

4/13/2004

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1 4/13/04 Judge Casey take 1 of national versus Ashcroft TKR*PB,
2 an an, an an an good morning.

3 THE COURT: I might just say at the outset that we
4 went over the portions of the transcript that I believe the
5 government intends to read this morning from deposition and I
6 would urge both sides to take a little more pains in reviewing
7 materials that you submit to the Court. You're all more
8 professional than what you submitted last night, at least I
9 hope so.

10 I am not going to bother to go into each and every
11 one, but suffice it to say, one is, includes a question which
12 was perfectly appropriate that was objected to by the plaintiff
13 in the transcript but then the government lawyer rephrased it,
14 all of which is absurd to submit to me because the question was
15 clearly appropriate in the first place in any event it was
16 rephrased and it was answered.

17 This is just a waste of time. You have the rulings on
18 the questions and let's get on with the transcript.

19 MR. LANE: Your Honor, if I can, Shawn Lane for the
20 government.

21 At this time what we would like to do is since we have
22 a live witness here, Dr. Kanwaljeet Anand, we would like to
23 present him as the government's next witness, if that's okay
24 with the Court.

25 THE COURT: I'm always happy to have someone who is

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1 breathe engine that box and is giving something live and fresh
2 and not off a transcript.

3 MR. LANE: I suspected as much, your Honor. Thank
4 you.

5 THE COURT: That's fine, go RAOEURT ahead.

6 MR. LANE: The government's next witness, we call
7 Dr. Kanwaljeet Anand to the stand.

8 THE DEPUTY CLERK: Doctor, please take the witness
9 stand and remain standing. Raise your right hand.

10
11
12 THE DEPUTY CLERK: State and spell your full name
13 slowly for the record.

14 THE WITNESS: My name is Can wall SKWRAOET, K A N W A
15 L J E E T, is Anand. A N A N D.

16 THE DEPUTY CLERK: Thank you, Doctor. Please be
17 seated.

18 THE COURT: Doctor, can you give me that, how do you
19 spell your last name again?

20 THE WITNESS: An in and, A N A N D.

21 THE COURT: Anand am I pronouncing it correct.

22 THE WITNESS: Yes, that's correct.

23 THE COURT: Mr. Lane, you may inquire.

24 MR. LANE: Thank you, your Honor.

25 DIRECT EXAMINATION

1
2 BY MR. LANE:
3 Q. Good morning, Dr. Anand.
4 A. Good morning.
5 Q. Could you tell Court what you do for a living?
6 A. I'm a pediatrician trained in the care ever critically ill
7 infants and children.
8 Q. And are you licensed to practice medicine?
9 A. Yes, in the state of Arkansas.
10 Q. Are you board certified in any particular medical
11 speciality?
12 A. I'm a diplomat of the American board of pediatrics and the
13 sub board of critical care medicine.
14 Q. And what is critical care medicine?
15 A. Critical care medicine is the subspecialty within
16 pediatrics that is focused on the care of critically ill
17 children, severely ill infants and children.
18 Q. And why are you testifying here today?
19 A. I was asked to provide an opinion on this case by the U.S.
20 Department of Justice based on my background and experience and
21 research.
22 Q. And on what topic?
23 A. My area of training and research has been on the topic of
24 development of the pain system during fetal and neo natal life.
25 Q. Doctor, before we get into your opinion on that issue, I

1 would like to ask you some questions about your background.
2 Where did you go to medical school?
3 A. I went to medical school in Mo hat ma Gandhi PH*E moral
4 medical college in a place called indoor, I N D O R E, in
5 India.
6 Q. What degree did you get from that college?
7 A. I get an MBBS, which stands for bachelor of medicine,
8 bachelor of surgery, and is equivalent to the MD degree in the
9 U.S.
10 Q. What did you do after medical school?
11 A. I did one year internship partly at PHA* hath are aa since
12 she been SKROU HAPT, .
13 A. S Y E S H W A N T R A O, Hospital, in inn door, and
14 partlyality hindu RA*U hospital in Del Hi.
15 Q. What did you do after your internship?
16 A. I did another year of postgraduate training in pediatrics
17 which was part of a three year residency program, but I just
18 completed one year.
19 Q. And why was that?
20 A. During that year I had applied for the Rhodes scholarship
21 from India and received that scholarship so I left my pediatric
22 residency and went to University of Oxford.
23 Q. What degree did you receive from Oxford?
24 A. From University of Oxford I received the D Phil degree
25 which is doctor of philosophy.

