

**2013 OK 93**  
**IN THE SUPREME COURT OF THE STATE OF OKLAHOMA**

Terry Cline, in his official capacity as )  
Oklahoma Commissioner of Health, )  
Lyle Kelsey, in his official capacity as )  
Executive Director of the Oklahoma )  
State Board of Medical Licensure and )  
Supervision, Catherine V. Taylor, )  
in her official capacity as the President )  
of the Oklahoma State Board of )  
Osteopathic Examiners, )

Petitioners, )

v. )

Oklahoma Coalition for Reproductive )  
Justice, on behalf of itself and its )  
Members and Nova Health Systems, )  
d/b/a Reproductive Services, on )  
behalf of itself, its staff, and its patients, )

Respondents. )

**FILED**  
**SUPREME COURT**  
**STATE OF OKLAHOMA**

OCT 29 2013

MICHAEL S. RICHIE  
CLERK OF  
THE APPELLATE COURTS

Case No. 111,939

**FOR OFFICIAL  
PUBLICATION**

**CERTIFIED QUESTIONS OF LAW FROM THE  
SUPREME COURT OF THE UNITED STATES**

¶ 0 On December 4, 2012, this Court issued a memorandum opinion, finding House Bill 1970, 2011 Okla. Sess. Laws 1276, facially unconstitutional pursuant to the U.S. Supreme Court's decision in Planned Parenthood v. Casey, 505 U.S. 833 (1992). See Okla. Coal. for Reprod. Justice v. Cline, 2012 OK 102, 292 P.3d 27. The Attorney General filed a Petition for Certiorari with the U.S. Supreme Court on March 4, 2013. On June 27, 2013, the U.S.

Supreme Court granted certiorari in the case and certified two questions of law to the Supreme Court of Oklahoma.

### **CERTIFIED QUESTIONS ANSWERED**

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## **PER CURIAM**

¶ 1 The Supreme Court of the United States certified two questions of Oklahoma law under the Revised Uniform Certification of Questions of Law Act, 20 O.S. 2011 §§ 1601–1611:

Whether H.B. No. 1970, Section 1, Chapter 216, O.S.L. 2011 prohibits: (1) the use of misoprostol to induce abortions, including the use of misoprostol in conjunction with mifepristone according to a protocol approved by the Food and Drug Administration; and (2) the use of methotrexate to treat ectopic pregnancies.

We answer both certified questions in the affirmative.

### ***Procedural Background***

¶ 2 In May of 2011, the Governor signed House Bill 1970, 2011 Okla. Sess. Laws 1276, into law.<sup>1</sup> The Respondents challenged the bill in Oklahoma

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<sup>1</sup> Section 1, Subsection C, of H.B. 1970 provides:

C. No physician who provides RU-486 (mifepristone) or any abortion-inducing drug shall knowingly or recklessly fail to provide or prescribe the RU-486 (mifepristone) or any abortion-inducing drug according to the protocol tested and authorized by the U.S. Food and Drug Administration and as authorized in the drug label for the RU-486 (mifepristone) or any abortion-inducing drug.

Section 1, Subsection A, defines “abortion-inducing drug” as:

County District Court. The District Court found H.B. 1970 was unconstitutional and issued a permanent injunction, prohibiting enforcement of H.B. 1970. The Attorney General appealed the order and filed a Motion to Retain in this Court. We retained the case and issued a memorandum opinion on December 4, 2012, in Case No. 110,765, affirming the district court's decision. We found H.B. 1970 was facially unconstitutional pursuant to the U.S. Supreme Court's decision in Planned Parenthood v. Casey, 505 U.S. 833 (1992). See Okla. Coal. for Reprod. Justice v. Cline, 2012 OK 102, 292 P.3d 27. On January 15, 2013, the Chief Justice issued the mandate in Case No. 110,765.<sup>2</sup>

¶ 3 The Attorney General filed a Petition for Certiorari with the U.S. Supreme Court on March 4, 2013. The U.S. Supreme Court Clerk filed a letter in Case No. 110,765 on March 14, 2013, advising this Court that a petition for certiorari review of the order in Case No. 110,765 had been filed on March 4, 2013. The Attorney General has not asked this Court to suspend the effectiveness of mandate in Case No. 110,765.

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[A] medicine, drug, or any other substance prescribed or dispensed with the intent of terminating the clinically diagnosable pregnancy of a woman, with knowledge that the termination shall with reasonable likelihood cause the death of the unborn child. This includes off-label use of drugs known to have abortion-inducing properties, which are prescribed specifically with the intent of causing an abortion, such as misoprostol (Cytotec), and methotrexate. This definition does not apply to drugs that may be known to cause an abortion, but which are prescribed for other medical indications, such as chemotherapeutic agents or diagnostic drugs;

H.B. 1970 was codified at 63 O.S. 2011 § 1-729a.

<sup>2</sup> See Mandate, Okla. Coal. for Reprod. Justice v. Cline, No. 110,765 (Jan. 15, 2013). This Court "can take judicial notice of *its own records* in litigation interconnected with a case before it." Robinson v. Texhoma Limestone, Inc., 2004 OK 50, ¶ 13, 100 P.3d 673, 677.

¶ 4 On June 27, 2013, the U.S. Supreme Court granted certiorari in the case and certified two questions of law to this Court. See Terry Cline et al. v. Okla. Coal. for Reprod. Justice et al., No. 12-1094 (June 27, 2013). Further proceedings in the U.S. Supreme Court were reserved “pending receipt of a response from the Supreme Court of Oklahoma.” Id. The certified questions were filed in this Court on July 1, 2013, in Case No. 111,939. The briefs filed in the U.S. Supreme Court were included with the certification order. After the certified questions were filed, the Attorney General filed a request for briefing schedule. This Court entered a briefing schedule on July 16, 2013. Applications for amicus briefs were filed by several organizations, and this Court granted those applications on August 16, 2013. Briefing was completed on October 2, 2013.

***This Court Has Jurisdiction to Answer the Certified Questions***

¶ 5 Petitioners sought certiorari to the U.S. Supreme Court from Oklahoma Supreme Court Case No. 110,765, which has been mandated and is not before this Court at this time. Oklahoma Supreme Court Rule 1.16 permits a party to file a motion to suspend the effectiveness of mandate if the party contemplates the filing of a petition for certiorari in the U.S. Supreme Court and authorizes suspension of the effectiveness of the mandate until 1) expiration of time to file the petition; or 2) notice of final disposition by the U.S.

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Supreme Court.<sup>3</sup> Until a party makes a request to suspend the mandate pursuant to Rule 1.16 in Case No. 110,765, or upon final disposition by the U.S. Supreme Court, this Court will not suspend or recall the mandate in Case No. 110,765.<sup>4</sup>

¶ 6 The jurisdictional basis for a majority of this Court's decisions is derived from the jurisdiction conferred upon the Court by Oklahoma Constitution Article VII, § 4.<sup>5</sup> This section vests five types of jurisdiction in the Supreme Court: (1) appellate jurisdiction over all civil matters; (2) jurisdiction to

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<sup>3</sup> Okla. Sup. Ct. R. 1.16.

