1	IN THE UNITED STATES DISTRICT COURT
2	FOR THE WESTERN DISTRICT OF OKLAHOMA
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4	CHARLES F. WARNER, et al.,
5	Plaintiffs,
6	vs. Case No. CIV-14-665-F
7	KEVIN J. GROSS, et al.,
8	Defendants.
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12	TRANSCRIPT OF COURT'S RULING
13	BEFORE THE HONORABLE STEPHEN P. FRIOT
14	UNITED STATES DISTRICT JUDGE
15	DECEMBER 22, 2014
16	3:00 P.M.
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24	Proceedings recorded by mechanical stenography; transcript
25	produced by computer-aided transcription.

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1	(PROCEEDINGS HAD DECEMBER 22, 2014.)
2	THE COURT: Good afternoon. We're here in Civil
3	14-665, Charles Warner and others v. Kevin Gross and others,
4	for the Court's ruling on the motion of four plaintiffs for
5	preliminary injunction. Counsel will please give your
6	appearances.
7	MS. GHEZZI: Patti Ghezzi for the plaintiffs that are
8	represented by the Federal Public Defender's Office in the
9	Western District.
10	MS. HENRICKSEN: Lanita Henricksen for Andrew,
11	Warner, Hancock, Jackson, and Glossip.
12	MR. AUTRY: David Autry for James Coddington, Your
13	Honor.
14	THE COURT: We have Arizona counsel present by
15	telephone?
16	MS. KONRAD: Yes. Robin Konrad and Dale Baich for
17	Plaintiff Tremane Wood.
18	MR. HADDEN: John Hadden for state defendants, Your
19	Honor.
20	MR. STEWART: Aaron Stewart for state defendants,
21	Your Honor.
22	THE COURT: I'll now make my findings of fact and
23	conclusions of law with respect to the preliminary injunction
24	which has been motion for preliminary injunction which has
25	been filed by Plaintiffs Charles Warner, Richard Glossip, John

Grant, and Benjamin Cole.

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The hearing on these plaintiffs' motion for preliminary injunction was held on December 17, 18, and 19, 2014. The plaintiffs were ably represented by Dale A. Baich, Robin C. Konrad, Patti P. Ghezzi, and Randy A. Bauman. The defendants were ably represented by John D. Hadden, Aaron J. Stewart, and Jeb E. Joseph. Over three full days of hearings, generating 694 pages of transcript, the plaintiffs called 14 witnesses, including several Oklahoma Department of Corrections employees, and the defendants called three witnesses.

My scheduling of the hearing on the motion for preliminary injunction, as well as my scheduling of the preparatory steps leading to the hearing was driven by the fact that these four movants are scheduled for execution beginning in the case of Charles Warner on January 15, 2015. That necessitated a rather compressed schedule. Even though the schedule was compressed, there were certain essential steps that, although unfolding on a tight time schedule, certainly could not be eliminated. As an example, it was my conclusion that the plaintiffs ought to have the benefit of discovery as thorough and searching as was possible under the circumstances. And I believe that plaintiffs have indeed had the benefit of thorough discovery.

Plaintiffs' discovery began, at least in terms of substantial discovery, with the production of thousands of pages of documents that were either turned over to the

Department of Public Safety or generated by the Department of Public Safety in the investigation that was conducted by that agency following the Lockett execution. That included thousands of pages of interview transcripts as well as numerous original source documents.

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And before I go any further, I will say again that I applaud the diligence and professionalism of counsel on both sides. For the reasons I have described, among others, this has been, to put it mildly, a very demanding case, especially in the run-up to the three-day hearing last week. Although there were some instances in which I had to referee discovery disputes on fairly short notice, I can say without hesitation that preparation for the preliminary injunction hearing proceeded with less rancor than there would have been if counsel on both sides had not made every effort as true professionals to bring the matter to this stage with a hard focus on the merits and with minimal diversions unrelated to the merits.

I am making my findings of fact and conclusions of law in this setting, on the record, as permitted by Rule 52(a). As I said at the end of the day last Friday, I could take another five or six working days to turn out a polished 35- or 40-page order, but I am certain that the parties and their counsel would rather have those five or six days back so that they can prepare for the next stage of this litigation, which will

necessarily unfold between now and January 15, 2015. I will assure all concerned, however, that even though I am ruling from the bench rather than taking another five or six days to produce a formal written order, the findings and conclusions that I'm about to express are made with all of the thought process that would ultimately go into a formal order.

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Many of the matters that I'm about to address involve mixed issues of fact and law. For that reason, it is in some ways a bit artificial to speak in terms of findings of fact separately from conclusions of law. I will separate the two as much as I reasonably can. However, I am confident that the parties, as well a reviewing court, will be able to discern the difference between my factual findings and my legal conclusions. In any event, of course, to the extent that I express a legal conclusion as if it were a matter of fact, it should be regarded as a legal conclusion and vice versa.

My exceedingly capable reporter is prepared to produce a transcript of my ruling in very short order. For ease of reference by the parties and by a reviewing court, the reporter has advised me that it would be permissible for me to insert headings into the transcript before the transcript is filed. I think that would be helpful to all concerned and I will do that before the transcript is filed. I assure you, however, that because of the very nature of a ruling from the bench, I will not make any change of any kind in the record of my ruling as

taken by the reporter.

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2 I'll now turn to the factual history of this matter.
3 FACTUAL HISTORY

Clayton Derrell Lockett, having been convicted of first degree murder and sentenced to death, was scheduled for execution at the Oklahoma State Penitentiary on April 29, 2014. The execution took place as scheduled and it achieved the intended result, the death of Clayton Lockett, but the execution was ineptly performed in some ways, as I will discuss later in these findings.

As a result of the extraordinary events surrounding the Clayton Lockett execution, Governor Mary Fallin issued Executive Order 2014-11 on April 30, 2014, which mandated an independent investigation of the events leading up to and during the execution of Clayton Lockett. The Governor's Executive Order appointed Michael Thompson, the Secretary of Safety and Security and the Commissioner of the Department of Public Safety to lead the independent investigation process.

The Department of Public Safety investigation culminated in the issuance of a report entitled "The Execution of Clayton D. Lockett," Case Number 14-0189SI, by the Oklahoma Department of Public Safety. On September 16, 2014, that report was filed in this action as Docket Entry Number 49-1.

The DPS report contains a wealth of information, much of it favorable to the plaintiffs. The plaintiffs have asserted

that for their purposes there are some relevant facts that are not included in the DPS report. That is correct. An encyclopedic report on the Lockett execution would probably have run to 3- or 400 pages rather than 29 pages. But the DPS report, which resulted from a thorough investigation that was conducted with noticeable professional integrity by Captain Jason Holt, under the supervision of Commissioner of Public Safety Michael Thompson, certainly provided plaintiffs with a good starting point. By the time Captain Holt left the witness stand last Thursday, no one in this courtroom could have doubted the seriousness with which he took his assignment to lead the DPS investigation.

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Meanwhile, this action was filed on June 25, 2014. As indicated by the allegations on pages 6 to 9 of the original complaint in this case, Docket Entry Number 1, the plaintiffs' original complaint in this action centered substantially on the events surrounding the execution of Clayton Lockett. The same is true of the amended complaint, Docket Entry Number 75, filed on October 31, 2014.

The amended complaint also focuses on the revised execution protocol adopted by the Oklahoma Department of Corrections, DOC Policy OP-040301, with an effective date of September 30, 2014. I will discuss that revised protocol as relevant to the issues now before the Court later in my findings.

THESE FOUR PLAINTIFFS

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Although the plaintiffs in this action include all or nearly all of the death row inmates in the state of Oklahoma, the matter now before the Court is the motion for preliminary injunction which was filed by four of those plaintiffs; namely, Charles Frederick Warner, Richard Eugene Glossip, John Marion Grant, and Benjamin Robert Cole. The four plaintiffs who have filed this motion for preliminary injunction have been scheduled for execution, respectively, on January 15, January 29, February 19, and March 5, 2015.

As recounted by the Oklahoma Court of Criminal Appeals at 144 P.3d 838, Charles Warner raped and murdered an 11-month-old baby girl on August 22, 1997, an assault which resulted in, among other injuries, two skull fractures including a depressed fracture and two fractures of the baby girl's left jaw.

As recounted by the Oklahoma Court of Criminal Appeals, 157 P.3d 143, on January 7, 1997, Richard Glossip hired Justin Sneed to kill Barry Van Treese, the employer of Glossip and Sneed, which Sneed proceeded to do by bludgeoning Van Treese to death with a baseball bat.

As recounted by the Oklahoma Court of Appeals, 95 P.3d 178, on November 13, 1988, John Grant, then an inmate at the Connor Correctional Center in Hominy, Oklahoma, murdered Gay Carter, a food service supervisor at the Connor Correctional Center, by stabbing her 16 times with a shank.

As recounted by the Oklahoma Court of Criminal Appeals, 164 P.3d 1089, on December 20, 2002, Benjamin Cole murdered his nine-month-old daughter, Brianna Cole, by snapping her spine in half and inflicting other fatal injuries because she would not stop crying.

All four of the moving plaintiffs have reached the end of their trial, direct review and collateral review process, which includes, in the case of Messrs. Warner and Glossip, two trials, two first degree murder convictions, and two sentences of death.

I will now address the facts surrounding the Lockett execution.

THE LOCKETT EXECUTION

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Although a number of preliminary steps took place earlier in the day on April 29, 2014, and even before that day, the final sequence of events leading to the execution of Clayton Lockett began at approximately 5:22 p.m. on April 29 when Lockett was placed onto the execution table and strapped down. Earlier in the day, Lockett had twice refused visits from his attorneys. He had also cut himself twice on April 29 at "the bend of the elbow," as described by Warden Trammell. That is as page 225 of the transcript.

The execution of Clayton Lockett was the first Oklahoma execution using midazolam. The protocol called for the administration of 100 milligrams of midazolam, 40 milligrams of

vecuronium bromide, and 200 milliequivalents of potassium
chloride. Midazolam had been added to the protocol
approximately two weeks before the Lockett execution after it
was determined that pentobarbital would not be available for
the Lockett execution.

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On April 29, before the blinds between the execution chamber and the reviewing rooms were raised, the execution team had worked for nearly an hour trying to establish intravenous access to Lockett's cardiovascular system. Postmortem examination revealed at least a dozen needle puncture marks on Lockett's body indicating at least that many attempts to establish IV access.

The first member of the execution team who was involved in securing intravenous access to Lockett's cardiovascular system was an emergency medical technician licensed as a paramedic. The paramedic attempted, without success, to establish IV access in the typical location in the crook of Lockett's left arm. Three attempts to establish IV access at that location were unsuccessful.

Next a physician member of the execution team attempted to establish IV access through Lockett's left jugular vein.

Although it appeared momentarily that this attempt had been successful, that success was short-lived. At the same time, the paramedic attempted to establish IV access by way of Lockett's right arm. Three attempts to do so were

unsuccessful. After those unsuccessful attempts, the physician sought unsuccessfully to establish IV access through Lockett's left subclavian vein and the paramedic attempted to establish IV access in two locations on Lockett's right foot.

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The physician next attempted to establish IV access by way of Lockett's right femoral vein. The physician and the paramedic concluded that this attempt had been successful. To facilitate right femoral IV access, the physician asked for a longer catheter so that they could attempt to establish IV access through Lockett's femoral vein.

