apply to any prior authorization	65
(C) Both of the following apply to any prior authorization	66
requirements or utilization review measures contained in a	
health insuring corporation policy, contract, or agreement	67
subject to this section and any coverage denials made pursuant	68
to those requirements or measures with respect to opioid	69
analgesic drug products:	70
(1) Prior authorization requirements or utilization review	71
measures shall not be any more restrictive for abuse-deterrent	72
opioid analgesic drug products than for opioid analgesic drug	73
products that are not abuse-deterrent opioid analgesic drug	74
products.	75
(2) Prior authorization requirements or utilization review_	76

measures shall not require treatment with an opioid analgesic	77
drug product that is not an abuse-deterrent opioid analgesic	78
drug product in order to access an abuse-deterrent opioid	79
analgesic drug product.	80
(D) This section shall not be construed to prevent a	81
health insuring corporation from applying utilization review	82
measures to abuse-deterrent opioid analgesic drug products,	83
including prior authorization requirements or nonopioid	84
analgesic drug step therapy, provided that the same utilization	85
review measures are applied to all opioid analgesic drug	86
products.	87
(E) If a health insuring corporation measures the	88
efficiency, quality of care, or clinical performance of a	89
prescriber, including through the use of patient satisfaction	90
surveys, it shall not penalize the prescriber, financially or	91
otherwise, for either of the following actions:	92
(1) Prescribing an abuse-deterrent opioid analgesic drug	93
product;	94
(2) Deciding not to prescribe any opioid analgesic drug	95
product.	96
Sec. 3923.851. (A) As used in this section:	97
(1) "Abuse-deterrent opioid analgesic drug product" means	98
a brand or generic opioid analgesic drug product approved by the	99
United States food and drug administration with abuse-deterrence	100
labeling claims indicating its abuse-deterrent properties are	101
expected to deter or reduce its abuse.	102
(2) "Opioid analgesic" has the same meaning as in section	103
3719.01 of the Revised Code.	104

Page 4

(3) "Prescriber" has the same meaning as in section	105
4729.01 of the Revised Code.	105
	100
(B) Notwithstanding section 3901.71 of the Revised Code,	107
an individual or group policy of sickness and accident insurance	108
or a public employee benefit plan that is delivered, issued for	109
delivery, or renewed in this state and covers opioid analgesic	110
drug products as part of providing any coverage of prescription	111
drugs shall provide access to abuse-deterrent opioid analgesic	112
drug products in the drug formulary or other list of covered	113
drugs that applies under the policy or plan.	114
(C) Both of the following apply to any prior authorization	115
requirements or utilization review measures contained in a	116
sickness and accident insurance policy or public employee	117
benefit plan subject to this section and any coverage denials	118
made pursuant to those requirements or measures with respect to	119
made parbaane to enobe requiremente or medbared with respect to	± ± 2
opioid analgesic drug products:	120
· · · · · · · · · · · · · · · · · · ·	-
opioid analgesic drug products:	120
opioid analgesic drug products: (1) Prior authorization requirements or utilization review	120 121
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent	120 121 122
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent opioid analgesic drug products than for opioid analgesic drug	120 121 122 123
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent opioid analgesic drug products than for opioid analgesic drug products that are not abuse-deterrent opioid analgesic drug	120 121 122 123 124
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent opioid analgesic drug products than for opioid analgesic drug products that are not abuse-deterrent opioid analgesic drug products.	120 121 122 123 124 125
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent opioid analgesic drug products than for opioid analgesic drug products that are not abuse-deterrent opioid analgesic drug products. (2) Prior authorization requirements or utilization review	120 121 122 123 124 125 126
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent opioid analgesic drug products than for opioid analgesic drug products that are not abuse-deterrent opioid analgesic drug products. (2) Prior authorization requirements or utilization review measures shall not require treatment with an opioid analgesic	120 121 122 123 124 125 126 127
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent opioid analgesic drug products than for opioid analgesic drug products that are not abuse-deterrent opioid analgesic drug products. (2) Prior authorization requirements or utilization review measures shall not require treatment with an opioid analgesic drug product that is not an abuse-deterrent opioid analgesic	120 121 122 123 124 125 126 127 128
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent opioid analgesic drug products than for opioid analgesic drug products that are not abuse-deterrent opioid analgesic drug products. (2) Prior authorization requirements or utilization review measures shall not require treatment with an opioid analgesic drug product that is not an abuse-deterrent opioid analgesic drug product in order to access an abuse-deterrent opioid	120 121 122 123 124 125 126 127 128 129
opioid analgesic drug products: (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent opioid analgesic drug products than for opioid analgesic drug products that are not abuse-deterrent opioid analgesic drug products. (2) Prior authorization requirements or utilization review measures shall not require treatment with an opioid analgesic drug product that is not an abuse-deterrent opioid analgesic drug product in order to access an abuse-deterrent opioid analgesic drug product.	120 121 122 123 124 125 126 127 128 129 130