1 Q. While at Oxford, what did you study?
2 A. The D Phil was from the faculty of medicine in O*FPL
3 forward university and my work was primarily focused on
4 research on the hormonal and metabolic responses of newborn
5 babies undergoing surgery.
6 Q. When you use the term hormonal response, what do you mean?
7 A. Hormonal means the release of hormones or the release of
8 endocrine mediators as a result of the surgical operation that
9 is done.
10 Q. You also used the term metabolic response, what do you mean
11 by that?
12 A. Many of the hormones produce changes in metabolism, changes
13 in the way the body handles sugars and fats and amineo addieds
14 and things like that so the metabolic effects of those hormonal
15 responses is very important.
16 Q. What were the results of your research while you were at
17 Oxford?
18 A. To summarize a number of different studies, basically my
19 research showed initially that babies have a massive hormonal
20 and metabolic response to a surgical operation. This was
21 contrary to our expectations and that was the initial finding
22 from the research.
23 Q. When you say contrary to your expectations, could you
24 explain that for us?
25 A. Yes.

1 We had postulated that premature babies and even
2 full-term newborns would be unlikely to mount a significant
3 response to a surgical operation because they have imKHER
4 endocrine glands, these are the glands that release the
5 hormones and they have immature metabolic processes in their
6 liver or other body organs. So, we had thought that there W-B
7 a very dampened response from new born infants.
8 On the contrary, it was a huge response, three to five
9 times that of the adult patient undergoing similar types of
10 surgery.
11 Q. Did you reach any ultimate conclusions as a result of your
12 research while at Oxford?
13 A. Yes.
14 After these results came out we postulated that this
15 might be partly due to the basic physiology of the infant or
16 partly due to the management that was provided during the
17 surgical operation. And we did some clinical studies where we
18 gave potent anesthetics to babies and we found that their
19 stress response was markedly decreased as a result of giving
20 deep anesthesia.
21 At that time the common -- the commonly prevalent
22 medical practice was that newborn babies were operated upon
23 without the benefit of anesthesia, they were given muscle
24 relaxants and they were paralyzed and operations were done
25 under minimal or no anesthesia at that time. And we showed

1 that giving anesthesia can reduce their stress response and can
2 improve their clinical outcome.

3 Q. For your research and studies at Oxford, what was the age
4 of the premature infants that were studied?

5 A. The premature infants we studied were from 25 weeks of
6 gestation and higher.

7 Q. Did you do any clinical work while at Oxford?

8 A. Yes.

9 While I was at University of Oxford for three years I
10 was working in the neo natal ICU, the intensive care unit,
11 where premature and full-term babies are cared for.

12 Q. After your time at Oxford, what did you do, professionally?

13 A. I received my doctorate and then moved to Harvard medical
14 school and Boston Children's Hospital for post-doctoral
15 fellowship.

16 Q. What did your post-doctoral fellowship consist of?

17 A. I was in the department of anesthesia at that time and
18 continued this line of investigation because I was intrigued by
19 the fact that babies given adequate anesthesia have fewer
20 complications in the post-operative period the days after the
21 surgery.

22 So, I wanted to study a population of babies that
23 are critically ill and we studied babies that were undergoing
24 open heart surgery and we devised some nuances at the timing
25 techniques and found that by giving them deep anesthesia we

1 could decrease their stress hormone response but we could also
2 improve their survival and markedly lower the complications
3 after the operation.

4 Q. You used the term deep anesthesia, what do you mean by
5 that?

6 A. There are various different levels of anesthesia. You can
7 have a light anesthetic which barely suppresses the responses
8 to painful stimulation and SRUBG a deep anesthetic where there
9 is a much greater suppression of the physiological and
10 autonomic systems. These are systems within the body that help
11 to maintain the body dynamic balance called homeo state EUS.