A one-and-a-quarter-inch 14-gauge angiocatheter was used for this purpose. The one-and-a-quarter-inch needle was inserted and was "positional," meaning the patency of the IV flow was dependent upon relatively precise positioning of the catheter. This one-and-one-quarter-inch catheter was taped in place. The use of the one-and-a-quarter-inch catheter was clearly inappropriate, a failure that is made all the more inexplicable by the fact that a central line IV kit was available.

After the physician and the paramedic concluded that femoral IV access had been established, Warden Trammell covered Lockett's body with a sheet. For the purpose of preserving Lockett's privacy, his genital area was covered. This was an improvident decision. From that point until it appeared that there may be a problem with intravenous flow, Lockett's genital

area remained covered and the IV access point could not be observed.

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There was approximately a 23-minute delay in the beginning of the execution. This delay was evidently due to the difficulties in establishing acceptable IV access. The blinds were raised at approximately 6:23 p.m. After the blinds were raised, Warden Trammell read the death warrant and Lockett was asked whether he had any last words. He had none.

The execution team began to push the midazolam into the IV manifold. Administration of the midazolam was followed by administration of the vecuronium bromide and potassium chloride, both by way of the right femoral IV access point. Confirmation of continuous IV flow was, to put it mildly, hampered by the fact that the execution team put a hemostat on the IV line and then they covered the IV injection access point with a sheet with the result that, in the words of Captain Holt at page 408 of the transcript, they had "covered the one and stopped the other," which made it impossible to confirm that the IV flow continued or even that it could continue.

At approximately 6:30, the physician performed a consciousness check and determined that Lockett was conscious. At approximately 6:33, Lockett was determined to be unconscious and the vecuronium bromide was pushed, followed by a saline flush and the potassium chloride. Shortly after that, Lockett began to move. He raised his head and was heard to say, "man"

or "oh, man." By the account of Edith Shoals, which I find to be credible, Lockett said, "This shit is fucking with my mind." That is at page 204 of the transcript. Lockett was heard to mumble "something is wrong," and he moved his shoulders and head forward. By the account of Jeanetta Boyd, which was also credible, Lockett was heard to say, "The drugs aren't working." That is at page 182 of the transcript. After these movements and verbalizations, the blinds were closed at the direction of Warden Trammell.

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The physician recognized that there was a problem. The physician lifted the sheet and recognized that the IV had infiltrated, meaning that the IV fluid had leaked into the tissue surrounding the IV access point. The physician noted an area of swelling under Lockett's skin. It was smaller than a tennis ball but larger than a golf ball. It is evident from the autopsy photographs and from the testimony that a bulge of that size, unmistakably indicating a serious infiltration problem, could have and should have been noticed at a significantly earlier stage of the execution process.

Vecuronium bromide is a potent paralytic agent. The intravenous administration of a massive dose of vecuronium bromide, as was called for by the lethal injection protocol that governed Lockett's execution, would have resulted in complete paralysis. Lockett would have been unable to breathe, speak, or raise his head. During this phase of his execution,

Lockett was surely experiencing all of the mental pain that is inevitable in the execution process as well as serious physical discomfort if not serious physical pain.

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Postmortem toxicology confirmed the presence of midazolam as well as vecuronium and potassium in Lockett's femoral blood at 45 milliequivalents for the potassium and 460 nanograms per milliliter for the vecuronium. The vecuronium bromide was pushed after the midazolam but before the potassium chloride and the vecuronium obviously did not immediately have its intended paralytic effect. Not all of the potassium chloride was pushed, but the potassium chloride was behind the vecuronium bromide and the vecuronium bromide clearly did not flow into Lockett's cardiovascular system in the manner contemplated by the lethal injection protocol. Consequently, it is not possible to determine the extent to which Lockett suffered the searing pain that would result from an injection of potassium chloride into a sensate person.

Although I cannot and do not find that Lockett was not in pain during this part of the execution process, I do note that during the time that Lockett moved, vocalized, and raised his head and shoulders, of all of the verbalizations attributed to Lockett, at least two of which were complete sentences, the most specific, emphatic, and intelligible statement was, "This shit is fucking with my mind." Which may or may not be a statement that one would have expected Lockett to make if he

was feeling the searing pain that he certainly would have felt if he had been conscious while any substantial amount of sodium chloride was being delivered to his tissues.

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By all accounts, potassium chloride will cause a sensate individual to feel serious pain. It is clear that patent IV flow of potassium chloride into the cardiovascular system of Clayton Lockett, while conscious and sensate, would have had the extremely painful effect that has been described because patent IV flow would have delivered the substance throughout his body to the most sensitive receptors. What is not clear from the evidence is the extent to which that, in fact, occurred.

After the swelling in Lockett's groin was noted, the execution process was halted and the blinds were lowered at about 6:42 p.m., approximately 20 minutes after the midazolam was administered. Administration of the second syringe of potassium chloride was stopped. The paramedic assessed the situation and concluded that the IV catheter was no longer penetrating the vein. The physician attempted IV insertion into the left femoral vein. The needle penetrated the femoral artery rather than the femoral vein. No left femoral IV access was established.

At this juncture, Director Patton asked Warden Trammell two questions. First, do you have another viable vein? And, second, do you have any more chemicals to push? Warden

Trammell told Director Patton that there was no viable vein.

After that, the physician told the warden and the warden relayed to Director Patton that not enough drugs had entered Lockett's body to cause death.

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The autopsy of Lockett indicated that there were concentrations of midazolam in the tissue near the insertion site in Lockett's right groin area. This indicated that the drugs were not flowing intravenously into Lockett's cardiovascular system and that this problem had existed as early as the stage at which the midazolam was being administered.

At 6:56 p.m., after two conversations with Governor Fallin's counsel, Director Patton terminated the execution process, although the administration of the drugs had been stopped at about 14 minutes before that. Witnesses were escorted out of the viewing room. At 7:06 p.m., Lockett was pronounced dead. He died as a result of the lethal injection. The drugs that were intended to be lethal had their intended effect.

As to the opportunity for the witnesses to see what was happening, the blinds between the execution chamber and the viewing rooms were raised and lowered once during Lockett's execution. The individuals who viewed the execution room from the viewing room had been seated in the viewing room by 6 p.m. At approximately 6:23 p.m., after Director Patton received

approval from the governor's office to proceed with the execution, the blinds were raised. The blinds were still up when the physician inspected the femoral IV insertion site and concluded that there was a problem with IV access. At 6:42 p.m., when the administration of the second syringe of potassium chloride was stopped, the blinds were lowered. The blinds remained down from that point until Lockett was pronounced dead.

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The autopsy disclosed wounds consistent with the IV access attempts that I have described. The autopsy also indicated that both midazolam and vecuronium bromide were found in the psoas muscle indicating that those chemicals had been distributed throughout Lockett's body. The concentration of midazolam found in Lockett's blood was greater than the concentration required to render an average person unconscious.

I will now make my findings with respect to the revision of the lethal injection protocol and Department of Corrections practice under that protocol.

THE REVISION TO THE PROTOCOL AND DOC PRACTICE UNDER THAT PROTOCOL

As I have noted, the DOC adopted a new protocol entitled "Execution of Offenders Sentenced to Death" with an effective date of September 30, 2014. Including attachments, the revised protocol is 55 pages long.

I do not consider it necessary to go into a detailed

1 side-by-side comparison of the revised protocol versus the 2 previous protocol, but it is safe to say that with respect to 3 the matters that are most relevant here, specifically the procedures for establishing IV access to the offender's 4 5 cardiovascular system, the procedure for administering the 6 chemicals, and the procedures for dealing with mishaps or 7 unexpected contingencies, the new protocol is noticeably more detailed. The revised protocol also includes detailed 8 9 provisions with respect to training and pre-execution 10 preparation of the members of the execution team. The new protocol is in evidence as Plaintiffs' Exhibit 68. 11 12 A copy of the new protocol is also in the record at Docket Entry Number 55-1 filed on October 1, 2014. 13 14 The new protocol provides for several teams to participate 15 in and complete the execution process, including the 16 Intravenous Team, as indicated on page 7. The IV Team consists 17 of a team leader and one or more physicians, physician 18 assistants, nurses, emergency medical technicians, paramedics, 19 or a military corpsman, or other certified or licensed 20 personnel, including those trained in the U.S. Military. The team leader and members shall be "currently certified or 2.1 licensed within the United States." 22 2.3 Practitioners in some of those categories, such as physicians, must be licensed and others, like military 24 25 corpsman, are credentialed by certification. The new protocol

provides that a central femoral venous line shall not be used unless the person placing the line is currently certified or licensed within the United States to place a central femoral line. That is on page 27. That is not a particularly meaningful requirement because there is no licensing or certification specific to that procedure.

The team leader and members are selected by the director of the DOC on the basis, among other things, of the proposed team member's qualifications, training, experience, and professional licenses or certification, as indicated on page 7.

I'll now address training under the revised protocol.

TRAINING UNDER THE REVISED PROTOCOL

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Warden Anita Trammell acknowledged that after the Lockett execution she realized that the training of the execution team had been, in point of time, "up to bringing the offender into the execution chamber." She testified that "the training should have gone beyond that." I agree with that comment. That is at page 158 of the transcript. The new protocol does call for significantly more training.

For execution team members, the new protocol requires "ten training scenarios within the 12 months preceding the scheduled execution." That is on pages 9 and 10. The training section of the protocol provides for "multiple training scenarios," including but not limited to contingency plans for issues with execution equipment or supplies, issues with offender IV

access, including alternate IV access sites, issues if the offender is not rendered unconscious after administration of execution chemicals, and unanticipated medical or other issues concerning the offender or an execution team member, all as covered on page 10.

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Two days prior to the day of execution, the division manager is required to schedule and conduct "on-site scenario training sessions, modifying practices as warranted," as indicated on page 22. The training section of the revised protocol also requires that the IV Team members "shall participate in at least one training session with multiple scenarios within one day prior to the scheduled execution," as indicated on page 10. The H-Unit Section Chief is required to be trained in determining whether there is a problem with IV flow.

This fall, the execution team has been training one day a week for five or six hours in each instance. The director is included in these training sessions. They train for a minimum of six scenarios each time and sometimes address seven or eight scenarios. The training scenarios include, for instance, situations in which there is a problem with the IV manifold and drills involving access to the femoral vein.

In addition to the training sessions that have been conducted, a training session will be held with the IV Team within 24 hours before the next execution in which the IV Team

will address a scenario in which the inmate regains consciousness after having been pronounced unconscious.

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Thirty-five days prior to the day of execution, the protocol requires that the offender's medical condition "be assessed in order to identify any necessary accommodations or contingencies that may arise from the offender's medical condition or history," as indicated on page 16. This includes examination for "concerns for establishing or maintaining IV lines," as indicated on page 17.

The offender's telephone privileges are terminated at 9 p.m. on the day prior to the day of execution, except for calls from the offender's attorney of record and others as approved by the division manager. Likewise, visitation is terminated at 9 p.m. on the day before the day of execution, except that two hours of in-person visitation with up to two attorneys of record is permitted as long as that visitation ends two hours prior to the scheduled execution or earlier, if necessary, to begin preparing the offender for the execution, as indicated on page 22.

