



## **New Georgia Law pertaining to use of the Prescription Drug Monitoring Program (PDMP)- Enrollment Required by January 1, 2018**

The purpose of this communication is to inform you of a new Georgia law requiring use of the Prescription Drug Monitoring Program. This new law is a result of bill HB 249 which Governor Deal signed on May 4, 2017. It impacts all Georgia physicians / prescribers requiring use of the Prescription Drug Monitoring Program (PDMP). The bill was designed to address opioid abuse in the state. The resulting law requires reviewing PDMP information on a patient when prescribing a controlled substance (see list below\*) or a benzodiazepine beginning July 1, 2018.

### DPH Transition

H.B. 249 moves the Prescription Drug Monitoring Program (PDMP) from the Georgia Drugs and Narcotics Agency to the Georgia Department of Public Health (DPH).

### **Mandatory PDMP Enrollment**

**By January 1, 2018, every prescriber in Georgia who has a DEA registration number must enroll as a PDMP user. If a prescriber obtains a DEA license after January 1, 2018, they must enroll with the PDMP within 30 days.**

### PDMP Review Delegation

Under H.B. 249, a prescriber may delegate their authority to check the PDMP to two members of their medical practice staff. For a prescriber to delegate this authority to an unlicensed or unregistered staff member (i.e., an office manager or medical assistant), the staff member must submit to an annual registration

process that will be administered by the Georgia Board of Pharmacy. A health care facility (e.g., hospital or ambulatory surgery center) may select two employees to serve as delegates per shift or rotation. At hospitals that provide emergency services, each prescriber may designate two individuals who are employed by that hospital per shift or rotation.

**Any unauthorized use of PDMP data by a delegate can result in civil or criminal liability for the prescriber. Delegates may only use PDMP data for the purpose of providing medical care or to inform the prescriber of a patient's potential use, misuse, abuse or underutilization of a prescribed medication.** The delegation of your access to the PDMP has potential liability so if you chose to delegate, please choose your delegates carefully and with clarity of responsibility for the authorized use of accessed information.

### **Mandatory PDMP Review**

Between January 1, 2018 and May 31, 2018, the DPH must randomly test the PDMP to confirm it is accessible and operational 99.5 percent of the time.

Once DPH has certified that the PDMP's accessibility and operational standards have been met, the prescriber or their delegate must review the information from the PDMP when prescribing a controlled substance (see list below\*) or a benzodiazepine beginning July 1, 2018. This review is limited to the first time the prescriber issues a prescription for the given patient and at least every 90 days thereafter.

\* Source' paragraph (1) or (2) of Code Section 16-13-26 Paragraph 1-2

### **Exemptions from the requirement to check the database are as follows...**

- Prescriptions for no more than a three-day supply of a covered substance and no more than 26 pills
  
- The patient is in a hospital or health care facility, including – but not limited to – a nursing home, an intermediate care home, a personal care home, or a hospice program that provides patient care and whereby the prescriptions are to be administered and used by a patient on the premises of the facility

- The patient has had outpatient surgery at a hospital or ambulatory surgical center and the prescription is for no more than a 10-day supply of a covered substance and no more than 40 pills
- The patient is terminally ill or under the supervised care of an outpatient hospice program
- The patient is receiving treatment for cancer

### **Medical Record Notations**

**Prescribers or their delegate must make a notation in the patient's medical record that the PDMP was consulted and identify the individual who conducted the PDMP patient search. If the PDMP does not allow the prescriber/delegate to gain access to the patient's information for any reason, the prescriber/delegate should note the time and date and the prescriber/delegate's name in the patient's medical record.**

**Prescribers may now include PDMP prescription information in a patient's electronic health or medical record.**

### **GCMB Accountability**

If a prescriber fails to check the PDMP as outlined above, he or she will be held administratively accountable to the Georgia Medical Composite Board; however, the prescriber may not be held civilly liable or criminally responsible.

### **Required Prescribing Information**

When prescribing an opiate, opioid, opioid analgesic, or opioid derivative, the prescriber must provide the patient with information on the drug's addictive risks and the options that are available for safely disposing of any unused medications. This information may be provided in either verbal or written form.