<sup>4</sup> Although the Attorney General's failure to move to suspend the effectiveness of mandate is not fatal to our exercise of jurisdiction in this case, litigants practicing before this Court must conform to the rules and procedures of this Court. The file also indicates that no one from the Attorney General's office has filed an entry of appearance in Case No. 111,939 as required by Oklahoma Supreme Court Rule 1.5, which provides that "[a]ll parties to any proceeding in the appellate courts shall immediately, but no later than filing the first document in the appellate court, file an Entry of Appearance on the forms set forth in Rule 1.301, by counsel or an unrepresented party representing himself or herself." Okla. Sup. Ct. R. 1.5. "When no counsel enters a formal appearance on behalf of an appellate party this Court possesses the discretion to list as counsel the lawyer who has signed and submitted a brief or motion for that party." State ex rel. Okla. Bd. of Medical Licensure and Supervision v. Pinaroc, 2002 OK 20, n.1, 46 P.3d 114, 116 n.1.

<sup>5</sup> The Oklahoma Constitution, Article VII, § 4 provides:

The appellate jurisdiction of the Supreme Court shall be co-extensive with the State and shall extend to all cases at law and in equity; except that the Court of Criminal Appeals shall have exclusive appellate jurisdiction in criminal cases until otherwise provided by statute and in the event there is any conflict as to jurisdiction, the Supreme Court shall determine which court has jurisdiction and such determination shall be final. The original jurisdiction of the Supreme Court shall extend to a general superintendent control over all inferior courts and all Agencies, Commissions and Boards created by law. The Supreme Court, Court of Criminal Appeals, in criminal matters and all other appellate courts shall have power to issue, hear and determine writs of habeas corpus, mandamus, quo warranto, certiorari, prohibition and such other remedial writs as may be provided by law and may exercise such other and further jurisdiction as may be conferred by statute. Each of the Justices or Judges shall have power to issue writs of habeas corpus to any part of the State upon petition by or on behalf of any person held in actual custody and make such writs returnable before himself, or before the Supreme Court, other Appellate Courts, or before any District Court, or judge thereof in the State. The appellate and the original jurisdiction of the Supreme Court and all other appellate courts shall be invoked in the manner provided by law.

Okla. Const. art. VII, § 4.

determine whether the Court of Criminal Appeals or the Supreme Court has jurisdiction over a controversy; (3) superintending control jurisdiction; (4) jurisdiction to issue writs of habeas corpus, mandamus, quo warranto, certiorari, prohibition, and such other remedial writs as may be provided by law; and (5) further jurisdiction conferred by statute.<sup>6</sup>

¶ 7 This Court may also exercise jurisdiction that arises independent of Article VII, § 4, and one example of this occurs when the Court answers a certified question from a federal court. In Bonner v. Oklahoma Rock Corp., we said:

This court needs no explicit grant of jurisdiction to answer certified questions from a federal court; such power comes from the United States Constitution's grant of state sovereignty. By answering a state-law question certified by a federal court, we may affect the outcome of federal litigation, *but it is the federal court who hears and decides the cause*. "Except in matters governed by the Federal Constitution or by Acts of Congress, the law to be applied in any case is the law of the state." Certification assures that federal courts are apprised of the *substantive norms of the Oklahoma legal system*.

1993 OK 131, n.3, 863 P.2d 1176, 1178, n.3 (citations omitted).

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<sup>6</sup> See 20 O.S. 2011 § 1602.

***H.B. 1970 prohibits the use of misoprostol to induce abortions, including the use of misoprostol in conjunction with mifepristone according to a protocol approved by the Food and Drug Administration and prohibits the use of methotrexate to treat ectopic pregnancies***

¶ 8 The U.S. Supreme Court certified two questions of law under the Revised Uniform Certification of Questions of Law Act, 20 O.S. 2011 §§ 1601–1611:

Whether H.B. No. 1970, Section 1, Chapter 216, O.S.L. 2011 prohibits: (1) the use of misoprostol to induce abortions, including the use of misoprostol in conjunction with mifepristone according to a protocol approved by the Food and Drug Administration; and (2) the use of methotrexate to treat ectopic pregnancies.

The certified questions are questions of statutory interpretation.<sup>7</sup> The meaning of statutory language presents a pure question of law. W.R. Allison Enters., Inc. v. Compsource Okla., 2013 OK 24, ¶ 10, 301 P.3d 407, 410. Unresolved questions of state law may be answered by this Court if certified questions are presented in accordance with the Revised Uniform Certification of Questions of Law Act, 20 O.S. 2011 §§ 1601–1611. Section 1602 outlines the discretionary power afforded this Court under the Act:

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<sup>7</sup> Curiously, although the Attorney General has not issued an opinion interpreting H.B. 1970, the Attorney General states:

[I]t should not be overlooked that this interpretation comes from the Attorney General, whose opinion “is binding upon the state officials whom it affects.” Thus, this interpretation of the law is not Petitioners’ “best guess” as to how the law will be interpreted and enforced; it is in fact how it *will* be enforced.

Petitioners’ Brief in Chief at 23, n.41 (citations omitted).

**The Supreme Court of Oklahoma “alone has the power to authoritatively determine the validity or invalidity of a statute.”** State ex rel. York v. Turpen, 1984 OK 26, ¶ 10, 681 P.2d 763, 767 (emphasis added).



The Supreme Court . . . may answer a question of law certified to it by a court of the United States . . . if the answer may be determinative of an issue in pending litigation in the certifying court and there is no controlling decision of the Supreme Court or Court of Criminal Appeals, constitutional provision, or statute of this state.

20 O.S. 2011 § 1602.

¶ 9 In 1996, a U.S. manufacturer filed a new drug application for mifepristone.<sup>8</sup> The FDA approved the application for mifepristone in 2000. According to mifepristone's FDA-approved final printed label, an informational document providing guidance about a drug's indications, precautions, and dosage, the protocol for administration of mifepristone for the termination of pregnancy requires three office visits by the patient.<sup>9</sup> During the first office visit, the patient is given 600 mg of mifepristone orally. Two days later, the patient returns to the office and is given 400 µg (0.4 mg) of misoprostol orally. Two weeks later, the patient returns to the office for a third visit to verify the procedure was successful. Mifepristone's FDA-approved label states mifepristone can be administered through forty-nine days of pregnancy.

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<sup>8</sup> "In answering a certified question, the Court does not presume facts outside those offered by the certification order. Although we will neither add nor delete facts, we may consider uncontested facts supported by the record." McClure v. ConocoPhillips Co., 2006 OK 42, n.3, 142 P.3d 390, 392, n.3. Although the record on appeal to the U.S. Supreme Court is not before this Court, the facts recited are not disputed by the parties. Additionally, neither party disputes that these facts are included in the record, and neither party has provided a citation to the record indicating evidence to the contrary exists.

<sup>9</sup> The FDA does not design or test the proposed protocol and does not conduct its own clinical trials; rather, FDA experts scrutinize submissions by the drug's sponsor, and other interested parties, concerning the safety and efficacy of the drug. See Petitioners' Brief in Chief at app. 2-3; see also Planned Parenthood v. Dewine, 696 F.3d 490, 495 (6th Cir. 2012).

¶ 10 After FDA approval of mifepristone, additional clinical trials led to the development of new protocols for administering mifepristone. The practice of providing approved medications using regimens different from that described in the medication's final printed label is known as an "off-label use," or an "evidence-based regimen." The FDA has stated that evidence-based regimens are common, permissible, and can be required by good medical practice.<sup>10</sup>

¶ 11 Evidence-based regimens for administering mifepristone vary from the protocol in mifepristone's FDA-approved label in three ways. First, the evidence-based regimens allow women to take one-third the dosage of mifepristone at the first office visit. Second, the evidence-based regimens allow a woman to self-administer the second drug, misoprostol, in the privacy of her own home rather than at a medical facility. Third, evidence-based regimens extend the effective use of mifepristone from forty-nine days to sixty-three days into the pregnancy.