opioid analgesic drug products, including prior authorization	134
requirements or nonopioid analgesic drug step therapy, provided	135
that the same utilization review measures are applied to all	136
opioid analgesic drug products.	137
(E) If a sickness and accident insurer or public employee	138
benefit plan measures the efficiency, quality of care, or	139
clinical performance of a prescriber, including through the use	140
of patient satisfaction surveys, it shall not penalize the	141
prescriber, financially or otherwise, for either of the	142
following actions:	143
(1) Prescribing an abuse-deterrent opioid analgesic drug	144
product;	145
(2) Deciding not to prescribe any opioid analgesic drug	146
product.	147
<u>producer</u>	± 1 /
Sec. 5164.091. (A) As used in this section:	148
Sec. 5164.091. (A) As used in this section:	148
Sec. 5164.091. (A) As used in this section:	148 149
Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the	148 149 150
Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the United States food and drug administration with abuse-deterrence	148 149 150 151
Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the United States food and drug administration with abuse-deterrence labeling claims indicating its abuse-deterrent properties are	148 149 150 151 152
Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the United States food and drug administration with abuse-deterrence labeling claims indicating its abuse-deterrent properties are expected to deter or reduce its abuse.	148 149 150 151 152 153
Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the United States food and drug administration with abuse-deterrence labeling claims indicating its abuse-deterrent properties are expected to deter or reduce its abuse. (2) "Opioid analgesic" has the same meaning as in section	148 149 150 151 152 153 154
<pre>Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the United States food and drug administration with abuse-deterrence labeling claims indicating its abuse-deterrent properties are expected to deter or reduce its abuse. (2) "Opioid analgesic" has the same meaning as in section 3719.01 of the Revised Code.</pre>	148 149 150 151 152 153 154 155
Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the United States food and drug administration with abuse-deterrence labeling claims indicating its abuse-deterrent properties are expected to deter or reduce its abuse. (2) "Opioid analgesic" has the same meaning as in section 3719.01 of the Revised Code. (3) "Prescriber" has the same meaning as in section.	148 149 150 151 152 153 154 155 156
Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the United States food and drug administration with abuse-deterrence labeling claims indicating its abuse-deterrent properties are expected to deter or reduce its abuse. (2) "Opioid analgesic" has the same meaning as in section 3719.01 of the Revised Code. (3) "Prescriber" has the same meaning as in section 4729.01 of the Revised Code.	148 149 150 151 152 153 154 155 156 157
Sec. 5164.091. (A) As used in this section: (1) "Abuse-deterrent opioid analgesic drug product" means a brand or generic opioid analgesic drug product approved by the United States food and drug administration with abuse-deterrence labeling claims indicating its abuse-deterrent properties are expected to deter or reduce its abuse. (2) "Opioid analgesic" has the same meaning as in section 3719.01 of the Revised Code. (3) "Prescriber" has the same meaning as in section 4729.01 of the Revised Code. (B) With respect to the medicaid program's coverage of	148 149 150 151 152 153 154 155 156 157 158