The revised protocol requires that an electrocardiograph and a backup electrocardiograph be on site available for use during the execution.

The execution team must prepare a complete set of the required chemicals. In addition, "An additional complete set of the necessary chemicals shall be obtained and kept available

in the chemical room," as indicated on page 38.

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As is set forth on pages 39 through 41 of the revised protocol, the new protocol gives the director four alternatives with respect to the combination of drugs to be used in the lethal injection process. These alternatives are set forth in Chart A, Chart B, Chart C, and Chart D.

Chart A calls for the administration of 5,000 milligrams of pentobarbital in a one-drug procedure. Chart B provides for the administration of 5,000 milligrams of sodium pentothal, again, in a one-drug procedure. Chart C provides for the administration of 500 milligrams of midazolam and 500 milligrams of hydromorphone. Hydromorphone is a narcotic analgesic. Finally, Chart D provides for the administration of 500 milligrams of midazolam, 100 milligrams of vecuronium bromide, and 240 milliequivalents of potassium chloride.

Rocuronium bromide will apparently be used in the upcoming executions, but it is not materially different from vecuronium bromide, aside from what Dr. Katz referred to as "a dosing change," which apparently is not in controversy.

The protocol provides that the director shall have the sole discretion to determine which chemicals will be used for the scheduled execution. This decision is required to be provided to the offender in writing ten calendar days before the scheduled execution date, as indicated on page 41. If it is necessary to use a compounded drug, the compounded drug

"shall be obtained from a certified or licensed compounding pharmacist or compounding pharmacy in good standing with their licensing board." The protocol requires a qualitative analysis of the compounded drug to be performed no more than 30 days before the execution date. The protocol also requires that the decision to use compounded drugs be provided to the offender in writing not less than ten calendar days before the scheduled execution, as indicated on page 41.

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The new protocol provides for the insertion of a primary IV catheter and a backup IV catheter. The primary line is referred to as the "A line." The secondary line is referred to as the "B line." IV access is established at two points, with the A line and the B line. If all goes as planned, the drugs would be entirely injected through the A line. If there is a problem with the A line or with IV access through the A line, then the B line is a backup. The complete second set of execution drugs is for use in the B line or otherwise, if need The preferred site for IV access is the "arm veins near the joint between the upper and lower arm," as indicated on page 26. If the IV Team is unable to establish an IV at a preferred site, the IV Team members are authorized to establish an IV at an alternative site, including a central femoral venous line. The protocol states that the IV Team shall be allowed as much time as necessary to establish viable IV sites, but that after one hour of unsuccessful IV attempts, the

director must contact the governor or the governor's designee to advise of the status and "potentially request a postponement of the execution," as indicated at pages 26 and 27.

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After insertion of the IV needle, the IV Team is required to test the viability of the IV site with a low-pressure saline drip through the IV tubing. If necessary, a Heparin lock may be attached to the IV needle as an alternative to the saline drip as indicated on page 27.

The protocol contains detailed provisions for monitoring the condition of the offender during the execution process, including a requirement that a microphone be affixed to the offender's shirt to enable the execution team "to hear any utterances or noises made by the offender throughout the procedure," as indicated on page 43.

For the purpose of monitoring the offender's cardiac status, the protocol requires that execution team members "attach the leads from the electrocardiograph to the offender's chest once the offender is secured. The IV Team leader shall confirm that the electrocardiograph is functioning properly. A backup electrocardiograph shall be on site and readily available if necessary. Prior to and on the day of the execution, both electrocardiograph instruments shall be checked to confirm that they are functioning properly," as indicated on page 43.

Finally, on the subject of monitoring, the protocol

requires that the IV Team leader "monitor the offender's level of consciousness and electrocardiograph readings utilizing direct observation, audio equipment, camera and monitor, as well as any other medically approved methods deemed necessary by the IV Team leader. The IV Team leader shall be responsible for monitoring the offender's level of consciousness," as indicated on page 43.

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Although the presence of a physician during the execution process is not required under the revised protocol, a physician has been selected as the IV Team leader for the executions of the four plaintiffs who seek a preliminary injunction. The second member of the IV Team will be a paramedic. Warden Trammell testified that the consciousness check will be performed by the physician.

The protocol requires that the IV catheter remain visible to the H-Unit Section Team Chief throughout the execution procedure and that the H-Unit Section Team Chief remain in the room with the offender "in a position sufficient to clearly observe the offender and the primary and backup IV sites for any potential problems." The Section Team Chief is required to immediately notify the IV Team leader and the director if any problem is observed, as indicated at page 44.

The H-Unit Section Team Chief is required to observe the offender during the injection process "to look for signs of swelling or infiltration at the IV site, blood in the catheter

and leakage from the lines, and other unusual signs or symptoms," as indicated on page 27. For that reason, it falls to the H-Unit Section Team Chief to determine whether it is necessary to use an alternative IV site. When an alternative IV site is used, the team members who administer the chemicals are required to administer a full dose of the execution drugs through the alternative site, as indicated on page 27.

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To facilitate the level of scrutiny required by the new protocol, the operations room adjacent to the execution chamber now has two video monitors with feeds from two cameras. Both cameras have tilt, turn, and zoom capability. One camera is located near the head of the gurney and the other camera is located near the foot of the gurney. With these cameras, the IV Team members can monitor the point of IV access on the offender's body.

The lapel microphone on the prisoner provides a continuous audio feed into the operations room where most of the participants in the execution process will be during the execution. This microphone comes on at the beginning of the execution and is not turned off until the offender is pronounced dead. There is a separate microphone over the offender's head. This microphone feeds to the viewing room and to the overflow room. This microphone is turned on when the blinds are raised and is turned off after the offender makes his final statement, if he chooses to make one. It is then

turned back on for the doctor to announce the results of the consciousness check and immediately turned off. It is then turned on for the announcement of the time of death.

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There are three telephones in the operations room: one to maintain contact with the governor's office, one to maintain contact with the Attorney General's Office, and one for the purposes of communications within the Oklahoma State Penitentiary. In addition, there is an intercom between the execution chamber and the operations room.

With respect to the actual administration of the lethal injection chemicals, the new protocol has one procedure for administration of the chemicals specified in Charts A, B, and C and a slightly different procedure for administration of the three-drug sequence specified in Chart D, as indicated at pages 44 to 46.

With respect to all four chemical charts, the IV Team leader is required to enter the room where the offender is located "to physically confirm the offender is unconscious by using all necessary and medically appropriate methods," as indicated on pages 44 and 46.

If there is a delay in losing consciousness, the IV Team is required to inform the director so that the director can determine how to proceed or whether to "start the procedure over at a later time or stop," as indicated on pages 45 and 46. The director has the discretion to instruct the execution team

to administer additional doses of the chemical, as indicated, again, on pages 45 and 46.

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The execution facility is now equipped with an electrocardiograph machine to monitor the offender's blood pressure, oxygen saturation, and heart activity during the execution process. A backup machine is available in the event there is a problem with the primary machine. The DOC also purchased an ultrasound machine for use, if need be, in locating a deep vein and the stock of surgical supplies has been improved, to include a newly acquired assortment of IV needles. The newly acquired assortment of IV needles was shown to Captain Holt during his recent tour of the execution facility.

The revised protocol calls for an after-action review following the execution. This after-action review includes discussion of "any unique or unusual events," as well as "opportunities for improvement and successful procedures," as indicated on page 31.

I will now make my findings with respect to the unavailability of sodium thiopental and pentobarbital.

UNAVAILABILITY OF SODIUM THIOPENTAL AND PENTOBARBITAL

In paragraph 31 of their amended complaint, plaintiffs proffer sodium thiopental used in a single-drug protocol as their alternative to midazolam, as indicated at Docket Entry Number 75 at page 7. At the trial last week, pentobarbital was

also mentioned several times.

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Pentobarbital and sodium thiopental are powerful barbiturates. A massive dose of either of these drugs is lethal. Which is why, as long as they were available, they had a well-established record of successful use in execution by lethal injection, even in drug combinations in which it was not necessary that they have lethal effect.

Pentobarbital and sodium thiopental are both unavailable to the Department of Corrections. Attempts to procure pentobarbital and sodium thiopental have been unsuccessful. Ιt is a judicially noticeable fact that the Lockett execution was preceded by a storm of litigation involving the state and federal district courts as well as the Oklahoma Supreme Court and the Oklahoma Court of Criminal Appeals. The litigation included intense efforts to force the disclosure of the sources of the lethal injection drugs as noted in Lockett v. Evans, 2014 Westlaw 1584517, a decision from the Oklahoma Supreme Court on April 21, 2014. Former Oklahoma Department of Corrections General Counsel Michael Oakley, who retired shortly before the Lockett execution, testified quite believably that "the vendor, because of pressure in the litigation, decided that he didn't want to sell us the pentobarbital any longer." That is at page 296 of the transcript.

Director Patton cannot think of anything he could have

done differently in his efforts to get these drugs and the

Court credits this testimony. The DOC talked to numerous

pharmacies, including compounding pharmacies, in its efforts to

procure pentobarbital and sodium thiopental, either

commercially manufactured or compounded. These efforts were

not successful. Sodium thiopental and pentobarbital are

6 certainly known alternatives, but it is equally clear that 7 they're not available to the DOC.

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I will now analyze the evidence derived from the execution of Clayton Lockett.

ANALYSIS OF EVIDENCE DERIVED FROM THE EXECUTION OF CLAYTON LOCKETT

Dr. Eric D. Katz, a well-qualified emergency physician, had some credible criticisms of the process by which Clayton Lockett was executed. IV access to Clayton Lockett's right femoral vein was established, as I have said, with a one-and-a-quarter-inch 14-gauge angiocatheter. The one-and-a-quarter-inch 14-gauge angiocatheter was not the appropriate equipment to use to accomplish this task. And the use of a catheter of that size substantially increased the risk of serious difficulties in establishing patent intravenous access by way of Clayton Lockett's right femoral vein. It is now common in medical practice to use ultrasound equipment for guidance in establishing intravenous access to the subclavian, internal jugular, or femoral veins.

Dr. Katz also commented with respect to the requirement in

the revised protocol that the person placing a central femoral line be "currently certified or licensed within the United States to place a central femoral line." As I have previously mentioned, Dr. Katz pointed out that this specific task is not one for which certification or licensure is available.

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Dr. Katz also commented on the medications and techniques which would be necessary to reverse the effect of midazolam, vecuronium, and potassium. Aside from the medications that would be required to reverse the effects of these drugs, reversal of the effects of vecuronium would require a ventilator and reversal of the effects of a high dose of midazolam would likely require endotracheal intubation, in other words, a breathing tube, with supplies for ventilator assistance. Dr. Katz pointed out that these medications and this equipment necessary for resuscitation of an individual affected by midazolam, vecuronium bromide, and potassium chloride were not available at the time of Clayton Lockett's execution.

Dr. Joseph I. Cohen and Dr. Joni McClain, both well-qualified pathologists, testified with respect to their respective autopsy examinations of the body of Clayton Lockett. Dr. Cohen was called by the plaintiffs and Dr. McClain, who performed the independent autopsy at the Southwestern Institute of Forensic Sciences in Dallas, was called by the defendants. Their observations from their actual examinations did not

substantially differ. Dr. Cohen conducted his autopsy in Tulsa on May 14, 2014. Dr. Cohen's autopsy followed the autopsy performed by the Southwestern Institute of Forensic Sciences and the partial examination that was performed by the Oklahoma State Medical Examiner.