¶ 12 Both the protocol in mifepristone's FDA-approved label and the evidence-based regimens require mifepristone be used *in conjunction with* misoprostol to induce an abortion. Misoprostol has not been approved by the

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<sup>10</sup> *Dewine*, 696 F.3d at 496 ("[I]t is standard medical practice in the United States for physicians to prescribe FDA-approved drugs in dosages and for medical indications that were not specifically approved—or even contemplated—by the FDA, particularly where the alternative use is supported by adequate study.").

FDA for use in abortions but has been approved by the FDA to treat ulcers.

The FDA-approved label for misoprostol is silent on abortion-related uses.

¶ 13 Although the most common evidence-based regimens involve some combination of mifepristone and misoprostol, other evidence-based regimens involve the use of methotrexate. Methotrexate is also a drug frequently used by physicians to terminate early ectopic pregnancies without surgery. Ectopic pregnancies pose grave health risks, and surgical intervention can result in serious complications, including future infertility, organ damage, and death. Methotrexate was approved by the FDA to treat neoplastic diseases, psoriasis, and rheumatoid arthritis. The FDA-approved label for methotrexate is silent on abortion-related uses.

¶ 14 In 2011, the Legislature passed H.B. 1970. Section 1, Subsection C, of H.B. 1970 provides:

C. No physician who provides RU-486 (mifepristone) or any abortion-inducing drug shall knowingly or recklessly fail to provide or prescribe the RU-486 (mifepristone) or any abortion-inducing drug according to the protocol tested and authorized by the U.S. Food and Drug Administration and as authorized in the drug label for the RU-486 (mifepristone) or any abortion-inducing drug.<sup>11</sup>

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<sup>11</sup> Title 63 O.S. Supp. 2010 § 1-729a regulated the specific drug RU-486 (mifepristone) prior to the passage of H.B. 1970. Section 1-729a was originally enacted by Senate Bill 1902, 2010 Okla. Sess. Laws 1086, and provided specific restrictions regarding the distribution and use of RU-486 (mifepristone). It required the prescribing physician to have certain qualifications and prescribe the medication under specific conditions, but it made no mention of drug labels and did not apply to other substances.

H.B. 1970 made several significant changes to § 1-729a. For example: 1) it extended the existing restrictions on RU-486 (mifepristone) to "any abortion-inducing drug" and defined that term to include "a medicine, drug, or any other substance prescribed or dispensed with the intent of terminating the clinically

To determine the meaning of H.B. 1970, we first look to the plain language of the statute. W.R. Allison, 2013 OK 24, ¶ 14, 301 P.3d at 411. “The Legislature is presumed to have expressed its intent in the text of the statute.” Id. The rules of statutory construction are employed “[o]nly where the legislative intent cannot be ascertained from the statutory language, *i.e.*, in cases of ambiguity or conflict.” McClure, 2006 OK 42, ¶ 12, 142 P.3d at 395.

¶ 15 Three times in Subsection C the phrase “RU-486 (mifepristone) or any abortion-inducing drug” is used. The Legislature’s use of the word “or” to separate the term “RU-486 (mifepristone)” from “any abortion-inducing drug” shows its intent to treat the terms as separate and distinct. In re J.L.M., 2005 OK 15, ¶ 7, 109 P.3d 336, 339 (“The Legislature’s use of the disjunctive word ‘or’ indicates its intent that either the custodial parent alone (with whom the child was living), or both parents, may be ordered to pay restitution.”); Corp. Comm’n v. Union Oil Co., 1979 OK 30, ¶ 8, 591 P.2d 711, 715 (“The use of the word ‘or’ to connect these phrases in [the statute] indicates that the grounds for relief connected thereby are disjunctive, and each is sufficient in itself to authorize the relief requested.”).<sup>12</sup>

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diagnosable pregnancy of a woman, with knowledge that the termination shall with reasonable likelihood cause the death of the unborn child”; 2) it added a definition for “drug label” to essentially reference FDA-approved guidelines for use of medications; and 3) in addition to earlier restrictions, it altered § 1-729a to require that RU-486 (mifepristone) and any “abortion-inducing drug” be provided or prescribed only “according to the protocol tested and authorized by the U.S. Food and Drug Administration and as authorized in the drug label for the RU-486 (mifepristone) or any abortion-inducing drug.”

<sup>12</sup> See also Hedrick v. Virginia, 513 S.E.2d 634, 640 (Va. 1999) (“[T]he use of the disjunctive word ‘or’ . . . signifies the availability of alternative choices.”); Resolution Trust Corp. v. United Trust Fund, Inc., 57 F.3d

¶ 16 Therefore, under H.B. 1970 if a physician wishes to provide or prescribe RU-486 (mifepristone), the physician must provide or prescribe RU-486 (mifepristone) according to the FDA-approved label for *RU-486 (mifepristone)*. If a physician wishes to provide or prescribe any abortion-inducing drug, the physician must provide or prescribe the abortion-inducing drug according to the FDA-approved label for *that abortion-inducing drug*.

¶ 17 Abortion-inducing drug is defined in Section 1, Subsection A, of H.B. 1970 as:

a medicine, drug, or any other substance prescribed or dispensed with the intent of terminating the clinically diagnosable pregnancy of a woman, with knowledge that the termination shall with reasonable likelihood cause the death of the unborn child. This includes off-label use of drugs known to have abortion-inducing properties, which are prescribed specifically with the intent of causing an abortion, such as misoprostol (Cytotec), and methotrexate. This definition does not apply to drugs that may be known to cause an abortion, but which are prescribed for other medical indications, such as chemotherapeutic agents or diagnostic drugs;

Misoprostol, when used in either the protocol described in the FDA-approved label for mifepristone or an evidence-based regimen, is an abortion-inducing drug as defined by subsection A because it is prescribed or dispensed with the intent of terminating the clinically diagnosable pregnancy of a woman, with

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1025, 1033 (11th Cir. 1995) (“[T]he disjunctive ‘or’ gives independent meaning to the words it separates.”); Knutzen v. Eben Ezer Lutheran Housing Ctr., 815 F.2d 1343, 1349 (10th Cir. 1987) (“[T]he use of a disjunctive in a statute and regulations indicates that alternatives were intended.”); Azure v. Morton, 514 F.2d 897, 900 (9th Cir. 1975) (“As a general rule, the use of a disjunctive in a statute indicates alternatives and requires that they be treated separately.”).

knowledge that the termination shall with reasonable likelihood cause the death of the unborn child. Similarly, methotrexate, when used either in an evidence-based regimen or to treat ectopic pregnancies, is an abortion-inducing drug as defined by subsection A because it too is prescribed or dispensed with the intent of terminating the clinically diagnosable pregnancy of a woman, with knowledge that the termination shall with reasonable likelihood cause the death of the unborn child.

¶ 18 The Attorney General argues that 63 O.S. 2011 § 1-730(A)(1) of the Public Health Code defines the term “abortion” to exclude the termination of ectopic pregnancies, so methotrexate can still be used off-label to treat ectopic pregnancies.<sup>13</sup> *But the operative term in H.B. 1970 is not the term “abortion,” but rather the new, separately defined term “abortion-inducing drug.”* The Legislature could have defined abortion-inducing drug to mean a medicine prescribed with the intent of causing an abortion. It did not. Instead, it defined it as a drug prescribed or dispensed with the intent of terminating the clinically diagnosable pregnancy of a woman, with knowledge that the termination shall with reasonable likelihood cause the death of the unborn

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<sup>13</sup> Section 1-730(A)(1) provides:

“Abortion” means the use or prescription of any instrument, medicine, drug, or any other substance or device intentionally to terminate the pregnancy of a female known to be pregnant with an intention other than to increase the probability of a live birth, to preserve the life or health of the child after live birth, to remove an ectopic pregnancy, or to remove a dead unborn child who died as the result of a spontaneous miscarriage, accidental trauma, or a criminal assault on the pregnant female or her unborn child.