12 So, a deep anesthetic allows the body to undergo major
13 types of surgery without having a huge stress response to that.

14 Q. While you were at Harvard in this post- doctorate research
15 fellowship, did you do any teaching?

16 A. Yes, I did.

17 I was in the department of anesthesia and I was
18 providing lecture OPBD neo natal anesthesia and analgesia.
19 Analgesia means pain relief in newborn infants as an instructor
20 in the department of anesthesia.

21 I was also running a bio statistics course for the
22 residents and fellows in the department of anesthesia.

23 Q. What did you do next, professionally?

24 A. After the completion of this post doc fellowship I applied
25 for residency training and was given a spot in the pediatric

1 residency training program at Boston Childrens' Hospital and
2 Harvard med.

3 Q. What did that pediatric residency training program consist
4 of?

5 A. It's a three-year training program which basingly allows
6 medical graduates to train in the area of pediatric care, so
7 it's different rotations, different disciplines of pediatrics
8 we get exposed to.

9 Q. While doing that pediatric residency training program did
10 you have any involvement in the issues of pain in premature
11 babies?

12 A. Yes, certainly.

13 It is during that time that my thoughts about what is
14 the real meaning of the stress response came to fruition and I
15 put forward the first synthesis of the published data regarding
16 the early development of the pain system, which was published
17 back in 1987.

18 Q. While at childrens' Hospital did you have any clinical
19 involvement in the treatment of pain?

20 A. I was a pediatric resident and I was certainly taking care
21 of patients in, as a trainee. So wherever I could make an
22 impact on the pain management of children I tried my best.

23 Q. After your residency at childrens' hospital, what did you
24 do next?

25 A. I then moved to Massachusetts Jen RAL Hospital, which is

1 still within the Harvard medical school system and did a
2 clinical training fellowship in neo natal and pediatric
3 intensive care units at mass general hospital.

4 Q. At the conclusion of that, where did you go?

5 A. After this two year clinical training fellowship I moved to
6 Emory University in Atlanta and I was appointed as an assistant
7 professor of pediatrics, anesthesiology and psychiatry at
8 Emory.

9 THE COURT: What year was that, Doctor?

10 THE WITNESS: That was in July of 1993.

11 Q. How long were you at Emory, Doctor?

12 A. I was at Emory for about four years, until September of
13 '97.

14 Q. And while at Emory, could you give the Court an idea of
15 what you did professionally?

16 A. Certainly.

17 I continued my research on pain in the newborn and
18 tried to develop some animal models because having convinced my
19 devil that there is the development of the pain system, I
20 wanted to investigate methods that we could safely employ
21 medicines to treat pain in early life.

22 and so developed some

23 animal experimental models in the newborn rat.

24 I was also working clinically as a pediatric
25 intensivist in Atlanta.

1 Q. When you say pediatric intensivist, what does that mean?

2 A. That means I was taking care of children who were admitted
3 to the intensive care unit at two different hospitals, one was
4 *EGen tell STO*PB, E G E L S T O N, childrens' hospital and the
5 other was Hughes spalding Childrens' hospital, both in Atlanta.

6 Q. In treating those patients, do us have to address the issue
7 of pain?

8 A. Certainly.

9 Critically ill infants and children require multiple
10 invasive procedures and monitoring lines and things like that
11 and we have to pay a lot of attention to their pain and
12 sedation requirements.

13 Q. After your approximately four years at Emory, what did you
14 do professionally?

15 A. I was then appointed as chief of critical care medicine and
16 the department of pediatrics at University of Arkansas for
17 medical sciences and so I moved to Little Rock and was working
18 at Arkansas childrens' hospital.

19 Q. Did you hold any other positions at Arkansas KH*EUPBZ'
20 hospital?

21 A. Yes.

22 I was heading the critical care division and I was
23 also the director for the fellowship training program in
24 pediatric critical care medicine.