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Dr. Cohen found numerous punctures and incisions consistent with the several reported attempts to gain IV access at several sites on Clayton Lockett's body, which was consistent with Dr. McClain's findings. Dr. Cohen found, and this is uncontradicted, that Lockett's veins were in good condition and suitable for establishing IV flow of the lethal injection drugs. Dr. Cohen concluded that the manner of death was judicial execution and that the mechanism of death was respiratory depression and cardiac dysrhythmias directly resulting from the administration of the lethal injection drugs during the execution process.

Commenting on the numerous fresh punctures observable in Clayton Lockett's body, Dr. Cohen stated that his findings, as well as the findings of the Oklahoma state medical examiner and the Dallas medical examiner, support the ineffective application of medical implements. This is accurate.

Dr. Cohen also opined that Clayton Lockett was not dehydrated during the execution and that he likely suffered conscious pain and suffering due to the failed attempts to establish IV access. I have already commented on the issue of

whether Lockett experienced pain beyond that inherent in the execution process. As I have discussed, Lockett may well have experienced significant pain, but any such conclusion is laden with an element of speculation.

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I will now comment on the nature and characteristics of midazolam and the other drugs at issue.

NATURE AND CHARACTERISTICS OF MIDAZOLAM AND THE OTHER DRUGS AT ISSUE

Plaintiffs called two witnesses who addressed the characteristics of midazolam; namely, Dr. Larry D. Sasich, a pharmacist who holds a Bachelor of Science degree in pharmacy and a doctoral degree in pharmacy; and Dr. David A. Lubarsky, a Professor of Anesthesiology at the University of Miami.

Dr. Sasich also commented on the characteristics of vecuronium

The defendants called Dr. Lee R. Evans, the holder of a doctoral degree in pharmacy, principally to testify with respect to the characteristics and effects of midazolam.

bromide and potassium chloride.

Now, several days before the hearing, a Daubert motion was filed with respect to the expert testimony of Dr. Lee Evans. I indicated at the pretrial conference that I would address that -- as permitted in a non-jury case, that I would address that after hearing his testimony by way of direct and cross-examination at trial. And I certainly did hear his testimony by way of direct and cross-examination. I reviewed the motion

in limine. I made careful note of the extent to which his testimony was challenged in the motion in limine and the subjects on which it was challenged.

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The Tenth Circuit has made it very clear that once a Daubert challenge is filed, the Court must make its findings on the record indicating its resolution of the Daubert challenge.

And I will now do that at this time.

DAUBERT RULING WITH RESPECT TO THE EXPERT TESTIMONY OF DR. LEE EVANS

On December 15, as I indicated, plaintiffs filed a motion in limine challenging some specific aspects of the proposed expert testimony of Dr. Lee Evans. That motion is at Docket Entry Number 161. For that reason, before making any findings based on the testimony given by the expert witnesses called by the parties, it is necessary to address the Daubert challenge as to Dr. Evans.

The Supreme Court's decisions in Daubert v. Merrell Dow Pharmaceutical, Inc., 509 U.S. 579, and Kumho Tire Company v. Carmichael, 526 U.S. 137, establish a gatekeeper function for trial judges under Rule 702 of the Federal Rules of Evidence. This is commented on at considerable length in the Tenth Circuit's two Goebel decisions, the first one of which is Goebel v. Denver and Rio Grande Western Railroad Company, 215 F.3d 1083, with the relevant discussion at page 1087, a decision from the Tenth Circuit in 2000. The gatekeeper

function "requires the judge to assess the reasoning and methodology underlying the expert's opinion and determine whether it is scientifically valid and applicable to a particular set of facts," as indicated in the first Goebel decision at page 1087.

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In Kumho, the Supreme Court elaborated on the Daubert gatekeeping function as applied to proposed expert testimony other than classic scientific testimony. The Court emphasized that even where the proposed expert testimony is not scientific in nature in the classical sense, the trial judge is nevertheless required to ascertain whether the expert "employs in the courtroom the same level of intellectual rigor that characterizes the practice of an expert in the relevant field," as indicated at page 152 of the Kumho decision.

In this case, the plaintiffs' motion in limine challenges both Dr. Evans' qualifications and his methodology, so it is necessary to analyze the matter under both parts of the Daubert and Rule 702 test.

The decision in Ralston, 275 F.3d 965, provides a good starting point with respect to evaluation of Dr. Evans' qualifications. The plaintiff in Ralston asserted that the warnings accompanying an implanted orthopedic nail were inadequate. The Court of Appeals affirmed the trial court's exclusion of the testimony of plaintiffs' expert, a board certified orthopedic surgeon, who was also an associate

1 professor of medicine. The expert's general credentials were 2 clearly as good as reasonably could have been expected, but she 3 had done no research specifically looking at the nail in question and had not drafted a warning for a surgical device. 4 5 Her general credentials, though seemingly impressive as general 6 credentials, were not good enough. "Merely possessing a 7 medical degree is not sufficient to permit a physician to testify concerning any medical-related issue," as the Court 8 9 discussed at page 970. The board certified orthopedic 10 surgeon's reliance on general principles and concepts, as the 11 Court put it, did not suffice. The controlling Tenth Circuit 12 cases exemplified by Ralston established that the expert's qualifications must be both adequate in a general qualitative 13 14 sense as required by Rule 702 and specific to the matters he 15 proposes to address as an expert. 16 Plaintiffs also challenge the reliability of Dr. Evans' 17 expert testimony. 18 Under Rule 702, an expert with the necessary

Under Rule 702, an expert with the necessary qualifications in the relevant field may give expert testimony, one, if the testimony is based on sufficient facts or data; two, if the testimony is the product of reliable principles and methods; and, three, the witness has applied the principles and methods reliably to the facts of the case, as indicated by Rule 702. And this is also generally discussed in the second Goebel decision, 346 F.3d 987, with the relevant portion at page 991.

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Daubert, of course, involved a proffer of expert testimony in a classical scientific discipline, epidemiology. The Court provided a non-exclusive list of five factors which were provided by the Court to guide trial court determinations of reliability. I will not repeat those lists — those factors here. Counsel are well aware of them.

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Kumho made it clear that the gatekeeper function applies even where the proposed expert testimony is outside the realm of science in the classical sense, as I have discussed. As noted by the Advisory Committee in commenting on the 2000 amendments to Rule 702, courts both before and after Daubert have found other factors relevant in determining whether expert testimony is sufficiently reliable to be considered by the jury. Those additional factors are listed in the comments to Rule 702 and I will not repeat them here.

In sum, the Daubert assessment of reliability is a determination of whether the conclusions to be expressed by an expert possessed of the necessary qualifications in the relevant field are the product of application of that expertise using recognized and supportable methodologies on the basis of adequate data which is rationally tied to the opinions which purport to be based on that data.

As indicated in the second Goebel decision, 346 F.3d at 992, "Under Daubert, any step that renders the analysis unreliable renders the expert's testimony inadmissible. This

is true whether the step completely changes a reliable methodology or merely misapplies that methodology." To the same general effect is Mitchell v. Gencorp, 165 F.3d 778, with the relevant discussion at page 782, a Tenth Circuit decision from 1999.

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I reject plaintiffs' challenge to Dr. Evans' qualifications. Dr. Evans' CV is in evidence as Defendants' Exhibit 35 and I will not repeat that information here. His qualifications go far beyond those of an everyday pharmacist and his clinical experience is an obvious adjunct of his academic attainments, at least as relevant to this case.

I might add that Dr. Sasich, called by the plaintiffs, freely went beyond pharmacological topics and expounded on physiology and to some extent clinical medical practice based substantially on his searches of literature he considered to be relevant. Dr. Evans' considerable qualifications satisfy me that he should be accorded the same leeway.

It is necessary to evaluate the reliability of Dr. Evans' expert testimony only to the extent that the portions of his testimony that I cite in this ruling have been made explicitly — have been explicitly challenged by plaintiffs in their motion in limine. For that reason, there is no need to engage in a reliability analysis of all of the matters testified to by Dr. Evans or discussed in his report and there is no need to dwell on the fact that he misplaced a decimal point in one of

his observations about the possible lethal effect of midazolam.

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Those aspects of Dr. Evans' testimony, upon which I principally rely, are his findings with respect to the risk that a 500 milligram dose of midazolam will fail to induce a state of unconsciousness and his criticisms of the contention that there is a ceiling effect that is relevant to the determination of whether the prisoner will experience pain after IV administration of 500 milligrams of midazolam. His commentary in part 7 of his report about brain-dead patients and involuntary movements is of no moment to my ruling. With respect to the issue of whether Lockett's movements were voluntary or involuntary, the fact is that he, in all probability, had far less than 500 milligrams of midazolam in his circulatory system at the time he moved after being pronounced unconscious.

I find that Dr. Evans was well-qualified to give the expert testimony that he gave and that to the extent that his testimony was challenged in the motion in limine and relied upon by the Court in this ruling, his testimony was the product of reliable principles and methods reliably applied to the facts of this case. The motion in limine with respect to Dr. Evans, Docket Entry Number 161, is denied.

I will now address the nature and characteristics of midazolam.

NATURE AND CHARACTERISTICS OF MIDAZOLAM

With respect to the actual characteristics, effects, and preferred clinical uses of midazolam, the experts on the two sides of this case were in substantial agreement.

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Midazolam is a short-acting benzodiazepine, which is most commonly used as a pre-anesthetic agent for routine medical procedures. Midazolam is approved by the Food and Drug Administration for sedation and induction of general anesthesia to be used before administration of other anesthetic agents. It can be used to alleviate patient apprehension, to eliminate the patient's memory of a procedure, and to induce anesthesia. It is not intended for use as a pain reliever. It is not an analgesic. In that respect, midazolam does not behave like an opiate or narcotic medications. It does have the effect of depressing the central nervous system at least when administered in a large dose. It begins to take effect quite quickly after introduction into the blood stream. It crosses the blood brain barrier and reaches maximum effects within 20 to 60 minutes.

The 500 milligram dosage of midazolam, as called for in Charts C and D of the revised protocol, is many times higher than a normal therapeutic dose of midazolam. When midazolam is administered in that quantity, it will result in central nervous system depression as well as respiratory arrest and cardiac rest. In the dosage called for by the revised protocol, midazolam, although not an analgesic, is highly

likely to render the person unconscious and insensate during the remainder of the procedure. Consequently, analgesia, from midazolam or otherwise, is not necessary.

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The proper administration of 500 milligrams of midazolam, as specified in Chart D, would make it a virtual certainty that any individual will be at a sufficient level of unconsciousness to resist the noxious stimuli which could occur from the application of the second and third drugs — or from the administration of the second and third drugs in Chart D, assuming that proper intravenous access has been established. The administration of a 500 milligram dose alone would be likely to cause death by respiratory arrest within an hour and probably closer to 30 minutes. This is because midazolam is water soluble. And as I have mentioned, it crosses the blood brain barrier very quickly.