Page 6

162 the program. (C) Both of the following apply to any prior authorization 163 requirements or utilization review measures under the medicaid 164 program and any coverage denials made pursuant to those 165 requirements or measures with respect to opioid analgesic drug 166 167 products: 168 (1) Prior authorization requirements or utilization review measures shall not be any more restrictive for abuse-deterrent 169 opioid analgesic drug products than for opioid analgesic drug 170 products that are not abuse-deterrent. 171 (2) Prior authorization requirements or utilization review 172 measures shall not require treatment with an opioid analgesic 173 drug product that is not an abuse-deterrent opioid analgesic 174 drug product in order to access an abuse-deterrent opioid 175 analgesic drug product. 176 (D) This section shall not be construed to prevent the 177 department from applying utilization review measures to abuse-178 deterrent opioid analgesic drug products, including prior 179 authorization requirements or nonopioid analgesic drug step 180 therapy, provided that the same utilization review measures are 181 applied to all opioid analgesic drug products. 182 (E) If the department measures the efficiency, quality of 183 care, or clinical performance of a prescriber, including through 184 the use of patient satisfaction surveys, it shall not penalize 185 the prescriber, financially or otherwise, for either of the 186 following actions: 187 (1) Prescribing an abuse-deterrent opioid analgesic drug 188 product; 189

(2) Deciding not to prescribe any opioid analgesic drug 190

product.

Sec. 5167.12. (A) When contracting under section 5167.10 192 of the Revised Code with a managed care organization that is a 193 health insuring corporation, the department of medicaid shall 194 require the health insuring corporation to provide coverage of 195 prescribed drugs for medicaid recipients enrolled in the health 196 insuring corporation. In providing the required coverage, the 197 health insuring corporation may use strategies for the 198 management of drug utilization, subject to the department's 199 approval-and, the limitations specified in division (B) of this 200 section, use strategies for the management of drug utilization 201 and the requirements specified in division (C) of this section. 202

(B) The department shall not permit a health insuring203corporation to impose a prior authorization requirement in the204case of a drug to which all of the following apply:205

(1) The drug is an antidepressant or antipsychotic. 206

(2) The drug is administered or dispensed in a standard
207
tablet or capsule form, except that in the case of an
antipsychotic, the drug also may be administered or dispensed in
a long-acting injectable form.

(3) The drug is prescribed by either of the following:

(a) A physician whom the health insuring corporation,
pursuant to division (C) of section 5167.10 of the Revised Code,
has credentialed to provide care as a psychiatrist;
214

(b) A psychiatrist practicing at a community mental health
 services provider certified by the department of mental health
 and addiction services under section 5119.36 of the Revised
 Code.

191

211

(4) The drug is prescribed for a use that is indicated on 219 the drug's labeling, as approved by the federal food and drug 220 administration. 221 (C) The department shall require a health insuring_ 222 corporation to comply with the requirements of section 5164.091 223 of the Revised Code as if the health insuring corporation were 224 the department. 225 226 (D) The department shall permit <u>authorize</u> a health 227 insuring corporation to develop and implement a pharmacy utilization management program under which prior authorization 228 through the program is established as a condition of obtaining a 229 controlled substance pursuant to a prescription. The 230 department's authorization under this division does not affect a 231 health insuring corporation's obligation to comply with the 232 prior authorization procedures that apply as a result of 233 division (C) of this section. 234 Section 2. That existing sections 1739.05 and 5167.12 of 235 the Revised Code are hereby repealed. 236 Section 3. Sections 1739.05 and 1751.691 of the Revised 237 238 Code, as amended or enacted by this act, apply only to arrangements, policies, contracts, and agreements that are 239 created, delivered, issued for delivery, or renewed in this 240 state on or after January 1, 2017. Section 3923.851 of the 241 Revised Code, as enacted by this act, applies only to policies 242 of sickness and accident insurance that are delivered, issued 243 for delivery, or renewed in this state on or after January 1, 244 2017, and only to public employee benefit plans that are 245 established or modified in this state on or after January 1, 246 2017. Sections 5164.091 and 5167.12 of the Revised Code, as 247 amended or enacted by this act, apply to the Medicaid program 248

beginning January 1, 2017, and to contracts that the Department	249
of Medicaid and Medicaid managed care organizations enter into	250
or renew on or after January 1, 2017.	251