25 Q. And how long did you hold those positions at Arkansas?

1 A. I held both of those administrative positions from
2 September '97 until July of 2003.

3 Q. And are you still at Arkansas?

4 A. Yes, I'm still at Arkansas.

5 Q. And ado you do there, currently?

6 A. I'm currently a professor of pediatrics, answers tease
7 jolly, pharmacology and neuro biology at University of Arkansas
8 for medical sciences. And I'm also pediatric intensivist at
9 Arkansas Childrens' hospital.

10 Q. And how long have you been doing those particular two jobs
11 at Arkansas?

12 A. Since September of '97.

13 Q. When you mentioned your position as a professor, you used
14 the term farm pharmacology, could you explain what that term
15 means?

16 A. Certainly.

17 Pharmacology is the study of drugs and what they do to
18 the living organism and how the living organism responds and
19 metabolizes those drugs.

20 Q. You also use the term neuro biology and what does that term
21 mean?

22 A. Neuro biology is the discipline for the study of the
23 peripheral and central nervous system within living organisms.

24 Q. When you say peripheral and central nervous system, can you
25 give a little more full explanation of what that is?

1 A. Sure.

2 I'm sorry to use this jargon but the peripheral
3 nervous system is simply the network of receptors and nerve
4 fibers that bring information from the periphery, from the body
5 organs from the surfaces, the skin and mucous membranes to the
6 central nervous system.

7 The central nervous systems consistss of the brain,
8 the brain stem and the spinal cord.

9 Q. In your current position, what do you teach?

10 A. I teach pediatric critical care to the residents and
11 fellows in our department and I also teach medical students
12 particularly the physiology and neuro biology of pain.

13 I also have some opportunities to teach anesthesia
14 residents and emergency medicine residents in the pain
15 management and those kinds of areas.

16 THE COURT: Doctor, what is the difference between
17 resident and a fellow?

18 THE WITNESS: A resident is someone who has completed
19 their medical school and is undergoing either a two or a
20 three-year postgraduate training in a major speciality like
21 pediatrics or anesthesiology.

22 And then after completion of the residency people can
23 go for a fellowship to get subspecialty training.

24 So, someone who is trained in pediatrics may then
25 further specialize in pediatric critical care or other

1 subspecialties of pediatrics.

2 THE COURT: Thank you.

3 Q. Doctor, at the University do you hold any particular
4 position in connection with neuro biology of pain?

5 A. Yes. I'm also the director of the pain neuro biology
6 laboratory which is located at Arkansas childrens' Hospital
7 research institute.

8 Q. And what does that do?

9 A. My lab is basically focused on the research looking at the
10 short-term and the long-term effects of pain during early life.

11 We also have ongoing studies on the effects of
12 different pain relieving medications and their impact on
13 lock-term behavior and so on.

14 That's the experimental part of the are search.

15 There are some clinical studies being done as well,
16 primarily looking at the effects of both short-term and
17 long-term of pain relieving medications using premature babies.

18 Q. You had mentioned some current research, what are you
19 currently involved in researching?

20 A. My clinical research mainly deals with the effects of
21 commonly used analgesic agents or pain relieving medications
22 such as morphine or TPEPB TA mill in premature babies. And we
23 are studying babies all the way from 23 weeks of gestation
24 onwards to 32, 33 weeks of gestation.

25 Q. You used the term analgesic, how does that relate to the

1 term anesthesia, is it the same or are there differences?

2 A. Well, analgesic simply means an intervention or a
3 medication used for relieving pain. Many of these medications
4 are also used in higher doses to produce a state of obtunded
5 consciousness in terms of anesthesia, and so many of the drugs
6 that are used for providing anesthesia in smaller doses can be
7 used as analgesics.

8 There is a considerable overlap, however there are
9 some drugs that are purely analgesics that cannot be used for
10 anesthesia and there are some drugs that are purely answer at
11 the timeicss so they cannot be used for analgesic.

12 So there is an overlap between the types of
13 medications used in these two areas.

14 Q. Doctor -RGS you had also mentioned that you currently care
15 for patients, could you describe what your current care is for
16 patients in your position?

17 A. Certainly.

18 I care for patients in the pediatric ICU at Arkansas
19 childrens' Hospital and my patients include the entire gamut of
20 severe illness during infancy and childbirth and adolescence.