63 O.S. 2011 § 1-730(A)(1).

child. The fact that the Legislature excludes ectopic pregnancies from the definition of abortion in § 1-730(A)(1), yet defines “abortion-inducing drug” without incorporating § 1-730(A)(1) or including similarly exclusionary language indicates the Legislature intended to ban the off-label use of methotrexate, including its use in the treatment of ectopic pregnancies.

¶ 19 The Attorney General states that “[w]hile the most common off-label protocols involve some combination of [mifepristone] and misoprostol, *other off-label protocols involve the use of methotrexate followed by misoprostol, and others yet involve the use of just misoprostol or just methotrexate.*” Petitioners’ Brief in Chief at 9, n.18 (emphasis added). The Legislature specifically referenced both misoprostol and methotrexate in the definition of an abortion-inducing drug: “This includes off-label use of drugs known to have abortion-inducing properties, which are prescribed specifically with the intent of causing an abortion, such as misoprostol (Cytotec), and methotrexate.” We find that both misoprostol and methotrexate are abortion-inducing drugs as the term is used in Subsection A; therefore, under the plain language of Subsection C of the statute, the off-label use of both misoprostol and methotrexate is prohibited.<sup>14</sup>

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<sup>14</sup> We find no merit to the Attorney General’s argument that an ectopic pregnancy is not a “true ‘pregnancy,’” so methotrexate can still be used off-label to treat ectopic pregnancies. Petitioners’ Brief in Chief at 22. Title 63 O.S. 2011 § 1-730(A)(4) defines an “unborn child” as the “unborn offspring of human beings from the moment of conception, through pregnancy, and until live birth including the human conceptus, zygote, morula, blastocyst, embryo and fetus.” And 63 O.S. 2011 § 1-730(A)(7) defines “conception” as “fertilization of the ovum of a female individual by the sperm of a male individual.” Further

¶ 20 FDA-approved labeling is “not intended to limit or interfere with the practice of medicine nor to preclude physicians from using their best judgment in the interest of the patient.”<sup>15</sup> In an often-cited bulletin specifically addressing the use of approved drugs for unlabeled indications, the FDA stated:

The FD&C Act does not, however, limit the manner in which a physician may use an approved drug. Once a product has been approved for marketing, a physician may prescribe it for uses or in treatment regimens or patient populations that are not included in approved labeling. Such “unapproved” or, more precisely, “unlabeled” uses may be appropriate and rational in certain circumstances, and may, in fact, reflect approaches to drug therapy that have been extensively reported in medical literature.

The term “unapproved uses” is, to some extent, misleading. It includes a variety of situations ranging from unstudied to thoroughly investigated drug uses. Valid new uses for drugs already on the market are often first discovered through serendipitous observations and therapeutic innovations, subsequently confirmed by well-planned and executed clinical investigation.

FDA Drug Bulletin 12:4-5, 1982.<sup>16</sup>

¶ 21 As Respondents correctly point out, and as the FDA recognizes, human progress is not static: medical research and advances do not stop upon a particular drug’s approval by the FDA. Researchers continue to perform

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discrediting this argument is the fact that the Legislature believed an ectopic pregnancy was a pregnancy having excluded the termination of ectopic pregnancies from the definition of “abortion” in 63 O.S. 2011 § 1-730(A)(1).

<sup>15</sup> Weaver v. Reagan, 886 F.2d 194, 198 (8th Cir. 1989).

<sup>16</sup> See also 59 Fed.Reg. 59,820, 59,821 (Nov. 18, 1994).



clinical trials, doctors continue to gain experience, and widespread use of a particular treatment allows the medical community to collect data about side effects, alternative doses, and potential new uses for treatments. Ninety-six percent of medication abortions in the United States are now provided according to a regimen different from the one described in mifepristone's FDA-approved label.<sup>17</sup> At the clinic operated by Respondent Reproductive Services, an evidence-based regimen for administering mifepristone is the most prevalent method for terminating early pregnancies, accounting for two-thirds of all abortions performed by the clinic, and the physicians at Reproductive Services have concluded that the protocol in the mifepristone FDA-approved label likely no longer meets the standard of care.<sup>18</sup> Both the American College of Obstetricians and Gynecologists and the World Health Organization have endorsed these alternate regimens as safer and more effective than the now-outdated regimen provided for in mifepristone's FDA-approved label.<sup>19</sup> "Good medical practice and the best interests of the patient

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<sup>17</sup> Respondents' Answer Brief at 7 (citing R. on Appeal, Tab 14, App. 4, ¶¶ 21–24). Neither side in this cause disputes that when the FDA originally approved mifepristone, it did so under a regulatory provision known as Subpart H, which allows the FDA to restrict distribution of an approved drug by its sponsor to ensure safe use. See 21 C.F.R. § 314.520. Although the FDA required mifepristone's sponsor to distribute the drug only under conditions where it would be provided by or under the supervision of a physician who was able to meet certain criteria, the FDA did not go so far as to require that administering physicians utilize mifepristone according only to the protocol described in the FDA-approved label.

<sup>18</sup> Respondents' Answer Brief at 8 (citing R. on Appeal, Tab 14, App. 7, ¶¶ 9, 14–15, 21).

<sup>19</sup> Respondents' Answer Brief at 7 (citing R. on Appeal, Tab 14, App. 4, Ex. B at 2).

require that physicians use legally available drugs, biologics and devices according to their best knowledge and judgment.”<sup>20</sup>

¶ 22 In other areas of the law, the Oklahoma Legislature has recognized the importance of allowing physicians to prescribe medications based on science and their medical judgment rather than dogmatic adherence to FDA labeling. Title 59 O.S. 2011 § 509(16) provides that unprofessional conduct for physicians includes, among other criteria:

Prescribing, dispensing or administering of controlled substances or narcotic drugs in excess of the amount considered **good medical practice**, or prescribing, dispensing or administering controlled substances or narcotic drugs without medical need in accordance with published standards.

59 O.S. 2011 § 509(16) (emphasis added).

While § 509(16) requires physicians only dispense certain drugs in amounts considered good medical practice, nowhere does it globally require physicians to dispense those drugs in accordance with their FDA-approved labels.

¶ 23 Title 63 O.S. 2011 § 1-2604 prevents health insurers from denying coverage for prescription drugs for cancer treatment merely because their use in the treatment of cancer or study of oncology is off-label. It provides:

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<sup>20</sup> United States Food and Drug Administration, Regulatory Information: “Off-Label” and Investigational Use Of Marketed Drugs, Biologics, and Medical Devices - Information Sheet, available at <http://www.fda.gov/RegulatoryInformation/Guidances/ucm126486.htm>.

**No individual policy of accident and health insurance issued which provides coverage for prescription drugs, nor any group blanket policy of accident and health insurance issued which provides coverage for prescription drugs shall exclude coverage of prescription drugs for cancer treatment or the study of oncology because the off-label use of such prescription drug has not been approved by the Federal Food and Drug Administration for that indication in one of the standard reference compendia, as defined in paragraph (d) of Section 1-1401 of Title 63 of the Oklahoma Statutes.**

Any coverage of a prescription drug required by this section shall also include provisions for coverage of **medically necessary** services associated with the administration of the prescription drug. . . .

