

1 SENATE BILL 569

2 **50TH LEGISLATURE - STATE OF NEW MEXICO - FIRST SESSION, 2011**

3 INTRODUCED BY

4 Eric G. Griego

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10 AN ACT

11 RELATING TO OPIOID ABUSE PREVENTION AND TREATMENT; ENACTING A
12 NEW SECTION OF THE NEW MEXICO DRUG, DEVICE AND COSMETIC ACT TO
13 PROVIDE FOR THE ESTABLISHMENT OF A PRESCRIPTION DRUG MONITORING
14 PROGRAM TO MONITOR THE PRESCRIBING AND DISPENSING OF OPIATES;
15 MANDATING THAT HEALTH FACILITIES THAT RECEIVE FINANCIAL
16 ASSISTANCE PURSUANT TO THE RURAL PRIMARY HEALTH CARE ACT HAVE
17 PHYSICIANS CERTIFIED AND AVAILABLE TO PROVIDE MEDICALLY
18 ASSISTED TREATMENT FOR OPIOID ADDICTION; REQUIRING THE HUMAN
19 SERVICES DEPARTMENT TO ESTABLISH A PROGRAM TO PROVIDE FREE
20 VOUCHERS TO INDIGENT PERSONS TO RECEIVE MEDICALLY ASSISTED
21 TREATMENT FOR OPIOID ADDICTION; ENACTING NEW SECTIONS OF THE
22 HEALTH CARE PURCHASING ACT AND THE NEW MEXICO INSURANCE CODE TO
23 PROVIDE THAT NO PRIOR AUTHORIZATION BE REQUIRED BY HEALTH
24 COVERAGE ENTITIES FOR THE PROVISION OF BUPRENORPHINE TO TREAT
25 OPIOID ADDICTION; MAKING AN APPROPRIATION.

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BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:

SECTION 1. A new section of the New Mexico Drug, Device and Cosmetic Act is enacted to read:

"[NEW MATERIAL] PRESCRIPTION MONITORING--RULEMAKING.--

A. The board shall establish and maintain a prescription drug monitoring program to monitor the prescribing and dispensing of opiates by practitioners in the state and to research the prescribing and dispensing of opiates. The board shall promulgate any rules necessary to implement the prescription drug monitoring program. The prescription drug monitoring program shall not interfere with the legal use of opiates. The prescription drug monitoring program shall be used to provide information to practitioners, dispensers and patients to help avoid the illegal use of controlled substances. This information shall include patient utilization reports prepared pursuant to board rules, which reports patient utilization of Schedule II or III controlled substances.

B. Before writing a prescription for a Schedule II or III controlled substance for a patient about whom the practitioner has a reasonable belief that the patient may be seeking the controlled substance for any reason other than the treatment of an existing medical condition for which it was prescribed, a practitioner, or other person authorized by the practitioner, shall obtain a patient utilization report

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1 regarding that patient's history for the preceding twelve
2 months. The practitioner shall review the patient utilization
3 report to assess whether the prescription for the controlled
4 substance is necessary and appropriate.

5 C. The board shall require an applicant for a
6 license to prescribe controlled substances to register in the
7 prescription drug monitoring program and agree to participate
8 in the program by providing and monitoring their patients'
9 controlled substance use in accordance with the provisions of
10 this section.

11 D. The board may issue a waiver from the provisions
12 of this section to a practitioner who is unable to access
13 patient utilization reports by electronic means. A
14 practitioner who is unable to access patient utilization
15 reports by electronic means shall obtain a waiver from the
16 board on an annual basis, until the practitioner is able to
17 access patient utilization reports by electronic means.

18 E. Unless a court of competent jurisdiction makes a
19 finding of gross negligence, malice or criminal intent, neither
20 the board nor any practitioner, dispenser or any person in
21 proper possession of information pursuant to the provisions of
22 this section shall be subject to civil liability,
23 administrative action or other legal or equitable relief for
24 any of the following acts or omissions:

25 (1) furnishing information pursuant to the

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1 provisions of this section;

2 (2) receiving, using or relying upon or not
3 using or relying upon information received pursuant to this
4 section;

5 (3) not furnishing information to the board;
6 or

7 (4) furnishing information that was factually
8 incorrect.

9 F. As used in this section:

10 (1) "dispenser" means a person that the board
11 has authorized to dispense or distribute to the ultimate user a
12 controlled substance or dangerous drug; "dispenser" does not
13 mean a pharmacy that dispenses or distributes any controlled
14 substance or dangerous drug for the purposes of:

15 (a) inpatient care;

16 (b) emergency department care for the
17 immediate use of a controlled substance; or

18 (c) discharge of a patient from a health
19 facility, where the pharmacy provides the patient with a supply
20 of the drug intended for use within a maximum of seventy-two
21 hours;

22 (2) "pharmacy" means a licensed place of
23 business where drugs are compounded or dispensed and
24 pharmaceutical care is provided; and

25 (3) "Schedule II or III controlled substance"

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1 means a drug or substance listed in Schedules II and III of the
2 Controlled Substances Act."