21 So, all the severe infections, major trauma, post
22 operative care. All those areas are -- all those patients are
23 provided care in the pediatric ICU.

24 Q. What are the age ranges for the patients that you treat?

25 A. Our patients can be anywhere from tiny newborn infants to

1 adolescents up to 16 or 18 years of age. Sometimes we have
2 older patients as well. Pretty clearly if they have had care
3 provided in childrens' Hospital since their infancy for chronic
4 conditions.

5 But generally from -- there are even some premature
6 infants that we care for in the pediatric ICU but mostly
7 full-term or older infants and children.

8 Q. When you say premature, what are the age ranges,
9 gestational age ranges for those infants?

10 A. Generally the patients that are admitted to the pediatric
11 ICU are premature infants that have been discharged from the
12 neo natal ICU and may have gone home for a day or two and they
13 have some recurrent illness, they can come back to the
14 pediatric ICU.

15 So there are some of the more mature premature
16 children.

17 Q. Do you at PHER anesthesia to your patients?

18 A. During the case of providing critical carry do provide, on
19 occasion, anesthesia if one of my patients requires a surgical
20 operation of some sort.

21 I will routinely provide deep sedation to these
22 children under *R undergoing invasive procedures.

23 And I also work on what's called the sedation service
24 where we provide deep sedation or light anesthesia to children
25 all over the hospital wherever they're having procedures

1 performed on them.

2 Q. Personally what's your role in administering anesthesia to
3 these patients?

4 A. It comprises of evaluating these patients, particularly
5 when I am on the sedation service I will go and evaluate these
6 patients, take a full history and physical exam, talk to the
7 parents, and then formulate a plan of action of how we are
8 going to give sedation and how we are going to avoid some of
9 the complications that may be associated with that.

10 I get the parents' consent after full information and
11 discussion and then I provide the drugs, I monitor the patient
12 during the sedation or light anesthesia and then at the end of
13 the procedure I will stay with the patient until they are fully
14 awake or they're back to their baseline.

15 And then after that I will go back and do a post
16 anesthetic evaluation.

17 So these are primarily the pre anesthetic, the
18 anesthesia and the post anesthetic evaluations are what is the
19 recommended practice for anesthesiologist.

20 Q. Are you a board certified anesthesiologist?

21 A. No, I'm not.

22 Q. What's the difference between, if any, between what you do
23 and what a board certified anesthesiologist does?

24 A. A board certified anesthesiologist is someone who has gone
25 through three years of residency training in anesthesiologist

1 and they deal with a wider range of answers at the timeics that
2 are provided. As a pediatric intensivist during training in
3 critical care medicine we also learn to provide deep sedation
4 and anesthesia but in a very limited number of drugs in a
5 limited area.

6 That's the limit of -- there is some degree of overlap
7 in the two types of training, although anesthesiology is a
8 separate discipline and incorporates a lot more than what is
9 taught in critical care.

10 Q. Doctor, I would like to ask you generally about your
11 research work. Have you published any articles?

12 A. Yes. I have about 200 -- a little over 200 publications so
13 far.

14 Q. And how many of those are peer-reviewed articles?

15 A. They're close, between 55 and 60 peer-reviewed articles.

16 Q. And could you tell the Court what's meant by the term
17 peer-reviewed article?

18 A. These are articles that are submitted by researchers to
19 leading medical journals. The journal Ed I tore will send the
20 manuscript to a number of reviewers who are experts in that
21 area of research and will solicit their comments and criticism
22 and then as authors we have to respond to those comments and
23 criticisms and sometimes this process of back and forth takes,
24 makes a lengthy period of time, after which the article is
25 accepted or rejected.

1 And if it is accepted, then it's published.

2 Usually there is a lag time between completing the
3 research studies and the actual pub indication, there is
4 sometimes two or three years after the studies have been
5 completed.

6 Q. Doctor, in general, what kind of topics do you write about
7 in the articles that you author?

8 A. My research has mainly focused on pain, particularly its -P
9 indication in early life and its management.

10 I have also dealt with some of the ethical issues
11 around pain and focused on strategies for the management of
12 pain in newborns and small children.