There were some noteworthy areas of agreement between Dr. Evans and Dr. Lubarsky with respect to the anesthetic effect of midazolam. Dr. Lubarsky testified that an IV dose of 500 milligrams of midazolam would produce unconsciousness in "no more than a couple of minutes." That is at page 117 of the transcript. As to the level of unconsciousness needed, for instance, to render a prisoner insensate for purposes of setting a femoral IV line, Dr. Lubarsky testified that "midazolam unconsciousness is actually sufficient." That is at page 133 of the transcript. This is noteworthy not because

midazolam was used for that purpose with Lockett but because setting a femoral line entails "digging deeper into the tissue," as described by Dr. Lubarsky, "a couple of inches below the skin's surface," as indicated at page 150 of the transcript.

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Plaintiffs contend that there is a certain dosage level beyond which incremental increases in midazolam dosage would have no corresponding incremental effect. In pharmacological terms, this is called "the ceiling effect." As described by Dr. Sasich and Dr. Lubarsky, midazolam has a ceiling effect which prevents an increase in dosage from having a corresponding incremental effect on anesthetic depth. Dr. Evans testified persuasively, in substance, that whatever the ceiling effect of midazolam may be with respect to anesthesia, which takes effect at the spinal cord level, there is no ceiling effect with respect to the ability of a 500 milligram dose of midazolam to effectively paralyze the brain, a phenomenon which is not anesthesia but does have the effect of shutting down respiration and eliminating the individual's awareness of pain. The dosage at which the ceiling effect may occur at the spinal cord level is unknown because no testing to ascertain the level at which the ceiling effect occurs has been documented.

The use of midazolam presents a risk of paradoxical reactions or side effects such as agitation, involuntary

movements, hyperactivity, and combativeness. According to the product label for midazolam, these reactions may be the result of inadequate or excessive dosing or improper administration of The likelihood that a paradoxical reaction will midazolam. occur in any particular instance is speculative, but it occurs with the highest frequency in low therapeutic doses. estimated that with a low therapeutic dose of midazolam there would be less than a 1 percent incidence of a paradoxical reaction. Dr. Sasich could not say whether the incidence of a paradoxical reaction in the Oklahoma inmate population would be toward the low end or the high end of the range of incidence of that effect as documented in the literature. No data are available to show what the paradoxical reaction would be or the likelihood of a paradoxical reaction would be with a 500 milligram IV dose of midazolam.

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The evidence falls well short of establishing that the risk of a paradoxical reaction at a 500 milligram IV dosage presents anything more than a mere possibility in any given instance that midazolam will fail to deliver its intended effect.

Based on the impressive record of pentobarbital and sodium thiopental, in a long series of executions by lethal injection in Oklahoma and other states, there is little room for doubt that pentobarbital and sodium thiopental would be preferable as the first drug in a three-drug protocol. With midazolam, there

may be some incrementally greater risk than with pentobarbital or sodium thiopental that the inmate will sense pain as a result of the injection of vecuronium bromide and the potassium chloride but will not have the ability to express the fact that he senses pain. How much greater that risk is, nobody knows, but some added element of risk of pain may be present with midazolam as opposed to pentobarbital or sodium thiopental.

Ironically, the very efficacy of pentobarbital and sodium thiopental for use in lethal injection has resulted in their unavailability for use by the Department of Corrections for lethal injection purposes. If either pentobarbital or sodium thiopental were available, Director Patton would have selected them rather than the midazolam protocols.

On this point, I am mindful that the Supreme Court in Baze v. Rees, 553 U.S. 35, a decision from 2008, made it clear that this Court is not to sit as "a board of inquiry charged with determining best practices for executions." That's at page 51 of the Baze decision.

I'll now comment with respect to the capabilities or properties of vecuronium bromide.

VECURONIUM BROMIDE

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Vecuronium bromide is a neuromuscular blocking agent. The accounts of the Lockett execution which indicate that he began to move after the beginning of the administration of vecuronium bromide indicate that at that point he had not been paralyzed

1 | by the neuromuscular blocking effect of vecuronium bromide.

From all the evidence before the Court, I conclude that the implementation of lethal injection per Chart D does not carry a substantial likelihood of inflicting severe pain.

We'll now take a ten-minute recess.

(RECESS HAD.)

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THE COURT: I'll now summarize the claims asserted by the plaintiffs for preliminary injunction purposes.

CLAIMS ASSERTED BY THE PLAINTIFFS FOR PRELIMINARY INJUNCTION PURPOSES

The amended complaint filed on October 31, 2014, pleads claims divided into eight counts, some of which, as might be expected, are interrelated. Five of those counts, specifically Counts 2, 4, 5, 7, and 8, are asserted for preliminary injunction purposes as shown by the motion for preliminary injunction at Docket Entry Number 92.

In Count 2, the preliminary injunction plaintiffs assert that the use of midazolam would constitute cruel and unusual punishment in violation of the Eighth and Fourteenth Amendments.

Count 4 asserts violations of the Eighth and Fourteenth

Amendments based on what plaintiffs describe as "unsound

procedures and inadequate training," as indicated at page 28 of

Docket Entry Number 75. Under this heading, plaintiffs assert

that the failure of Defendants Patton and Trammell "to seek out

expert assistance has resulted in execution procedures that create a substantial risk of severe pain, needless suffering, and a lingering death," as indicated at paragraph 137.

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Count 4 also complains that ultimate authority to supervise the execution process and make decisions about the process is vested in the Defendant Patton without "appropriate checks and balances to ensure against severe pain, needless suffering, and a lingering death during the execution process." That is at paragraph 155.

Plaintiffs further assert that as a result of these failures, among others, there is "a substantial risk that the procedures will not be administered as written." That's paragraph 156.

Referring to the revised protocol, plaintiffs assert in paragraph 160 that "if the attempted executions of plaintiffs are allowed to proceed in accordance with the deficient procedures identified above, plaintiffs will be subjected to cruel and unusual punishment in violation of the Eighth and Fourteenth Amendments to the United States Constitution."

In Count 5, under "Notice and Opportunity to be Heard, Right to Counsel and to Petition the Courts," plaintiffs point out that under the revised protocol, the offender will be notified only ten calendar days before the date of execution of the drugs to be used and as to whether they will be compounded. This is paragraph 163.

In this count, plaintiffs also assert that the revised protocol allows Defendant Patton "to deviate from any of those procedures at will and without notice, thereby making the written instrument virtually meaningless as a form of notice." That is paragraph 164. On this basis, plaintiffs conclude in Count 5 that by "failing to require and provide meaningful and effective notice of how plaintiffs will be executed," defendants are depriving the plaintiffs of their right to notice and an opportunity to be heard in violation of the Due Process Clause of the Fourteenth Amendment and are subjecting plaintiffs to cruel and unusual punishment in violation of the Eighth and Fourteenth Amendments, as indicated in paragraph 169.

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Count 7 is an Eighth Amendment claim predicated on "experimentation on captive human subjects." Plaintiffs assert that by "attempting to conduct executions with an ever-changing array of untried drugs of unknown provenance, using untested procedures, defendants are engaging in a program of biological experimentation on captive and unwilling human subjects," as indicated at paragraph 184. In support of this allegation, plaintiffs cite the experience with the execution of Clayton Lockett, which they assert was "a failure that produced severe pain, needless suffering, and a lingering death," at page — at paragraph 184.

Under this heading, plaintiffs further assert that the

defendants lack the scientific skills necessary to design an execution procedure that does not inflict severe pain, needless suffering, or a lingering death and that defendants have failed to test their lethal drugs and execution procedures on non-human animals before using them on captive and unwilling human subjects as pleaded at paragraphs 187 and 188. On this basis, plaintiffs assert that if the attempted executions of plaintiffs are allowed to proceed plaintiffs will be subjected to cruel and unusual punishment in violation of the Eighth and Fourteenth Amendments to the United States Constitution.

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In Count 8, plaintiffs assert a violation of their "right of access to information, to counsel, and to the courts."

Under this heading, plaintiffs assert a violation of their rights as a result of the defendants' "deliberate concealment of information that would enable plaintiffs to determine how defendants intend to carry out their death sentences, including by failing to disclose in advance of the execution details about the drugs used, the rationale for the selection of these drugs and their dosages, the qualifications and training of the persons administering them, and defendants' ability to respond and prepare for responding to complications," as pleaded at paragraph 195.

They assert a violation of their right to petition the government for redress of their grievances as well as a denial of their right to counsel, which they assert exists "during

1 every stage of any attempt to execute" the plaintiff. at paragraph 198. 2 As for violation of the right to counsel, 3 plaintiffs assert, in substance, that plaintiffs' counsel must be permitted to observe all steps of the execution process from 4 5 the time that the offender is brought into the execution 6 chamber until the offender is pronounced dead. That is at page 207. 7

Plaintiffs also assert that their counsel "must be able to communicate with the courts as to any problems or deviations that occur during the execution that impact plaintiffs' substantial rights." That is at paragraph 208. For these reasons, among others, plaintiffs assert in Count 8 that execution under the revised protocol would violate their rights under the First and Fourteenth Amendments to the United States Constitution and under 18 United States Code, Section 3599, as indicated in paragraph 218.

I will now address the standards for entry of a $\ensuremath{\mathsf{preliminary}}$ injunction.

STANDARD FOR ENTRY OF A PRELIMINARY INJUNCTION

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To establish that preliminary injunctive relief is appropriate, plaintiffs must demonstrate, first, that they will likely succeed on the merits of their claim; second, that without preliminary relief they will suffer irreparable harm; third, that the balance of equities tips in their favor; and, fourth, that entry of an injunction is in the public interest.

That is all as discussed by our Court of Appeals in Kikumura v.

Hurley, 242 F.3d 950, with the relevant discussion at page 955.

That's a decision from the Tenth Circuit in 2001.

The injunctive relief the plaintiffs seek here is not in the disfavored category, so there is no basis for the Court to apply the heightened standard that would govern if the injunctive relief sought by plaintiffs were in one of the disfavored categories. As indicated by the Court of Appeals in the Kikumura decision, citing Otero Savings & Loan Association v. Federal Reserve Bank of Kansas City, 665 F.2d 275, with the relevant discussion at page 278, a decision from the Tenth Circuit in 1981. When the other three requirements for a preliminary injunction are satisfied, "it will ordinarily be enough that the plaintiff has raised questions going to the merits so serious, substantial, difficult, and doubtful as to make them a fair ground for litigation." That is at page 955 of the Kikumura decision.

CONCLUSIONS OF LAW

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I will now state my conclusions of law. I don't think it would be particularly helpful to the parties or to a reviewing court for me to simply declare some abstract principles of law, most of which are well-established. What is decisive in this case is the application of those principles to the facts of this case. That will be my main focus. And that will require some discussion of the facts as I state my conclusions of law.

As I have already mentioned, the motion now before the Court involves five of the eight counts in the amended complaint; namely, Counts 2, 4, 5, 7, and 8. I will discuss those in the order in which they appear in the amended complaint.

I will first address Count 2, which is the Eighth Amendment claim relating to midazolam.

COUNT 2 - EIGHTH AMENDMENT - MIDAZOLAM

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In Count 2, plaintiffs assert that their Eighth Amendment right to be free of cruel and unusual punishment would be violated if, in the execution process, midazolam -- if, in the execution process, midazolam is used as provided in the revised protocol.

Other drugs, such as sodium thiopental, have been used as the first drug in the execution sequence with a longer record than midazolam has of reliably producing the desired effect.