63 O.S. 2011 § 1-2604 (emphasis added).

¶ 24 Title 63 O.S. 2011 §§ 5030.1–5030.5 provide authorization and guidelines for the Medicaid Drug Utilization Review Board. The board is charged to:

develop and recommend to the Oklahoma Health Care Authority Board a retrospective and prospective drug utilization review program for medical outpatient drugs to **ensure that prescriptions are appropriate, medically necessary, and not likely to result in adverse medical outcomes.**

63 O.S. 2011 § 5030.4(1) (emphasis added).

Nowhere in §§ 5030.1–5030.5, however, is the board constrained by uses authorized in the FDA-approved labels for prescription drugs in making its determinations. Instead, the statute uses the term “medically necessary” in deference to the knowledge and experience of physicians exercised in the practice of medicine.

¶ 25 In contrast to the deference physicians receive regarding treatment decisions in almost all other areas of medicine, H.B. 1970 requires a physician to provide or prescribe mifepristone, misoprostol, and methotrexate according only to their respective FDA-approved drug labels.<sup>21</sup> It is undisputed that the FDA-approved label for mifepristone requires a dosage level no longer considered medically necessary. It is also undisputed that misoprostol has not been FDA-approved for abortion-related uses, and methotrexate has not been approved for either abortion-related uses or for treating ectopic pregnancies. The use of misoprostol in the protocol described in the mifepristone FDA-approved label is an off-label use prohibited by the terms of H.B. 1970, and the use of methotrexate in treating ectopic pregnancies is an off-label use also prohibited by H.B. 1970. **H.B. 1970 effectively bans all medication abortions.**

### ***Conclusion***

¶ 26 The role of the physician is to heal the sick and the injured, and physicians are required to undergo rigorous training to develop the required knowledge and experience to perform that role well. Physicians must inform their patient of the risks involved in any treatment, and together with the patient, must determine the best course of treatment. Part of the Hippocratic

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<sup>21</sup> Abortion is the only area of medicine where it appears the Oklahoma Legislature has seen fit to restrict a physician's use of certain practices. See also 63 O.S. 2011 § 1-745.3; 63 O.S. 2011 § 1-745.5; 63 O.S. 2011 § 1-745.5(A).

Oath requires Physicians to “follow that system of regimen which, according to my ability and judgment, I consider for the benefit of my patients, and abstain from whatever is deleterious and mischievous.”<sup>22</sup>

¶ 27 When the district court originally found H.B. 1970 unconstitutional, it correctly concluded that:

[t]he Act’s restriction of the use of the drug RU-486 or “any other abortion inducing drug, medicine or other substance” in the manner and to the regimen set forth in the medication FPL when used for abortion is **so completely at odds with the standard that governs the practice of medicine** that it can serve no purpose other than to prevent women from obtaining abortions and to punish and discriminate against those who do.

Okla. Coal. for Repro. Justice v. Cline, No. CV-2011-1722, slip op., ¶ 7 (Dist. Ct. Okla. Cnty. May 11, 2012) (emphasis added). The plain language of the statute and the manner in which H.B. 1970 restricts the long-respected medical discretion of physicians in the specific context of abortion compels an affirmative answer to both of the questions asked, a position entirely consistent with our decision to affirm the ruling of the district court: **H.B. 1970 prohibits the use of misoprostol to induce abortions, including the use of misoprostol in conjunction with mifepristone according to a protocol approved by the Food and Drug Administration and prohibits the use of methotrexate to treat ectopic pregnancies.**

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<sup>22</sup> Grant H. Morris, Dissing Disclosure: Just What the Doctor Ordered, 44 Ariz. L. Rev. 313 (2002) (quoting 20 Encyclopedia Americana 217 (int’l ed., deluxe libr. ed. 1993)).

## **CERTIFIED QUESTIONS ANSWERED**

¶ 28 REIF, V.C.J., KAUGER, WINCHESTER, EDMONDSON, TAYLOR,  
COMBS and GURICH, JJ., concur.

¶ 29 COLBERT, C.J. and WATT, J., not voting.

2013 OK 91  
IN THE SUPREME COURT OF THE STATE OF OKLAHOMA

DAVID F. WIDNER, INDIVIDUALLY AND AS  
TRUSTEE OF THE ROBERT HAROLD  
WIDNER TRUST, AND NORMA JEAN  
WIDNER CLEMENTS, INDIVIDUALLY AND  
AS TRUSTEE OF THE ROBERT HAROLD  
WIDNER TRUST,

Plaintiffs/Appellees,

v.

ENERLEX, INC.,

Defendant/Appellant.

**FILED**  
SUPREME COURT  
STATE OF OKLAHOMA

OCT 29 2013

MICHAEL S. RICHIE  
CLERK OF  
THE APPELLATE COURTS

No. 109,787

FOR OFFICIAL  
PUBLICATION

**ON WRIT OF CERTIORARI TO THE  
COURT OF CIVIL APPEALS, DIVISION III**

¶10 Defendant/appellant offered to purchase plaintiffs'/appellees' mineral interests in Craig County, Pottawatomie County, and Seminole County. At the time, plaintiffs/appellees did not know that the Seminole County mineral interests were included in a pooling order or that proceeds had accrued under the pooling order. Defendant/appellant admitted it knew about the pooling order and the accrued proceeds but did not disclose these facts in making the purchase offer. Plaintiffs/appellees signed the mineral deeds which defendant/appellant provided and subsequently discovered the pooling order, the production, and the accrued proceeds. Plaintiffs/appellees sued defendant/appellant in Seminole County for rescission and damages, alleging misrepresentation, deceit and fraud. The Honorable Timothy Olsen, District Judge, presiding, entered summary judgment in favor of plaintiffs/appellees. The Court of Civil Appeals reversed the summary judgment. We previously granted certiorari review.

**OPINION OF THE COURT OF CIVIL APPEALS VACATED;  
SUMMARY JUDGMENT OF THE TRIAL COURT AFFIRMED.**

John L. Randolph, Jr., Tulsa, Oklahoma, for defendant/appellant.

Matthew H. McBee, Poteau, Oklahoma, for plaintiffs/appellees.

**TAYLOR, J.**

¶1 David F. Widner and Norma Jean Widner Clements (plaintiffs) sued Enerlex, Inc. (defendant) seeking rescission of mineral deeds and tort damages. The issues presented are 1) whether, under the facts and circumstances in this case, the defendant owed the plaintiffs a duty to disclose the pooling order, the production, and the accrued mineral proceeds when it made an unsolicited offer to purchase their mineral interests in Seminole County and provided the mineral deeds to be executed, and if so, 2) whether rescission is an appropriate remedy. We recently addressed similar issues in *Croslin v. Enerlex, Inc.*, 2013 OK 34, 308 P.3d 1041. *Croslin* held that Enerlex owed a duty to Croslin to disclose the production and the accrued mineral proceeds when it offered to purchase his mineral interest and provided the mineral deed conveying the mineral interest and assigning accrued mineral proceeds, if any, and that rescission was an appropriate remedy for breach of the disclosure duty.

¶2 Plaintiffs initiated this suit in the district court on the same day the *Croslin* case was filed. This case and the *Croslin* case proceeded in the district court as companion cases, and by order, this appeal was made a companion with the *Croslin* appeal. As in *Croslin*, plaintiffs contended that the undisputed facts clearly showed the defendant misrepresented material facts, the misrepresentation gave rise to a legal obligation to disclose all material facts about the mineral interests, and defendant's failure to disclose all known material facts constituted constructive fraud.