13 Q. Have you written any chapters in textbooks, Doctor?

14 A. Yes, certainly.

15 Multiple chapters in textbooks of anesthesiology, of
16 pain management and other areas.

17 Q. Doctor, I think we are getting a little bit of feedback from
18 the microphones, if you can move back a tiny bit, that's
19 perfect.

20 Doctor, do you have any involvement in the national
21 institutes of mental health?

22 A. Yes. I'm part of the maternal fetal research network that
23 is fund by the NIH, in particular the NIM STKPWHRRBGSZ what is
24 that?

25 A. This is a collection of researchers from multiple different

1 countries and we are all focused on the area of fetal
2 physiology, fetal stress response, maternal stress and how that
3 has an impact on the fetus.

4 And we have regular meetings every six months and
5 sometimes more frequently than that to discuss this evolving
6 area of research.

7 Q. Do you perform any peer-review of scientific papers?

8 A. Yes, certainly.

9 I'm on the editorial board for the journal called
10 critical care medicine. Until last year I was also on the
11 editorial board for a journal called biology of the neo
12 TPHA*EUT.

13 But in addition to these two assignments, I do perform
14 peer review for a number of different journals in pediatrics,
15 in anesthesiology, in neuro biology, in behavioral neuro
16 science. All those areas of journals.

17 I routinely perform a lot of peer review.

18 Q. Doctor, have you testified as an expert witness before?

19 A. Yes.

20 As stated in my expert report I have testified in four
21 different cases in the past four years as an expert witness.

22 Q. In what areas have you testified?

23 A. I have been classified as an expert in the area of
24 pediatric critical care and in the pharmacology of answers at
25 the timic and analgesic drugs.

1 Q. Have you ever testified, putting aside this case, in any
2 case involving the regulation of abortion?
3 A. No, I have not.
4 MR. LANE: Your Honor, if I may approach?
5 THE COURT: You may.
6 Q. Doctor, you have been handed what's been marked Government
7 Exhibit C-5, can you tell the Court what that document is?
8 A. Yes. This is a printout of my curriculum vitae.
9 Q. And does an up-to-date and current copy of your CV?
10 A. This is a copy of the CV that was supplied in January of
11 this year and there have been a few additions since then.
12 Q. What are those additions?
13 A. There have been a couple of papers that have been published
14 since then or have been accepted for publication since January.
15 I have also received an award from the royal college
16 of pediatrics and child health in the UK.
17 MR. LANE: Your Honor, at this time we would like to
18 offer Government Exhibit C-5 into evidence.
19 THE COURT: Any objection.
20 MS. WIGMORE: No objection, your Honor.
21 THE COURT: It will be received.
22 (Government's Exhibit C-5 received in evidence
23 MR. LANE: At this time, your Honor, the government
24 would like to tender Dr. Anand in the fields of fetal O
25 knowledge fetal pain and pharmacology of anesthetic drugs.

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1 MS. WIGMORE: Your Honor, this is Ms. Wigmore, as you
2 know we have objected to the testimony of Dr. Anand on the
3 basis that it's irrelevant, not reliable, that it should not be
4 admissible under federal Rule of evidence 403 and we have also
5 challenged his his call physicians to he have as an expert
6 about anesthesiology.
7 We understand the Court has denied that motions but we
8 simply wanted it stated for the record.
9 THE COURT: It wasn't in the record already,
10 Ms. Wigmore.
11 MS. WIGMORE: It may be I want to make sure that we
12 preserve.
13 THE COURT: You knew full well it was in the record,
14 did you not?
15 MS. WIGMORE: In the trial it has not been mentioned
16 so I just wanted to make sure.
17 THE COURT: It is in the record of the case though.
18 MS. WIGMORE: Thank you before and we also object to
19 the characterization of his expertise as in fetal pain.
20 He has testified about expertise in pain and stress
21 responses but not studies about fetal pain in particular, so we
22 object to that.
23 THE COURT: Thank you your arrow there but your
24 objections are noted and overruled.
25 You may proceed, sir.

