1	ENGROSSED HOUSE
2	BILL NO. 2931 By: Mulready of the House
2	and
3	Griffin of the Senate
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7	An Act relating to controlled dangerous substances; amending 63 O.S. 2011, Section 2-309, as last amended
8	by Section 1, Chapter 323, O.S.L. 2013 (63 O.S. Supp. 2017, Section 2-309), which relates to the Uniform
9	Controlled Dangerous Substances Act; requiring electronic prescribing for all scheduled drugs;
10	providing exceptions; modifying time period for certain exception; deleting prohibition concerning
11	hydrocodone refills and restrictions on dispensing or distributing Schedule V substances; deleting
12	restrictions related to the dispensing of paregoric; modifying certain definition; directing counties with
13	certain populations to comply with electronic prescription requirements by certain date; and
14	providing an effective date.
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16	
17	BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:
18	SECTION 1. AMENDATORY 63 O.S. 2011, Section 2-309, as
19	last amended by Section 1, Chapter 323, O.S.L. 2013 (63 O.S. Supp.
20	2017, Section 2-309), is amended to read as follows:
21	Section 2-309. A. 1. Except for dosages medically required
22	for a period not to exceed forty-eight (48) hours which are
23	administered by or on direction of a practitioner, other than a
24	pharmacist, or medication dispensed directly by a practitioner,

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other than a pharmacist, to an ultimate user, no controlled 1 2 dangerous substance included in Schedule II, which is a prescription 3 drug as determined under regulation promulgated by the Board of 4 Pharmacy, may shall be dispensed without the written an electronic 5 prescription of a practitioner; provided, that in emergency situations, as prescribed by the Board of Pharmacy by regulation, 6 7 such drug may be dispensed upon oral prescription reduced promptly 8 to writing and filed by the pharmacist in a manner to be prescribed 9 by rules and regulations of the Director of the Oklahoma State 10 Bureau of Narcotics and Dangerous Drugs Control.

2. Electronic prescribing may <u>shall</u> be utilized for Schedules
 II, III, IV, and V, subject to the requirements set forth in 21 CFR,
 Section 1311 et seq.

14 3. The transmission of written prescription by practitioner to 15 dispensing pharmacy by facsimile or electronic transmission with 16 electronic signature is permitted only under the following 17 conditions:

18	a.	for Schedule II drugs, the original prescription must
19		be presented and verified against the facsimile at the
20		time the substances are actually dispensed, and the
21		original document must be properly annotated and
22		retained for filing, except:
23		(1) home infusion pharmacy may consider the facsimile
24		to be a "written prescription" as required by

1		Section 2-101 et seq. of this title and as
2		required by Title 21 U.S.C., Section 829(a). The
3		facsimile copy of the prescription shall be
4		retained as an original prescription, and it must
5		contain all the information required by Section
6		2-101 et seq. of this title and 21 CFR, Section
7		1306.05(a), including date issued, the patient's
8		full name and address, and the practitioner's
9		name, address, DEA registration number, and
10		signature. The exception to the regulations for
11		home infusion/IV therapy is intended to
12		facilitate the means by which home infusion
13		pharmacies obtain prescriptions for patients
14		requiring the frequently modified parenteral
15		controlled release administration of narcotic
16		substances, but does not extend to the dispensing
17		of oral dosage units of controlled substances,
18	(2)	the same exception is granted to patients in Long
19		Term Care facilities (LTCF), which are filled by
20		and delivered to the facility by a dispensing
21		pharmacy, and
22	(3)	an An electronic prescription with electronic
23		signature may serve as an original prescription,
24		