For that reason, it is not necessary to look past sodium thiopental to say with considerable confidence that if all of the — if all of the potentially usable anesthetic and sedative agents produced by the pharmaceutical industry were equally available to the DOC, it is not likely that midazolam would be the first choice. Director Patton's testimony makes that clear. This makes it especially important to proceed with a thorough understanding of the standard established by the Supreme Court in Baze v. Rees.

Baze v. Rees, 553 U.S. 35, a decision, as I have mentioned, from 2008, came to the Supreme Court from Kentucky where the protocol called for the administration of 3,000 milligrams of sodium thiopental, 50 milligrams of pancuronium bromide, and 240 milliequivalents of potassium chloride. Kentucky protocol before the Supreme Court in Baze v. Rees provided for IV insertion by "qualified personnel having at least one year of professional experience." That is 553 U.S. at page 45. In practice, Kentucky used a certified phlebotomist and an emergency medical technician to perform the venipunctures necessary for the catheters, as indicated at page The protocol allowed up to one hour within which to establish both primary and secondary peripheral intravenous sites in the arm, leg, hand, or foot of the inmate, as indicated also on page 45. In Baze, the Court noted as a preliminary matter that "it

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In Baze, the Court noted as a preliminary matter that "it is uncontested that failing a proper dose of sodium thiopental that would render the prisoner unconscious, there is a substantial constitutionally unacceptable risk of suffocation from the administration of pancuronium bromide and pain from the injection of potassium chloride," as discussed at page 53. In Baze, the prisoner asserted, among other things, that the protocol was deficient because it was "possible that the IV catheters will infiltrate into surrounding tissue causing an inadequate dose to be delivered to the vein because of

inadequate facilities and training and because Kentucky has no reliable means of monitoring the anesthetic depth of the prisoner after the sodium thiopental has been administered."

And that is at page 54.

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Under the Kentucky protocol, the warden and deputy warden, who apparently were not subject to any particular training requirements, were charged with the responsibility to "watch for any problems with the IV catheters and tubing," as discussed at pages 45 and 46. In the Kentucky procedure, the physician was prohibited from participating in conducting the execution other than to certify the cause of death, as also discussed on page 46.

It also fell to the warden and the deputy warden, "through visual inspection," to determine whether the prisoner had become unconscious within 60 seconds following the delivery of the sodium thiopental to the primary IV site, as discussed on page 45.

The plurality opinion in Baze was written by Chief Justice Roberts and was joined by Justices Kennedy and Alito. Justices Stevens, Scalia, Thomas, and Breyer concurred in the judgment. As indicated by the concurring opinion of Justice Thomas, with whom Justice Scalia joined, Justices Scalia and Thomas would find an Eighth Amendment violation only if a method of execution "is deliberately designed to inflict pain," as discussed on page 94. The Supreme Court's decision in Marks v.

United States, 430 U.S. 188, from 1977, tells us at page 193 that where there is a fractured decision of the Supreme Court, the Court's holding is the position taken by those members of the Court who concurred in the judgment on the narrowest grounds. Since the position taken in Baze by Justices Scalia and Thomas was noticeably less exacting than the position taken by the Chief Justice and Justices Kennedy and Alito in the plurality opinion, the Court's holding in Baze is to be found in the plurality opinion.

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Before getting into the holdings in Baze, one other aspect of that decision should be noted. In Baze, the petitioner's arguments centered mainly on the asserted risk of improper administration of sodium thiopental, which would potentially leave the prisoner conscious when the second and third chemicals are administered. Counsel for the petitioner in Baze acknowledged at oral argument that proper administration of the first drug, sodium thiopental, would eliminate any meaningful risk that the prisoner would experience pain from the subsequent injections of the second and third drugs, as discussed on page 49.

In contrast, in the matter now before this Court, plaintiffs assert both the risk of maladministration of the drugs and that the first drug, midazolam, is, in any event, unreliable as an anesthetic agent. The holdings in Baze are nevertheless instructive and certainly binding on this Court

because regardless of the source of the risk, the Baze decision was all about risk and how we evaluate that risk for Eighth Amendment purposes.

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After a detailed analysis of the Kentucky lethal injection protocol, viewed in light of the Supreme Court's precedence, the plurality in Baze summarized the applicable standard as follows: "A stay of execution may not be granted on grounds such as those asserted here unless the condemned prisoner establishes that the state's lethal injection protocol creates a demonstrated risk of severe pain. He must show that the risk is substantial when compared to the known and available alternatives. A state with a lethal injection protocol substantially similar to the protocol we uphold today would not create a risk that meets this standard," as this Court stated at page 61.

In closing argument in this case last Friday, plaintiffs argued that Baze is distinguishable from this case in a way that makes the holding in Baze with respect to "known and available alternatives" inapplicable in this case. I disagree. It is true that this case involves both the risk that the first drug will not have its intended effect even if delivered in a massive IV dose and the risk that, due to a deficient technique, the massive IV dose will not, in fact, be delivered as intended.

In the section of the plurality opinion on page 61 in

1 which the Supreme Court made reference to comparison with 2 "known and available alternatives," the Court was speaking 3 broadly in terms of whether the prisoner had established "that the state's lethal injection protocol creates a demonstrated 4 5 risk of severe pain." In stating that the prisoner "must show 6 that the risk is substantial when compared to the known and available alternatives," the Court did not differentiate 7 between the two types of risks. 8

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On this point, I would also note that this reading of Baze is also borne out by common sense. It is extremely unlikely that the Supreme Court would establish a constitutional doctrine that would enable a condemned inmate to block his execution on Eighth Amendment grounds with no consideration by the Court of alternatives which by way of comparison demonstrate the constitutional unacceptability of the risk complained of by the prisoner. Logic tells us that alternatives are relevant in determining whether there is a constitutionally impermissible quantum of risk regardless of the source of the risk.

In reaching its conclusions in Baze, the Court made several other observations which are of varying degrees of relevance on the facts of this case.

First, it is significant, at least at a high level of generality, that the Court in Baze noted that the Supreme Court had never invalidated a state's chosen procedure for carrying

out a sentence of death as the infliction of cruel and unusual punishment, as discussed at page 48. The Court also stated unequivocally that simply because an execution method may result in pain, either by accident or as an inescapable consequence of death, does not establish the sort of objectively intolerable risk of harm that qualifies as cruel and unusual within the meaning of the Eighth Amendment, as discussed at page 50.

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Elaborating on that point, also on page 50, the Court told us that an isolated mishap alone does not give rise to an Eighth Amendment violation precisely because such an event, while regrettable, does not suggest cruelty or that the procedure gives rise to a substantial risk of serious harm. Because Baze, like the case now before this Court, is really all about risk and how we evaluate and attribute significance to that risk, it is also helpful to bear in mind the risks which individually and collectively were insufficient to support granting relief to the petitioner in Baze.

The petitioner in Baze asserted that there was a risk of improper administration of the lethal injection drugs because the doses were difficult to mix into solution and to load into the syringes, the protocol failed to establish a rate of injection, there was a risk of infiltration of drugs into the surrounding tissue, Kentucky's execution facilities and training of the execution team were inadequate, and there was

no reliable means of monitoring the anesthetic depth the prisoner had reached, all as discussed on page 54. These risks, individually and collectively, were insufficient to support a grant of relief in Baze.

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As a necessary corollary to its main holding in Baze, the Court also stated unequivocally that "an inmate cannot succeed on an Eighth Amendment claim simply by showing one more step the state could take as a fail-safe for other independently adequate measures, " as discussed on pages 60 and 61. Supreme Court expressly rejected the notion that federal courts should, in effect, sit as "boards of inquiry charged with determining best practices for executions with each ruling supplanted by another round of litigation, touting a new and improved methodology, as discussed at page 51. The Court expressly noted that the best practices approach "calling for the weighing of relative risks without some measure of deference to a state's choice of execution procedures would involve the courts in debatable matters far exceeding their expertise, " as discussed in note 2 on page 51.

In her dissenting opinion, Justice Ginsburg lamented that there were several shortcomings in the Kentucky protocol that, in her view, regrettably did not make any difference to the majority. This included the lack of safeguards to determine whether the inmate was unconscious before injection of the second and third drugs, the fact that only the warden and the

1 deputy warden remained in the execution chamber after placement 2 of the catheters, the lack of any medical training on the part 3 of the warden and the deputy warden, the reliance only on visual observation to determine whether the inmate appeared to 4 5 be unconscious, and the failure to use reflex tests or noxious 6 stimulus to determine whether the prisoner was unconscious. 7 Those matters were discussed by Justice Ginsburg on pages 114 and 118. 8 9 The Tenth Circuit has had more than one opportunity to apply the principles established in Baze. The best example 10 would be the decision in Pavatt v. Jones, 627 F.3d 1336, a 11 Tenth Circuit decision from 2010. 12 In Pavatt, at pages 1338 and 39, the Tenth Circuit said --13 14 and here I am leaving out citations and internal quotations. 15 "In Baze, the Court acknowledged that subjecting individuals to 16 a risk of future harm, not simply inflicting pain, can qualify 17 as cruel and unusual punishment. However, the Court emphasized 18 to establish that such exposure violates the Eighth Amendment, 19 the conditions presenting the risk must be sure or very likely 20 to cause serious illness and needless suffering and give rise 2.1 to sufficiently imminent dangers." 22 In Pavatt, the Tenth Circuit also noted that in Baze the 2.3 Supreme Court held that simply because an execution method may

result in pain, either by accident or as an inescapable

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objectively intolerable risk of harm that qualifies as cruel and unusual. Likewise, the Tenth Circuit noted that in Baze the Supreme Court held that a stay of execution may not be granted "unless the condemned prisoner establishes that the state's lethal injection protocol creates a demonstrated risk of severe pain and that the risk is substantial when compared to the known and available alternatives." That is at page 1339 of the Pavatt decision quoting Baze at page 61. On this point, it is noteworthy that in Pavatt the prisoner, Jeffrey Matthews, was asserting a drug-related risk, not a risk of maladministration.

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One of the things that is clear from the Baze decision, especially as that decision was applied by the Tenth Circuit in Pavatt, is that if the risk asserted by the prisoner is very speculative at all that speculative element will drain away the constitutional significance of the risk. A good example of this is the treatment of an Arizona prisoner's claims by the Arizona District Court, the Ninth Circuit, and the Supreme Court.

In Landrigan v. Brewer, 2010 West Law 4269559, from the District of Arizona on October 25th of 2010, the plaintiff asserted that there was an unconstitutional risk of harm flowing from the state's proposed use of drugs from a foreign source that was not approved by the FDA. The plaintiff asserted that the foreign supply of sodium thiopental might be

contaminated with toxins that could cause pain and could fail to properly anesthetize the plaintiff resulting in excruciating pain when the second and third drugs are administered.

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The district court agreed that the prisoner had raised significant issues about the efficacy of the non-FDA-approved sodium thiopental and granted a stay of execution. was affirmed by the Ninth Circuit at 625 F.3d 1144. The Supreme Court promptly vacated the stay in a one-paragraph opinion. Brewer v. Landrigan, 131 Supreme Court 445, from 2010. Of note here, the Supreme Court, quoting its decision in Baze, in part, said that "there is no evidence in the record to suggest that the drug obtained from a foreign source is unsafe. The district court granted the restraining order because it was left to speculate as to the risk of harm. But speculation cannot substitute for evidence that the use of the drug is sure or very likely to cause serious illness and needless suffering." Those are the words of the Supreme Court.