3 SECTION 2. A new section of the Health Care Purchasing
4 Act is enacted to read:

5 "[NEW MATERIAL] NO PRIOR AUTHORIZATION REQUIREMENT FOR
6 BUPRENORPHINE.--Group health coverage, including any form of
7 self-insurance, offered, issued or renewed under the Health
8 Care Purchasing Act that provides coverage for medically
9 assisted treatment of opioid addiction using buprenorphine
10 shall not require a prior authorization before providing access
11 to buprenorphine. Nothing in this section shall be construed
12 to preclude a group coverage provider from monitoring the
13 prescribing of buprenorphine through audits or other means."

14 SECTION 3. A new section of the Rural Primary Health Care
15 Act is enacted to read:

16 "[NEW MATERIAL] STATE-FUNDED HEALTH FACILITIES--
17 REQUIREMENT FOR BUPRENORPHINE MEDICALLY ASSISTED OPIOID
18 ADDICTION TREATMENT.--A health facility that has at least two
19 full-time-equivalent physicians on staff and that receives
20 financial assistance or employs any staff member who receives
21 financial assistance pursuant to the Rural Primary Health Care
22 Act shall have on staff at least one physician who is certified
23 and available to provide buprenorphine medically assisted
24 treatment for opioid addiction."

25 SECTION 4. A new section of the Public Assistance Act is

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1 enacted to read:

2 "[NEW MATERIAL] OPIOID ADDICTION--VOUCHER PROGRAM FOR
3 MEDICALLY ASSISTED TREATMENT.--The department shall establish
4 and implement a program whereby the department provides to
5 uninsured individuals with incomes below two hundred percent of
6 the federal poverty level vouchers to permit the bearer to
7 receive free medically assisted treatment for opioid addiction
8 when prescribed by a physician who is certified to prescribe
9 buprenorphine. The vouchers shall fund buprenorphine treatment
10 for a total duration of six months or less for each patient,
11 with a total daily dose not to exceed sixteen milligrams."

12 SECTION 5. A new section of Chapter 59A, Article 22 NMSA
13 1978 is enacted to read:

14 "[NEW MATERIAL] NO PRIOR AUTHORIZATION REQUIREMENT FOR
15 BUPRENORPHINE.--An individual or group health insurance policy,
16 health care plan or certificate of health insurance that is
17 delivered, issued for delivery or renewed in this state that
18 provides coverage for medically assisted treatment of opioid
19 addiction using buprenorphine shall not require a prior
20 authorization before providing access to buprenorphine.
21 Nothing in this section shall be construed to preclude an
22 insurer from monitoring the prescribing of buprenorphine
23 through audits or other means."

24 SECTION 6. A new section of Chapter 59A, Article 23 NMSA
25 1978 is enacted to read:

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1 "[NEW MATERIAL] NO PRIOR AUTHORIZATION REQUIREMENT FOR
2 BUPRENORPHINE.--A blanket or group health insurance policy or
3 contract that is delivered, issued for delivery or renewed in
4 this state that provides coverage for medically assisted
5 treatment of opioid addiction using buprenorphine shall not
6 require a prior authorization before providing access to
7 buprenorphine. Nothing in this section shall be construed to
8 preclude an insurer from monitoring the prescribing of
9 buprenorphine through audits or other means."

10 **SECTION 7.** A new section of the Health Maintenance
11 Organization Law is enacted to read:

12 "[NEW MATERIAL] NO PRIOR AUTHORIZATION REQUIREMENT FOR
13 BUPRENORPHINE.--An individual or group health maintenance
14 organization contract that is delivered, issued for delivery or
15 renewed in this state that provides coverage for medically
16 assisted treatment of opioid addiction using buprenorphine
17 shall not require a prior authorization before providing access
18 to buprenorphine. Nothing in this section shall be construed
19 to preclude a health maintenance organization from monitoring
20 the prescribing of buprenorphine through audits or other
21 means."

22 **SECTION 8.** A new section of Chapter 59A, Article 47 NMSA
23 1978 is enacted to read:

24 "[NEW MATERIAL] NO PRIOR AUTHORIZATION REQUIREMENT FOR
25 BUPRENORPHINE.--An individual or group health insurance policy,

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1 health care plan or certificate of health insurance that is
2 delivered, issued for delivery or renewed in this state that
3 provides coverage for medically assisted treatment of opioid
4 addiction using buprenorphine shall not require a prior
5 authorization before providing access to buprenorphine.
6 Nothing in this section shall be construed to preclude a health
7 care plan issuer from monitoring the prescribing of
8 buprenorphine through audits or other means."

9 SECTION 9. APPROPRIATION.--Five million dollars
10 (\$5,000,000) is appropriated from the general fund to the human
11 services department for expenditure in fiscal year 2012 to fund
12 a program that provides vouchers for medically assisted
13 treatment for opioid addiction to individuals who are indigent
14 and uninsured when prescribed by a physician who is certified
15 to prescribe buprenorphine. Any unexpended or unencumbered
16 balance remaining at the end of fiscal year 2012 shall revert
17 to the general fund.