2Section 1311 et seq.,-and3b. for drugs in Schedules III and IV, a facesimile copy of4a written, signed preseription transmitted directly by5the preseribing practitioner to the pharmacy can serve6as an original preseription. Electronic preseribing7may be utilized for Schedules III and IV subject to8the same requirements as set forth in 21 CFR, Section91311 et seq.104. Prescriptions shall be retained in conformity with the11requirements of this section and Section 2-307 of this title. No12prescription for a Schedule II substance may be refilled.135. The electronic prescription requirement provided for in this14section shall not apply to prescriptions for controlled dangerous15substances issued by any of the following:16a. a person licensed to practice veterinary medicine,17b. a practitioner who experiences temporary technological18or electrical failure or other extenuating19circumstance that prevents the prescription from being10transmitted electronically; provided, however, that18the practitioner documents the reason for this19circumstance that prevents the prescription from being20transmitted electronically; provided, however, that21the practitioner documents the reason for this22exception in the medical record of the patient,23c. a practitioner, other than a pharmacist, who dispenses24directly to an ultimate user, <th>1</th> <th></th> <th>subject to the requirements set forth in 21 CFR,</th>	1		subject to the requirements set forth in 21 CFR,
4 a written, signed prescription transmitted directly by 5 the prescribing practitioner to the pharmacy can serve 6 as an original prescription. Electronic prescribing 7 may be utilized for Schedulco III and IV oubject to 8 the same requirements as set forth in 21 CFR, Section 9 1311 et seq. 10 4. Prescriptions shall be retained in conformity with the 11 requirements of this section and Section 2-307 of this title. No 12 prescription for a Schedule II substance may be refilled. 13 5. The electronic prescription requirement provided for in this 14 section shall not apply to prescriptions for controlled dangerous 15 substances issued by any of the following: 16 a. a person licensed to practice veterinary medicine, 17 b. a practitioner who experiences temporary technological 18 or electrical failure or other extenuating 19 circumstance that prevents the prescription from being 18 or electrical failure or other extenuating 19 circumstance that prevents the prescription from being 20 transmitted electronically; provided, however, that 21 the practitioner, ot	2		Section 1311 et seq. , and
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19circumstance that prevents the prescription from being20transmitted electronically; provided, however, that21the practitioner documents the reason for this22exception in the medical record of the patient,23c.a practitioner, other than a pharmacist, who dispenses	17	<u>b.</u>	a practitioner who experiences temporary technological
20transmitted electronically; provided, however, that21the practitioner documents the reason for this22exception in the medical record of the patient,23c.a practitioner, other than a pharmacist, who dispenses	18		or electrical failure or other extenuating
21 <u>the practitioner documents the reason for this</u> 22 <u>exception in the medical record of the patient,</u> 23 <u>c. a practitioner, other than a pharmacist, who dispenses</u>	19		circumstance that prevents the prescription from being
22 <u>exception in the medical record of the patient,</u> 23 <u>c.</u> <u>a practitioner, other than a pharmacist, who dispenses</u>	20		transmitted electronically; provided, however, that
23 <u>c.</u> <u>a practitioner</u> , other than a pharmacist, who dispenses	21		the practitioner documents the reason for this
	22		exception in the medical record of the patient,
24 <u>directly to an ultimate user</u> ,	23	<u>C.</u>	a practitioner, other than a pharmacist, who dispenses
	24		directly to an ultimate user,

1	<u>d.</u>	a practitioner who orders a controlled dangerous
2		substance to be administered in a state-certified and
3		recognized hospital, nursing home, hospice facility,
4		outpatient dialysis facility, residential care
5		facility or correctional facility,
6	<u>e.</u>	a practitioner who writes a prescription to be
7		dispensed by a pharmacy located on federal property,
8		provided the practitioner documents the reason for
9		this exception in the medical record of the patient,
10		or
11	<u>f.</u>	a prescriber that has received a waiver or extension
12		from the Oklahoma State Bureau of Narcotics and
13		Dangerous Drugs Control.
14	<u>6. Elect</u>	ronic prescriptions shall not be utilized under the
15	following cir	cumstances:
16	<u>a.</u>	prescriptions that have complicated directions,
17	<u>b.</u>	prescriptions that have directions that exceed one
18		hundred forty characters,
19		
20	<u>c.</u>	compound prescriptions containing two or more
20	<u>c.</u>	compound prescriptions containing two or more commercially available products or two or more active
	<u>c.</u>	
20	<u>c.</u> <u>d.</u>	commercially available products or two or more active
20 21		commercially available products or two or more active pharmaceutical ingredients,