Here, as in *Croslin*, the question as to the legal effect of the undisputed facts challenges plaintiffs' entitlement to summary judgment as a matter of law on the constructive fraud claim. We review questions of law *de novo*. *Id.* at ¶9, 308 P.3d at 1045. The trial court's grant of plaintiffs' claim for rescission and cancellation of the mineral deed is governed by principles of equity, and equity will cancel a deed where it is clear that an alleged false representation deceived the complainant and caused injury. *Id.* ¶10, 308 P.3d at 1045.

¶3 The following facts are undisputed. Robert Harold Widner (Widner) owned mineral interests in Seminole County, Oklahoma, at the time of his death in 2002. Plaintiffs are the sole heirs of Widner and the beneficiaries of the Robert Harold Widner Trust. Widner's Seminole County mineral interests were included in an Oklahoma Corporation Commission pooling order in Cause CD No. 200403170-T, Order No. 491924, dated July 6, 2004. By 2008, mineral proceeds in the amount of \$34,413.94 had accrued from production in Seminole County under the pooling order. The accrued funds were reported and transmitted to the State of Oklahoma for Widner's benefit pursuant to the statutory custodial taking of proceeds from pooled mineral interests owned by unknown or unlocated persons. See 52 O.S.2001, §§ 551, *et seq.*

¶4 Defendant is in the business of buying mineral interests. In 2008, defendant made unsolicited offers to the plaintiffs to buy their Craig County, Pottawatomie County, and Seminole County mineral interests. When it made the

offers, defendant knew, but did not disclose to plaintiffs, that the Seminole County mineral interests were included in an Oklahoma Corporation Commission pooling order; that the pooling order listed Widner and the Widner trust as parties; that the pooling order allowed Widner a \$60.00 per acre bonus and a 1/8th royalty; that there was production under the pooling order; and, that \$34,413.94 had been reported and transmitted for Widner's benefit to the Oklahoma Corporation Commission pursuant to the pooling order and 52 O.S.2001, § 552.<sup>1</sup>

¶5 Unaware of the pooling order, production, and accrued mineral proceeds, plaintiff David F. Widner sold his mineral interests to defendant for \$3,800.00 plus a \$250.00 signing bonus and, on November 5, 2008, executed the deed prepared by defendant. Also unaware of the pooling order, production, and accrued mineral proceeds, plaintiff Norma Jean Widner Clements sold her mineral interests to defendant for \$3,000.00 plus a \$250.00 signing bonus and, on November 13, 2008, executed the deed prepared by defendant. It is undisputed that plaintiffs would not have executed the mineral deeds had they known about the Seminole County production or accrued mineral proceeds.

¶6 The mineral deeds prepared by defendant in this case mirror those in *Croslin*. In each of the mineral deeds, the granting clause conveyed, transferred,

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<sup>1</sup> Pursuant to 52 O.S.2001, § 552, the undistributed mineral interest proceeds were reported and transmitted to the Oklahoma Corporation Commission, deposited in the Mineral Owner's Escrow Fund, and then, pursuant to 52 O.S.Supp.2003, § 554, the funds were transferred to the Mineral Owner's Fund in the State Treasury.

assigned, and delivered the "interest in and to all of the Oil, Gas, and any other classification of valuable substance . . . in and under and **that may be produced** from the following described lands. . . ." (Bold added.) The language that followed the granting clause explained its meaning:

it being understood and agreed that this transfer and assignment covers and **includes** that the grantee shall have, receive, and enjoy the herein granted undivided interests in and to **all royalties, accruals and other benefits, if any, from all Oil and Gas heretofore or hereafter run**, whether they be held therefore by any purchaser or other legal entity, or hereafter produced, sold and paid to the Grantee. The Grantor hereby irrevocably appoints and constitutes the Grantee as agent and attorney-in-fact for the limited purpose only of executing division and transfer orders and all other instruments necessary to make fully effective this assignment and conveyance so that the Grantee may act in Grantor's place and stead for such purpose.

(Bold added.)

¶7 By letter dated October 6, 2009, defendant advised plaintiff Norma Jean Widner Clements that it believed the State of Oklahoma was holding more than \$10,000.00 attributable to her former mineral interests and requested that she sign an agreement waiving her right to the mineral proceeds. The letter offered, that in return, "Enerlex will remit to you 25% of the funds received from the State of Oklahoma that accrued prior to your transfer to Enerlex. . . ." <sup>2</sup> Plaintiff did not

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<sup>2</sup> Under 60 O.S.2011, § 674.1, Enerlex was entitled to retain, at the most, 25 %, but not 75% as offered, of the funds held for Widner. Originally enacted in 1990 (1990 Okla.Sess.Laws, ch. 301, § 1), the language remains the same. Section 674.1 reads:

No person who:

1. informs a potential claimant of any unclaimed funds or other property, tangible or intangible, held pursuant to the Uniform Disposition of Unclaimed Property Act that such claimant may be entitled to claim such unclaimed property; or

execute defendant's proposed agreement. Instead, plaintiffs filed suit against defendant on October 23, 2009.

¶8 Plaintiffs' petition alleged that defendant, when it offered to purchase their mineral interests, had a duty to inform them of the pooling order, the production, and the accrued mineral proceeds; that defendant's failure to inform them constituted constructive fraud; and that defendant's deceitful and fraudulent actions amount to fraud and justify rescission, consequential damages, actual damages, and punitive damages. Defendant answered and counter-claimed that it was the rightful owner of the mineral interests and that plaintiffs filed suit in bad faith and slandered title of the mineral interests.

¶9 Plaintiffs moved for summary judgment. The trial court granted partial summary judgment in favor of plaintiffs on the false representation claim and ordered an accounting of royalty proceeds paid to defendant, leaving other issues unresolved. The parties agreed to the accounting and waived all other claims and issues for purposes of summary judgment, and the trial court entered the summary judgment order on appeal. Similar to its summary judgment in *Coslin*, the trial court

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2. files a claim for any funds or other property, tangible or intangible, on behalf of a claimant of such funds or property, shall contract for or receive from the claimant, for services, an amount that exceeds twenty-five percent (25%) of the value of the funds or property recovered.

If the funds or property involved are mineral proceeds, the amount for services shall not include a portion of the underlying minerals or any production payment, overriding royalty, or similar payment.

B. The provisions of this section shall apply to contracts executed on or after July 1, 1990.

concluded that: 1) a mineral interest purchaser has the duty to disclose production and failure to do so is a false representation, citing *Deardorf v. Rosenbusch*, 1949 OK 117, 206 P.2d 996; 2) a seller's constructive knowledge of production is not a defense to fraudulent misrepresentation, citing *Uptegraft v. Dome Petroleum Corp.*, 1988 OK 129, 764 P.2d 1350; and 3) the unclaimed property statutes and regulations place additional notice requirements upon one who claims funds in the Mineral Owners Escrow Fund based upon the transfer of a mineral interest. The trial court granted plaintiffs' claim for rescission and cancelled the mineral deeds, declared plaintiffs to be the rightful owners of the mineral interests, and directed plaintiffs to return the purchase money defendant paid them less any royalty proceeds that may have been paid to defendant.

¶10 Defendant appealed. The Court of Civil Appeals reversed the summary judgment, finding that defendant made no factual inducement, representation or misrepresentation that gave rise to a duty to disclose the pooled mineral interests or production. The Court of Civil Appeals also determined defendant had no duty to disclose the pooled mineral interests and accrued mineral proceeds to the plaintiffs under the unclaimed property statutes or the pooled mineral interests statutes. We previously granted plaintiffs' petition for certiorari review.