### **Neonatal Abstinence Syndrome Reporting**

A health care provider, coroner, or medical examiner must report all incidents of neonatal abstinence syndrome to DPH, which will submit an annual report – including findings and recommendations on how to reduce the number of infants

born with neonatal abstinence syndrome – to the president of the Georgia Senate, the speaker of the Georgia House of Representatives, and the chairs of both the Georgia House and Senate Health and Human Services committees.

### **Other Provisions**

Other H.B. 249 provisions include codifying Gov. Deal’s executive order to make naloxone available on an over-the-counter basis, annual inspections of narcotic treatment programs, and a requirement that calls for the Department of Community Health and the Department of Behavioral Health and Developmental Disabilities to report the number of patients who are enrolled in or have been discharged from a treatment program.

*O.C.G.A. § 16-13-26*

GEORGIA CODE  
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\*\*\* Current Through the 2016 Regular Session \*\*\*

TITLE 16. CRIMES AND OFFENSES  
CHAPTER 13. CONTROLLED SUBSTANCES  
ARTICLE 2. REGULATION OF CONTROLLED SUBSTANCES  
PART 1. SCHEDULES, OFFENSES, AND PENALTIES

O.C.G.A. § 16-13-26 (2016)

### **§ 16-13-26. Schedule II**

**The controlled substances listed in this Code section are included in Schedule II:**

(1) Any of the following substances, or salts thereof, except those narcotic drugs specifically exempted or listed in other schedules, whether produced directly or indirectly by extraction from substances of vegetable origin, or independently by extraction from substances of vegetable origin, or independently by means of chemical synthesis, or by combination of extraction and chemical synthesis:

(A) Opium and opiate, and any salt, compound, derivative, or preparation of opium or opiate, excluding naloxone hydrochloride, but including the following:

- (i) Raw opium;
- (ii) Opium extracts;
- (iii) Opium fluid extracts;
- (iv) Powdered opium;
- (v) Granulated opium;
- (vi) Tincture of opium;
- (vii) Codeine;
- (viii) Ethylmorphine;
- (ix) Hydrocodone;
- (x) Hydromorphone;
- (xi) Metopon;
- (xii) Morphine;
- (xiii) Oripavine;
- (xiv) Oxycodone;
- (xv) Oxymorphone;
- (xvi) Thebaine;

(B) Any salt, compound, isomer, derivative, or preparation thereof which is chemically equivalent or identical with any of the substances referred to in subparagraph (A) of this paragraph, except that these substances shall not include the isoquinoline alkaloids of opium;

(C) Opium poppy and poppy straw;

(D) Cocaine, coca leaves, any salt, compound, derivative, stereoisomers of cocaine, or preparation of coca leaves, and any salt, compound, derivative, stereoisomers of cocaine, or preparation thereof which is chemically equivalent or identical with any of these substances, but not including decocainized coca leaves or extractions which do not contain cocaine or ecgonine;

(2) Any of the following opiates, including their isomers, esters, ethers, salts, and salts of isomers, whenever the existence of these isomers, esters, ethers, and salts is possible within the specific chemical designation:

(A) Alfentanil;

(A.1) Alphaprodine;

(B) Anileridine;

(C) Bezitramide;

(D) Dihydrocodeine;

(E) Diphenoxylate;

(F) Fentanyl;

(G) Isomethadone;

(G.5) Levo-alphacetylmethadol (some other names: levomethadyl acetate, LAAM);

(H) Levomethorphan;

(I) Levorphanol;

(J) Methazocine;

(K) Methadone;

(L) Methadone-Intermediate, 4-cyano-2-dimethylamino-4, 4-diphenyl butane;

(M) Moramide-Intermediate, 2-methyl-3-morpholino-1, 1-diphenylpropane-carboxylic acid;

(N) Pethidine (meperidine);

(O) Pethidine-Intermediate-A, 4-cyano-1-methyl-4-phenylpiperidine;

(P) Pethidine-Intermediate-B, ethyl-4-phenylpiperidine-4-carboxylate;

(Q) Pethidine-Intermediate-C, 1-methyl-4-phenylpiperidine-4-carboxylic acid;

(R) Phenazocine;

(S) Piminodine;

(T) Racemethorphan;

(U) Racemorphan;

(U.1) Remifentanil;

(V) Sufentanil;

(V.1) Tapentadol;

(W) 4-anilino-N-phenethyl-4-piperidine (ANPP);

References;

HB 249

Georgia Code Section 16-13-26, 1,2.

MAG/ May 2017