1	e. prescriptions issued under approved research
2	protocols,
3	f. prescriptions that will be dispensed out-of-state, or
4	g. if the practitioner determines that an electronic
5	prescription cannot be issued in a timely manner and
6	the condition of the patient is at risk.
7	7. A pharmacist who receives a written, oral or facsimile
8	prescription shall not be required to verify that the prescription
9	falls under one of the exceptions provided for in paragraph 6 of
10	this subsection. Pharmacists may continue to dispense medications
11	from otherwise valid written, oral or facsimile prescriptions that
12	are consistent with current laws and regulations.
13	8. Practitioners must indicate in the health record of a
14	patient that an exception to the electronic prescription requirement
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	patient that an exception to the electronic prescription requirement
15	patient that an exception to the electronic prescription requirement was utilized.
15 16	patient that an exception to the electronic prescription requirement was utilized. B. 1. Except for dosages medically required for a period not
15 16 17	<pre>patient that an exception to the electronic prescription requirement was utilized. B. 1. Except for dosages medically required for a period not to exceed forty-eight (48) seventy-two (72) hours which are</pre>
15 16 17 18	<pre>patient that an exception to the electronic prescription requirement was utilized. B. 1. Except for dosages medically required for a period not to exceed forty-eight (48) seventy-two (72) hours which are administered by or on direction of a practitioner, other than a</pre>
15 16 17 18 19	<pre>patient that an exception to the electronic prescription requirement was utilized. B. 1. Except for dosages medically required for a period not to exceed forty-eight (48) seventy-two (72) hours which are administered by or on direction of a practitioner, other than a pharmacist, or medication dispensed directly by a practitioner,</pre>
15 16 17 18 19 20	<pre>patient that an exception to the electronic prescription requirement was utilized. B. 1. Except for dosages medically required for a period not to exceed forty-eight (48) seventy-two (72) hours which are administered by or on direction of a practitioner, other than a pharmacist, or medication dispensed directly by a practitioner, other than a pharmacist, to an ultimate user, no controlled</pre>
15 16 17 18 19 20 21	<pre>patient that an exception to the electronic prescription requirement was utilized. B. 1. Except for dosages medically required for a period not to exceed forty-eight (48) seventy-two (72) hours which are administered by or on direction of a practitioner, other than a pharmacist, or medication dispensed directly by a practitioner, other than a pharmacist, to an ultimate user, no controlled dangerous substance included in Schedule III or IV, which is a</pre>

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2. A written or oral Any prescription for a controlled
 dangerous substance in Schedule III or, IV or V may not be filled or
 refilled more than six (6) months after the date thereof or be
 refilled more than five times after the date of the prescription,
 unless renewed by the practitioner.

3. A written or oral prescription for any product containing
hydrocodone with another active ingredient shall not be refilled.
C. No controlled dangerous substance included in Schedule V may
be distributed or dispensed other than for a legitimate medical or

10 scientific purpose.

11 D. Except for dosages medically required for a period not to 12 exceed forty-eight (48) hours which are administered by or on 13 direction of a practitioner, other than a pharmacist, or medication 14 dispensed directly by a practitioner, other than a pharmacist, to an 15 ultimate user, tincture opium camphorated, commonly known as 16 paregoric, may not be dispensed without a written or oral 17 prescription. The refilling of a prescription for paregoric shall 18 be unlawful unless permission is granted by the prescriber, either 19 written or oral. 20 E. Whenever it appears to the Director of the Oklahoma State

Bureau of Narcotics and Dangerous Drugs Control that a drug not considered to be a prescription drug under existing state law or regulation of the Board of Pharmacy should be so considered because of its abuse potential, the Director shall so advise the Board of

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Pharmacy and furnish to the Board all available data relevant
 thereto.

F. D. "Prescription", as used herein, means a written or, oral 3 4 or electronic order by a practitioner to a pharmacist for a 5 controlled dangerous substance for a particular patient, which specifies the date of its issue, and the full name and address of 6 7 the patient; and, if the controlled dangerous substance is prescribed for an animal, the species of the animal+, the name and 8 9 quantity of the controlled dangerous substance prescribed;, the 10 directions for use; the name and address of the owner of the animal 11 and, if written, the signature of the practitioner.

12 G. E. No person shall solicit, dispense, receive or deliver any 13 controlled dangerous substance through the mail, unless the ultimate 14 user is personally known to the practitioner and circumstances 15 clearly indicate such method of delivery is in the best interest of 16 the health and welfare of the ultimate user.

17 F. Beginning November 1, 2018, the electronic prescription 18 requirement provided for in this section shall apply to all counties 19 having more than two hundred thousand (200,000) population according 20 to the latest Federal Decennial Census. All remaining counties 21 having less than two hundred thousand (200,000) population according 22 to the latest Federal Decennial Census shall comply with the 23 electronic prescription requirement provided for in this section on 24 or before November 1, 2019.

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1	SECTION 2. This act shall become effective November 1, 2018.
2	Passed the House of Representatives the 6th day of March, 2018.
3	
4	Presiding Officer of the House
5	of Representatives
6	Passed the Senate the day of, 2018.
7	rassed the senate the day of, 2010.
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9	Presiding Officer of the Senate
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