Thus, the Supreme Court's subsequent treatment of its decision in Baze makes it unmistakably clear that a speculative assertion of a risk of harm cannot substitute for a showing "that the use of the drug is sure or very likely to cause serious illness and needless suffering."

I'm also influenced by the Sixth Circuit's opinion in Cooey v. Strickland, 589 F.3d 210, a Sixth Circuit decision from 2009, in part because of the very thoughtful opinion

written for the Court by Circuit Judge Julia Smith Gibbons and in part because that case involving an Ohio execution arose against the backdrop of a serious mishap that had occurred in another execution in Ohio.

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Cooey v. Strickland was decided on December 7, 2009, one day before the plaintiff in that case, Kenneth Biros, was scheduled for execution. On September 15, 2009, slightly less than three months before the Sixth Circuit's decision in Cooey, the state of Ohio unsuccessfully attempted to execute Romell Broom. As explained by the Court of Appeals in Cooey, "The execution team was unable to find a vein on Broom's arm after repeated attempts over two hours. They attempted to insert the IV catheter into the crook of Broom's elbow, his wrists, over the knuckle of his first finger, and near his ankle. Twice the team managed to insert a catheter that was not secured properly and caused bleeding." That's 589 F.3d at page 224, note 3.

Citing the unsuccessful attempt to execute Biros, as well as other allegedly unconstitutional revisions to the Ohio protocol, Biros sought a stay of execution. He asserted that there was an undue risk of improper implementation of the Ohio protocol which would lead to severe pain, that Ohio employed untrained and insufficiently competent medical personnel, that there was a lack of supervision of the execution process by a licensed physician, and that there was a lack of a prescribed time limit within which to establish IV access, among other

complaints, as indicated on page 223.

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On the issue of the risk of maladministration, the unfortunate experience in the Broom execution figured heavily into the arguments advanced by Biros. As explained by the Fifth Circuit, "Biros relies heavily on Ohio's halted execution of Broom to distinguish his case from that of Baze," as stated on page 224.

The Sixth Circuit concluded that the unfortunate and very recent experience with the Broom execution, which, from the available information, could fairly be called a botched execution, did not take Biros' claim out of the realm of speculation. Citing Clemons v. Crawford, 585 F.3d 1119, an Eighth Circuit decision from 2009, the Sixth Circuit noted that the Eighth Circuit had "rejected the prisoner's claim that there was a substantial risk of pain due to incompetent personnel despite the fact that the Court had previously found that medical personnel administering the protocol, since removed, had been incompetent," as stated at page 225. The Sixth Circuit concluded that "for the same reasons, we cannot assume that the same misfortunes that befell Broom will befall Biros," as stated at page 225.

On the basis of the record before it, including the wholly unsuccessful attempt to execute Romell Broom, the Sixth Circuit held that "speculations or even proof of medical negligence in the past or in the future are not sufficient to render a

facially constitutionally sound protocol unconstitutional."

That is the Cooey decision at page 225.

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I conclude, as a matter of law, that the revised lethal injection protocol adopted by the Oklahoma Department of Corrections effective September 30, 2014, is facially constitutional when measured by the principles promulgated in Baze v. Rees, as further explained by our Court of Appeals in Pavatt v. Jones.

Citing the experience with the execution of Clayton

Lockett, these plaintiffs assert, in substance, that there is

not a constitutionally sufficient degree of assurance that the

revised protocol, even if it is constitutional on its face,

will be administered in a way which will avoid the infliction

of serious pain. This contention obviously requires the Court

to assess probabilities with respect to future events, some of

which are within the control of humans, albeit fallible humans,

and some of which are decidedly not within the control of

anyone.

I conclude on the basis of the evidence before me that plaintiffs have failed to establish that proceeding with the execution of these plaintiffs on the basis of the revised protocol presents a risk that is "sure or very likely to cause serious illness and needless suffering," amounting to "an objectively intolerable risk of harm," in the words of the Supreme Court at page 50 of the Baze decision.

In reaching this decision, I place considerable reliance, and I'm going to say again considerable reliance, on three aspects of the DOC's lethal injection protocol. The first is the requirement that both primary and backup IV access sites be established. The second is that confirmation of the viability of the IV sites is specifically required. The third is that the offender's level of consciousness must be monitored throughout the procedure.

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The predominant risk asserted by the plaintiffs in this case is that midazolam will not have its intended effect as the first drug in the series, thus leading to injection of vecuronium bromide or rocuronium bromide and potassium chloride into a sensate person either because of the asserted limitations of midazolam or because it is not effectively administered in a massive IV dose. The three safeguards to which I have referred will reduce the risk of injection of the second and third drugs into a sensate person to well below a level that would establish a right to relief under Baze v. Rees.

The conclusions I have reached establish that plaintiffs have failed to show a probability of success on the merits of Count 2. However, wholly apart from that, there is a separate reason for which plaintiffs have failed to establish a probability of success on the merits of Count 2.

In proposing that they be executed with a lethal dose of

1 sodium thiopental, as they assert at paragraph 31 of their 2 amended complaint, plaintiffs have failed to provide a 3 comparison with, in the words of the Supreme Court, a "known and available alternative." That is from page 61 of the Baze 4 5 On this issue, the burden of proof is immaterial. decision. 6 Because aside from plaintiffs' failure to demonstrate a known 7 and available alternative, the defendants have affirmatively shown that sodium thiopental and pentobarbital, the only 8 9 alternatives to which the plaintiffs have even alluded, are not available to the DOC. 10

In In Re Lombardi, 741 F.3d 888, an Eighth Circuit decision from earlier this year, the plaintiffs asserted that they were "not required to propose an alternative method of execution as an element of their Eighth Amendment claim."

That's at page 895. The Eighth Circuit responded, in my view correctly, that this was "a plain misreading of the Supreme Court's decision in Baze v. Rees and the Eighth Amendment," also at page 895.

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The Eighth Circuit, after quoting the passage in Baze, which requires comparison "to the known and available alternative," cited cases from the Fifth Circuit, Raby v. Livingston, 600 F.3d 552, with the relevant discussion at pages 560 and '61, a Fifth Circuit decision from 2010; and the Sixth Circuit, Cooey v. Strickland, which I have already discussed, for the proposition, with which I agree, that it is incumbent

upon a prisoner who challenges an execution protocol under Baze
to demonstrate that the risk created by the challenged protocol
is substantial when compared to the known and available
alternatives. Indeed, in the Raby decision, 600 F.3d at page
561, the Fifth Circuit described this as "the second step of
the Baze test."

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The only alternative short-acting barbiturate proposed by the plaintiffs as an alternative to midazolam is sodium thiopental, as alleged in paragraph 31 of the amended complaint. But sodium thiopental became unavailable in the United States long before these plaintiffs proffered it in this case as an alternative to midazolam. In Pavatt v. Jones, which I've already discussed, 627 F.3d at page 1338, the Tenth Circuit noted that "sodium thiopental is now effectively unobtainable anywhere in the United States, thus requiring Oklahoma and other death penalty states to revise their lethal injection protocols."

In Sepulvado v. Jindal, 729 F.3d 413, a Fifth Circuit decision from 2013, at page 416, the Fifth Circuit noted that sodium thiopental had been unavailable since 2010. In Chavez v. Florida, 742 F.3d 1267, at page 1274, an Eleventh Circuit decision from earlier this year, Chief Judge Carnes made the same observation in his concurring opinion in which he also noted that in 2013 the European Union threatened to limit the supply of propofol, which caused Missouri authorities to revise

Missouri's protocol. Judge Carnes concluded, I believe correctly, that "an alternative drug that its manufacturer or its distributor or the FDA will not allow to be used for lethal injection purposes is no drug at all for Baze purposes," as stated on page 1275.

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I conclude as a matter of law that the plaintiffs' reliance on sodium thiopental as an alternative to midazolam is altogether unavailing because there has been no showing that sodium thiopental is, in fact, an available alternative.

Plaintiffs have failed to establish a probability of success on the Eighth Amendment cruel and unusual punishment claim asserted in Count 2.

I will now turn to Count 4, the Eighth Amendment claim asserting unsound procedures and inadequate training.

COUNT 4 - EIGHTH AMENDMENT - UNSOUND PROCEDURES AND INADEQUATE TRAINING

With respect to execution procedures and training to prepare the execution team to competently perform an execution by lethal injection, the revised Oklahoma protocol is at least as protective of the prisoner's interests as the protocol which was before the Court in Baze.

The following aspects of the protocol which passed muster in Baze are noteworthy. The individuals inserting the IV catheters were only required to have at least one year of professional experience. The prisoner's state of consciousness

1 or unconsciousness is determined by the warden and the deputy 2 warden through visual inspection. The duty to watch for 3 problems with the IV catheters and the tubing fell to the 4 warden and the deputy warden. The list of professional 5 categories approved for IV insertion was identical to 6 Oklahoma's list, specifically a certified medical assistant, a 7 phlebotomist, an EMT, a paramedic, or a military corpsman. Kentucky protocol called for at least ten practice sessions per 8 The Kentucky protocol called for the IV Team to 9 year. 10 establish primary and backup lines and to prepare two sets of 11 lethal injection drugs. In Baze, the petitioners specifically faulted the Kentucky 12 13 protocol for lacking a systematic mechanism for monitoring 14 anesthetic depth, as indicated at page 58 of the Baze decision. 15 They maintained that the visual inspection performed by the 16 warden and deputy warden was an inadequate substitute for more 17 sophisticated procedures they proposed, such as the use of 18 various types of monitoring equipment, as discussed at page 59. 19

Moreover, the Supreme Court, citing the Tenth Circuit's decision in Hamilton v. Jones, 472 F.3d 814, with the relevant discussion at page 817, a Tenth Circuit decision from 2007, concluded that "the risks of failing to adopt additional monitoring procedures are thus even more remote and attenuated than the risks proposed by the alleged inadequacies of Kentucky's procedures designed to ensure the delivery of

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thiopental." That is at page 59.

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On this point, the plurality opinion rejected the dissent's argument that "rough and ready tests for checking consciousness, calling the inmate's name, brushing his eyelashes, or presenting him with strong noxious odors" was necessary in order to "materially decrease the risk of administering the second and third drugs before the sodium thiopental has taken effect," as discussed by the Court at page 60.

This led to the Supreme Court's conclusion that "an inmate cannot succeed on an Eighth Amendment claim simply by showing one more step the state could take as a fail-safe for other independently adequate measures. This approach would serve no meaningful purpose and would frustrate the state's legitimate interest in carrying out a sentence of death in a timely manner." That's at pages 60 and 61.

In Muhammad v. Crews, 2013 West Law 6844489, a decision from the Middle District of Florida about a year ago, late 2013, a decision which was affirmed at 739 F.3d 683 earlier this year, with certiorari denied, 134 Supreme Court 894, which was on January 7th of this year, the district court observed at star page 8 that "the Florida protocol requires that the execution team confirm that the inmate is unconscious after administration of the first drug, midazolam hydrochloride.

Thus, if done correctly, there is no substantial risk of harm

from administration of the second and third drugs."