¶11 Following the teachings of *Berry v. Stevens*, *Deardorf v. Rosenbusch*, and *Uptegraft v. Dome Petroleum Corp.*, *Croslin* found that knowledge of the accrued mineral proceeds undoubtedly motivated Enerlex to extend unsolicited

offers to purchase the mineral interest and that the "if any" language in the mineral deed prepared by Enerlex, indirectly if not directly, created a false impression that Enerlex did not know of any production or any accruals from all oil and gas heretofore run. *Croslin*, 2013 OK 34, ¶30, 308 P.3d at 1051. Upon review of the summary judgment record on appeal, we find *Croslin* governs the issues presented here.

¶12 As in *Croslin*, the language in defendant's mineral deed assigning the accruals of royalties, if any, from heretofore runs gave rise to a duty on the part of defendant to disclose the whole truth, including all material facts about the accrual of the mineral proceeds. Instead of disclosing the \$34,413.94 of accrued mineral proceeds to the plaintiffs, defendant remained silent and allowed plaintiffs to rely, to their detriment, on the false impression created by the "if any" language in the mineral deeds. As in *Croslin*, plaintiffs were entitled to summary judgment on the legal issue of defendant's disclosure duty as a matter of law. *Id.* at ¶32, 308 P.3d at 1051.

¶13 Similar to its goal in *Croslin*, defendant, in this case, wanted to spend a total of \$7,300.00 in cash and get \$34,413.94 in cash plus mineral interests in Craig County, Pottawatomie County, and Seminole County and future income. To accomplish its goal, defendant offered to purchase the mineral interests from plaintiffs for a total of \$7,300.00, and relying on plaintiffs' ignorance of the \$34,413.94 of accrued mineral proceeds, defendant provided plaintiffs mineral deeds

transferring both the mineral interests and the accrued mineral proceeds. Defendant obtained the mineral deeds from plaintiffs by false representation and suppression of the whole truth. Defendant is liable to plaintiffs for constructive fraud. Fraud in the procurement of a written instrument vitiates it in the hands of one seeking its benefit. *Id.* at ¶37, 308 P.3d at 1052. Rescission is an appropriate remedy for defendant's misrepresentation and constructive fraud.

**OPINION OF THE COURT OF CIVIL APPEALS VACATED;  
SUMMARY JUDGMENT OF THE TRIAL COURT AFFIRMED.**

ALL JUSTICES CONCUR.

2013 OK 89  
IN THE SUPREME COURT OF THE STATE OF OKLAHOMA

FILED  
SUPREME COURT  
STATE OF OKLAHOMA

IN REGARD TO THE APPLICATION  
OF KEITH ALLEN OSBURN,

OCT 29 2013

MICHAEL S. RICHIE  
CLERK OF  
THE APPELLATE COURTS

Appellee,

v.

Case No. 109,867

OKLAHOMA DEPARTMENT  
OF CORRECTIONS,

FOR OFFICIAL  
PUBLICATION

Appellant.

ON APPEAL FROM THE DISTRICT COURT OF  
GARFIELD COUNTY  
HONORABLE PAUL K. WOODWARD  
DISTRICT JUDGE

¶0 Appellee pled no contest to indecent exposure in 1998. Four months later, the Sex Offenders Registration Act (SORA), 57 O.S., § 581 et seq. was amended to include indecent exposure as an enumerated offense requiring registration. Appellant thereafter required Appellee to register as a sex offender. Appellee challenged his registration based on the fact the offense of indecent exposure was not an enumerated offense at the time of his conviction. The trial court agreed with Appellee and ruled he is not required to register. We affirm.

THE TRIAL COURT'S AUGUST 15, 2011, ORDER  
IS HEREBY AFFIRMED

Michael Thomas Oakley, Oklahoma Department of Corrections, Oklahoma City,  
Oklahoma, for Appellant

Grace K. Yates and Shawna N. Taylor, Holmes and Yates, Ponca City, OK,  
for Appellee

COMBS, J.:



## FACTS AND PROCEDURAL HISTORY

¶1 On or about, August 13, 1997, Appellee, Keith Allen Osburn (hereinafter, “Osburn”) was charged with one felony count of indecent exposure in violation of 21 O.S., § 1021 (A) (1) (CF 97-262; Kay County, Oklahoma). On June 22, 1998, Osburn pled no contest to the charge and was given a five-year sentence of which he was to serve ten weekends in the Kay County Jail with the remainder of his sentence suspended. Thereafter, in 2000, Osburn began registration under the Sex Offenders Registration Act (hereinafter “SORA”), 57 O.S., § 581 et seq.<sup>1</sup> Sometime after November 1, 2007, the Appellant, the Oklahoma Department of Corrections (hereinafter, “Department”), assigned Osburn a sex offender risk level of one.<sup>2</sup> This increased Osburn’s allegedly required SORA registration period to fifteen years from the date of the completion of his sentence. The Department determined his registration period will now end in June 2018.<sup>3</sup>

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<sup>1</sup> The record does not reflect why there was a delay before registration began. Under the Appellant’s viewpoint, Osburn should have been required to register when the law was amended to include indecent exposure as an enumerated offense which occurred on November 1, 1998, approximately four months after Osburn was convicted (HB 3144, 1998 Okla. Sess. Laws c. 347, §1).

<sup>2</sup> At the time of Osburn’s June 22, 1998, conviction, persons **who were required to register** under SORA only had to register for ten years following release from incarceration (Title 57 O.S. Supp. 1997, §§ 582 and 583 (C)). The ten-year registration period was extended on April 26, 2004, by requiring it to begin from the date of completion of the sentence (SB 1191, 2004 Okla. Sess. Laws c. 162, §1). Effective November 1, 2007, a sex offender risk level system was created which again increased the period of registration (HB 1760, 2007 Okla. Sess. Laws c. 261). It included three levels of risk to be assigned to sex offenders by the Oklahoma Department of Corrections (57 O.S. Supp. 2007, § 583 (C) and (D)). For a risk level of one, a person must register for fifteen years from the date of the completion of the sentence.

<sup>3</sup> The Department calculates this period by adding fifteen years to the date when his original five-year sentence was

¶2 On June 21, 2002, Osburn pled guilty to felony charges in two separate criminal cases for failure to register as a sex offender in violation of 57 O.S., § 587 (CF 2001-426 and CF 2002-226; Kay County, Oklahoma). In each case, the district court sentenced him to three years in the state penitentiary. The district court ordered each sentence suspended and the two sentences to be served consecutively. Neither case was appealed. On May 29, 2009, another Kay County case charging Osburn with failure to register was dismissed by the district court (CF 07-520).

¶3 On March 29, 2011, Osburn filed an Application to terminate his SORA registration in Garfield County, Oklahoma, the county of his residence. In his Application, he asserted he was eligible to apply under 57 O.S., § 583 (E) to end his SORA registration because he had been registered for ten years. At the time he filed his application, subsection E provided as follows:

E. Any person assigned a numeric risk level of one who has been registered for a period of ten (10) years and **who has not been arrested or convicted for any felony or misdemeanor offense** since being released from confinement, may petition the district court in the jurisdiction **where the person resides** for the purpose of removing the numeric risk level designation and allowing the person to no longer be subject to the registration requirements of the Sex Offenders Registration Act. (Emphasis added).

Title 57 O.S. Supp. 2010, § 583 (E).

¶4 On May 19, 2011, Osburn filed an Amended Application wherein he asserted the SORA registration requirements were “inaccurately” applied to him because he should never have been required to register in the first place. Osburn indicates this argument is based upon the constitutional prohibition against ex post facto laws.<sup>4</sup> When he was convicted on June 22, 1998, 57 O.S. Supp. 1997, § 582 specifically excluded indecent exposure from SORA registration.<sup>5</sup> Four months after his conviction, amendments to Section 582 became effective which included indecent exposure as an enumerated offense. This was accomplished by deleting the language which specifically excluded that crime from SORA registration.<sup>6</sup>

¶5 Osburn further asserted even if he was required to register he has already completed his ten years of registration. He argued he still can apply to be removed from SORA registration under Section 583 (E) because even though he was arrested and convicted after his indecent exposure conviction, those arrests and convictions were for failure to register under SORA and it is clear that at the time

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<sup>4</sup> Okla. Const. art. 2, § 15.

<sup>5</sup> Title 57 O.S. Supp. 1997, § 582; 1997 Okla. Sess. Laws c. 260, § 3 (eff. Nov. 1, 1997) which was in effect when he was convicted on June 22, 1998, of indecent exposure provided an exception from SORA registration for certain offenses which specifically included 21 O.S. § 1021 (A) (1) (indecent exposure). It stated:

“The provisions of the Sex Offenders Registration Act . . . shall apply to any person who, after November 1, 1989, has been convicted . . . for a crime or an attempt to commit a crime provided for in . . . **1021, except for a crime provided for in paragraph 1 of subsection A of Section 1021 . . .**” (Emphasis added).

<sup>6</sup> Title 57 O.S. Supp. 1998, § 582; 1998 Okla. Sess. Laws c. 347, §1 (eff. Nov. 1, 1998). The amendments struck through “~~except for a crime provided for in paragraph 1 of subsection A of Section 1021~~” thereby, deleting an exception to registration for crimes involving indecent exposure.

of his indecent exposure conviction the statute did not require him to register.

¶6 The Department filed an Objection to Amended Application alleging the trial court was without jurisdiction to hear Osburn's application. Title 57 O.S. Supp. 2010, § 583 (E) does not allow a person who has been "arrested or convicted for any felony or misdemeanor offense" to apply for a reduction to their level one registration period. The Department asserts Osburn had been arrested three times and convicted twice since his 1998 indecent exposure conviction. The Department further asserted Osburn may not collaterally attack the validity of his convictions for failure to register. The Department argued, even if his convictions could be challenged his arrests cannot. The language is clear that even an arrest will thwart eligibility for reduction under the statute. Therefore, it was asserted the trial court did not have jurisdiction to reduce Osburn's registration period.

¶7 On August 15, 2011, the trial court found SORA did not apply to Osburn when he was convicted in June 1998 and therefore he should never have been required to register. The court ordered Osburn was not subject to SORA's registration requirements. The Department appealed.

### STANDARD OF REVIEW

¶8 On appeal, this Court assumes "plenary independent and non-deferential authority to reexamine a trial court's legal rulings." *Kluver v. Weatherford Hospital Auth.*, 1993 OK 85, ¶14, 859 P.2d 1081, 1084. Issues of a statute's

constitutional validity and of its construction and application are questions of law reviewed *de novo*. *Gilbert v. Security Finance Corp. of Oklahoma, Inc.*, 2006 OK 58, ¶2, 152 P.3d 165, 171.

### ANALYSIS

¶9 On appeal, the Department alleges “the district court had neither jurisdiction over Appellant nor subject matter jurisdiction.” The Department asserts the trial court lacked jurisdiction over the Department because the court allegedly granted injunctive relief and the action was not brought in the proper venue. This assertion is based upon 12 O.S. 2011, § 133. That section of law provides in pertinent part:

Actions for the following causes must be brought in the county where the cause, or some part thereof arose: . . .

Second. An action against a public officer for an act done by him in virtue, or under color, of his office, or for neglect of his official duties.

Title 12 O.S. 2011, § 133.<sup>7</sup>

The Department also reiterates its argument that the court lacked subject matter jurisdiction because 57 O.S. Supp. 2010, § 583 (E) only allows a district court to reduce a level one registration period if the applicant had not been subsequently

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<sup>7</sup> The Department alleges Osburn’s challenge amounts to an attack on how the Department administers SORA and therefore the only proper venue, per 12 O.S. 2011, Section 133, is in Oklahoma County where the Department’s official residence is located. The Department cites three opinions of this Court; *Hillcrest Medical Center v. Lee*, 1978 OK 23, 575 P.2d 971, *Grand River Dam Authority v. State*, 1982 OK 60, 645 P.2d 1011, and *Office of Gov’r – Dept. of Indus. Development v. Dalton*, 1977 OK 34, 560 P.2d 971. In each case the governmental entity challenged venue at the trial court level based on Section 133. This Court held in those cases venue was proper where the entity resided based upon its interpretation of this section. The rationale being the decisional act of the public officer which gave rise to the cause of action emanates from the county of the official residence of the public officer. See *Dalton*, 1977 OK 34, ¶9.

arrested or convicted and Osburn had subsequent arrests and convictions.

¶10 Title 12 O.S. 2011, § 2012 (F) provides, “1. A defense of lack of jurisdiction over the person, [and] improper venue . . . **is waived . . . if omitted** from a motion that raises any of the defenses or objections which this section permits to be raised by motion . . . .” Subsection B of Section 2012 lists the types of defenses that can be made by motion which include “[l]ack of jurisdiction over the subject matter . . . [and] [f]ailure to state a claim upon which relief can be granted.” The record demonstrates the Department’s first filing at the trial court level consisted of an Objection to Amended Application wherein the Department asserted Osburn failed to state a claim upon which relief can be granted and requested the trial court to dismiss the action for want of jurisdiction. This argument challenging jurisdiction was based on the trial court’s subject matter jurisdiction and not personal jurisdiction. The Objection did not raise the defense of lack of personal jurisdiction or improper venue. Therefore, we find the Department waived any objection based upon personal jurisdiction or improper venue.

¶11 The dispositive issue here is whether an amendment to SORA which makes its provisions applicable to new offenses may be applied to persons convicted in Oklahoma of such offenses prior to the amendment. We find that it cannot be applied to such persons. Recently, this Court held SORA and its numerous amendments when viewed in their entirety have a punitive effect that outweighs

their non-punitive purpose and therefore a retroactive application of SORA's registration provisions would violate the ex post facto clause in the Oklahoma Constitution.<sup>8</sup> *Starkey v. Oklahoma Department of Corrections*, 2013 OK 43, ¶77, 350 P.3d 1004, 1030. Here we are not just dealing with an increase in the registration provisions of SORA; we are dealing with applying SORA in its entirety to someone convicted prior to the Act being applicable to his offense. For a person convicted in Oklahoma of an enumerated sex offense, the controlling provisions of SORA are those in effect upon the date of conviction. On June 22, 1998, Osburn's date of conviction, SORA did not apply to a person convicted of indecent exposure (21 O.S., § 1021 (A) (1)). Therefore, we affirm the trial court's holding that Osburn is not subject to the provisions of SORA.

**THE TRIAL COURT'S AUGUST 15, 2011, ORDER  
IS HEREBY AFFIRMED**

¶12 COLBERT, C.J., REIF, V.C.J., KAUGER, WATT, EDMONDSON, COMBS, and GURICH, JJ., concur.

¶13 WINCHESTER and TAYLOR, dissent.

¶14 WINCHESTER, J., dissenting:

I dissent for the same reasons that I stated in my dissent in *Starkey v. Oklahoma Department of Corrections*, 2013 OK 43, 305 P.3d 1004.

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<sup>8</sup> Okla. Const. art. 2, § 15.

¶15

**TAYLOR, J., dissenting:**

I dissent for the same reasons that I stated in my dissent in *Starkey v. Oklahoma Department of Corrections*, 2013 OK 43, 305 P.3d 1004.