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Although this is not a conclusion of law, an additional comment with respect to Warden Trammell may be appropriate here. My sense of the matter from the evidence, including listening very carefully to the testimony of Warden Trammell, is that the experience of the Lockett execution was in some ways repugnant to Warden Trammell. I am persuaded that Warden Trammell does not want a mishap like this to ever occur again, at least on her watch. Granting that Warden Trammell may, in some ways, be subject to criticism for playing what was arguably an overly passive role in the run-up to the Lockett execution, I quite easily find that her skills as an administrator have already manifested themselves in the training regimen that has been implemented since last September and will be very evident in the preparations for the upcoming executions.

Plaintiffs have not satisfied the Court that the lethal injection procedures and training regimen that are now in place present any substantial risk of serious harm within the meaning of the Court's holdings in Baze v. Rees. I accordingly conclude that plaintiffs have failed to establish a probability of success on the Eighth Amendment unsound procedures and inadequate training claim asserted in Count 4.

I now proceed to consider the claim asserted in Count 5 relating to notice and opportunity to be heard asserted under

1 | the Eighth and Fourteenth Amendments.

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COUNT 5 - NOTICE AND OPPORTUNITY TO BE HEARD - EIGHTH AND FOURTEENTH AMENDMENTS

The heart of this claim is plaintiffs' assertion that ten days is not constitutionally sufficient notice to the prisoner of the DOC's intentions with respect to the specific drug combination to be used. This proposition is essentially moot as to these plaintiffs because they have been given a minimum of several weeks' notice of the combination of drugs that will be administered. In any event, ten days' notice would be sufficient, even if not optimal.

This claim is foreclosed by the reasoning of the Fifth Circuit in Sepulvado v. Jindal, 729 F.3d 413, with the relevant discussion at pages 418 through 420, and the Eleventh Circuit in Wellons v. Commissioner, 754 F.3d 1260, with the relevant discussion at page 1267, another decision from earlier this year. I agree with the analysis of the courts in both of those cases and further elaboration is not necessary.

I will add, however, that the ten-day provision is not without a sound rationale. Inmates and others have succeeded in a number of instances in securing embargoes to cut off the supply of chemicals used in lethal injection. In some situations, eleventh-hour litigation has virtually become the norm. That eleventh-hour litigation is intended to forestall, after years of litigation on the merits as well as clemency

proceedings, the execution of a validly imposed sentence of death, a penalty which itself has repeatedly been held to be constitutional. Those who are charged with the responsibility to carry out a sentence of death by lethal injection need the ability to avail themselves of other options on relatively short notice.

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I accordingly conclude that plaintiffs have failed to establish a probability of success on the notice and opportunity to be heard claim asserted in Count 5.

I now turn to Count 7, the Eighth and Fourteenth Amendment claim asserting experimentation on human subjects.

COUNT 7 - EIGHTH AND FOURTEENTH AMENDMENTS - EXPERIMENTATION ON HUMAN SUBJECTS

In this count, plaintiffs complain that the use of untried drugs by way of untested procedures will cause them to experience severe pain, needless suffering, and a lingering death.

In the Baze decision, at page 62, the Supreme Court stated, and here I am leaving out an internal citation,
"Throughout our history, whenever a method of execution has been challenged in this Court as cruel and unusual, the Court has rejected the challenge. Our society has nonetheless steadily moved to more humane methods of carrying out capital punishment. The firing squad, hanging, the electric chair, and the gas chamber have each, in turn, given way to more humane

1 methods, culminating in today's consensus on lethal injection. 2 The broad framework of the Eighth Amendment has accommodated 3 this process toward more humane methods of execution and our 4 approval of a particular method in the past has not precluded legislatures from taking the steps they deem appropriate in 5 6 light of new developments to ensure humane capital punishment. 7 There is no reason to suppose that today's decision will be any different." That is from page 62 of the Baze decision. 8 9 Thus, as the Sixth Circuit pointed out in Cooey, to which 10 I've already referred more than once, 589 F.3d at page 229, "that the procedure has never before been used, does not itself 11 12 establish that the procedure is cruel and unusual. The Supreme Court has previously considered various modes of execution and 13 14 has yet to find one violative of the Eighth Amendment." 15 short, I conclude that the Eighth Amendment does not immunize 16 an individual from being the first person to be subjected to a 17 new method of execution. 18 Count 7 fails as a factual matter and as a matter of law. 19 As a factual matter, by plaintiffs' own count, execution with 20 midazolam as part of a three-drug protocol has been 21 accomplished 12 times. That's the plaintiffs' pleading at 22 Docket Entry Number 159, page 58. This is not a new method, at 2.3 least in the sense required for the Court to regard its use as human experimentation. 24 25 As a matter of law, the basic Baze test still controls.

These plaintiffs must establish that the state's lethal injection protocol creates a demonstrated risk of severe pain and that the risk is substantial when compared to the known and available alternatives. They have failed to do so.

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I accordingly conclude that plaintiffs have failed to establish a probability of success on the human experimentation claim asserted in Count 7.

I now turn to the Count 8 claim relating to a right of access to information, counsel, and the courts.

COUNT 8 - RIGHT OF ACCESS TO INFORMATION, COUNSEL, AND THE COURTS

This claim is based on the First and Fourteenth
Amendments. From plaintiffs' motion, Docket Entry Number 92 at
page 15, and their preliminary hearing brief, Docket Entry
Number 160 at page 7, it appears that plaintiffs assert a right
essentially to have counsel physically present as a legal
proctor of the IV insertion process. This conjures up an
untenable scene in which the prisoner's counsel is standing at
the gurney, cell phone in hand, ready to dictate the
information necessary to fill in the blanks on an emergency ex
parte motion for stay if he or she takes issue with any part of
the process as it unfolds.

The reality is that as execution by lethal injection is actually carried out, the prisoner's erstwhile right of access to the courts must, of necessity, give way to the execution

team's discharge of its duties as long as those who are carrying out the process are operating within the confines of a constitutionally sound lethal injection protocol. And I hasten to add that it would appear from plaintiffs' contention as to the very closeness of the scrutiny that they say is constitutionally required that protection of the identities of the execution team members would likely be impossible.

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On this claim, I agree with the reasoning of Judge Wake of the District of Arizona in Towery v. Brewer, 2012 Westlaw 592749, from the District of Arizona, February 23, 2012, at star page 18, a decision that was affirmed by the Ninth Circuit at 673 F.3d 650, and here I'm leaving out internal citations "Prisoners have a constitutional right of access and quotes. to the courts that is adequate, effective, and meaningful. However, this right quarantees no particular methodology but rather the conferral of a capability. The capability of bringing contemplated challenges to sentences or conditions of confinement before the courts. Consequently, an inmate who brings a Section 1983 claim based on his right of access to the courts must be able to show that the infringing act somehow defeated his ability to pursue a legal claim. That is, a prisoner must show he suffered an actual injury as a result of the defendant's actions." That's at pages 348 and 49. actual injury is actual prejudice with respect to contemplated or existing litigation such as the inability to meet a filing

deadline or to present a claim. The right of access does not create an abstract freestanding right but exists to vindicate other rights."

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No court has found a constitutional right for the prisoner to have counsel present to supervise the IV insertion process and I decline to be the first judge to so hold.

In Count 8, plaintiffs also assert a First Amendment right of access to information about their planned executions, as indicated in Docket Entry Number 92 at page 15, Docket Entry Number 160 at page 9, and Docket Entry Number 159 at pages 71 through 75.

I conclude that plaintiffs' reliance on the First

Amendment is misplaced. The interests that plaintiffs, as

prison inmates facing execution, would protect under this

heading are protected to the extent that they are protected at

all under the Due Process Clause of the Fourteenth Amendment

and perhaps arguably under the Eighth Amendment as made

applicable to the states through the Fourteenth Amendment.

Measured by these provisions, the plaintiffs' right of access

to information about their impending executions is adequately

protected by the revised protocol, as I have already discussed.

To the extent that plaintiffs seek to avail themselves of the public's right of access to executions in Oklahoma, I conclude that even if they had standing to tie their claims to the general public's right of access, their claim would fail

substantially for the reasons articulated last Friday by Judge Heaton in Oklahoma Observer v. Patton, Case Number Civil 14-0905, his order being Docket Entry Number 48 in that case.

I accordingly conclude that plaintiffs have failed to establish a probability of success on the right of access to information, counsel, and the courts as asserted in Count Number 8.

I now reach my conclusion.

CONCLUSION

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Plaintiffs have failed to establish any of the prerequisites to a grant of preliminary injunctive relief.

They have failed to establish a probability of success on the merits of any of the five claims they assert for preliminary injunction purposes, even under the relaxed standard articulated in Kikumura v. Hurley, 242 F.3d 950, with the relevant discussion at page 955.

Plaintiffs have failed to demonstrate that absent a preliminary injunction they would suffer any non-speculative irreparable harm.

As to the third factor, the balance of the equities does not tip in plaintiffs' favor. Plaintiffs have been successfully prosecuted, convicted, and sentenced to death in proceedings that have withstood decades of trials, direct review, and collateral review. The equities of the matter strongly favor bringing their cases at long last to a

conclusion by carrying out the penalty that the courts have determined to have been constitutionally imposed.

2.1

2.3

Finally, I conclude that entry of a preliminary injunction would not be in the public interest. It is well-settled that as the Supreme Court said in its unanimous decision in Nelson v. Campbell, 541 U.S. 637, with the relevant discussion at page 6844, the state has "a significant interest in meting out a sentence of death in a timely fashion."

And the Supreme Court also told us in Calderon v.

Thompson, 523 U.S. 538, a decision from 1998, at page 556 -and here again I'm omitting some citations and internal quotes.

"When lengthy federal proceedings have run their course and a
mandate denying relief has issued, finality acquires an added
moral dimension. Only with an assurance of real finality can
the state execute its moral judgment in a case. Only with real
finality can the victims of crime move forward knowing the
moral judgment will be carried out. To unsettle these
expectations is to inflict a profound injury to the powerful
and legitimate interest in punishing the guilty, an interest
shared by the state and the victims of crime alike."

The motion of Plaintiffs Charles Warner, Richard Glossip,
John Grant, and Benjamin Cole for preliminary injunction is
without merit. It is in all things denied. A brief written
order will be entered to memorialize this ruling.

I direct the parties to withdraw their exhibits. That is

1	routine for purposes of facilitating the parties getting the
2	exhibits to the Tenth Circuit for review. So I do direct the
3	parties to withdraw their exhibits. That is entirely separate
4	from any questions about ultimate public access to the
5	exhibits, which I have already addressed and which I hope the
6	Department of Corrections will itself address very quickly.
7	Court will be in recess.
8	(COURT ADJOURNED.)
9	CERTIFICATE OF OFFICIAL REPORTER
10	I, Tracy Washbourne, Federal Official Realtime Court
11	Reporter, in and for the United States District Court for the
12	Western District of Oklahoma, do hereby certify that pursuant
13	to Section 753, Title 28, United States Code that the foregoing
14	is a true and correct transcript of the stenographically
15	reported proceedings held in the above-entitled matter and that
16	the transcript page format is in conformance with the
17	regulations of the Judicial Conference of the United States.
18	Dated this 23rd day of December 2014.
19	
20	/S/ Tracy Washbourne
21	Tracy Washbourne, RDR, CRR
22	Federal Official Court Reporter
23	
24	
25	