1 MR. LANE: Thank you, your Honor.
2 BY MR. LANE:
3 Q. Doctor, can you give us a definition of what you mean by
4 pain?
5 A. Pain is an unpleasant sensory and emotional experience
6 associated with actual or potential tissue damage or *R are
7 described in terms of tissue damage.
8 Q. Doctor once TKPWREP I apologize for saying this but don't
9 get too close to the microphone?
10 THE COURT: The microphone, Doctor has a tendency, we
11 don't hear it as much up here but it tends to distort your
12 voice a little bit, so if you keep about three or four fingers
13 between your mouth and the microphone I think you will find
14 that at least for those out in the courtroom the projection of
15 your voice will be a little clearer.
16 THE WITNESS: Thank you. Thank you, your Honor.
17 MR. LANE: Thank you, your Honor.
18 Q. Is there a particular source for your definition of pain?
19 A. Yes. This is the official definition from the
20 international association for the study of pain.
21 MR. LANE: Your Honor, if I may approach?
22 THE COURT: You may.
23 Q. Doctor, I have handed you a binder that has a number of
24 exhibits inT most of which we won't get into here today, but I
25 would like to refer you to Government Exhibit P-7 in that

1 binder, if you could.
2 A. I can see it.
3 Q. And can you tell me what that document is?
4 A. This is a printout from the international association for
5 the study of pain which is titled IASP pain terminology.
6 Q. What is the international association for the study of
7 pain?
8 A. This is a highly respected world body with more than 7,000
9 members, all of whom are focused on the study of pain. It's a
10 multi disciplinary organization so there are pediatricians and
11 anesthesiologist and neuro PWHROPBLG KWREUFTs, fizz KWROL
12 gists, etc., different scientistss all focused on the study of
13 pain.
14 Q. Do you consider that to be an authoritative body?
15 A. Certainly do.
16 Q. RUZ a member of that body?
17 A. Yes, I am a member of IASP.
18 Q. How do you --
19 THE COURT: Are there any obstetricians who are
20 members?
21 THE WITNESS: Yes, there are. There are several
22 obstetricians who are members of the IASP as well.
23 THE COURT: Go ahead. I'm sorry. HRRPBLGTS
24 certainly, your Honor.
25 Q. Doctor, how did you become a member of that organization?

1 A. I applied for membership and I had to receive the
2 nomination of two other members that are in good standing at
3 the IASP.
4 And then my membership application was reviewed by the
5 committee on membership and they made their own inquiries and
6 after a few months my membership was confirmed.
7 Q. Doctor, looking at Exhibit P-7, I would like to refer you
8 to the second page of that document about halfway down the page
9 under the heading pain terms, there is the word pain listed in
10 some text after that.
11 What is that text that comes on to page 2?
12 A. That is the definition of pain that I had referred to a
13 minute ago from the IASP.
14 And if I can read that text, it says, pain is an
15 unpleasant sensory and emotional experience associated with
16 actual or potential tissue damage or described in terms of such
17 damage.
18 Q. By the term sensory experience, can you explain what's
19 meant by that?
20 A. Yes.
21 Pain is a bodily sensation. It's a sensation that is
22 transduced through receptors carried by nerve fibers and
23 is processed in the brain.
24 Q. This definition also refers to it as an emotional
25 experience; what's meant by that?

1 A. Pain is an emotional experience in so much that it is
2 characterized as unpleasant and noxious in character for the
3 most part. And so, it generates an affective or emotional
4 response for the person who is undergoing pain.
5 Q. Doctor, below the definition you just read a few seconds
6 ago there is a note that says the inability to communicate
7 verbally does not negate the possibility that an individual is
8 experiencing pain and is in need of appropriate pain-relief
9 leaving treatment.
10 Do you agree with that statement?
11 A. I certainly do.
12 I had a significant role to play in that note being
13 added to the definition.
14 Q. Could you explain your role in adding that note to the
15 definition?
16 A. Yes.
17 Back in the late '80s and early 90s when I was
18 trying to advocate for greater research on pain and analgesia
19 during early life I was faced with this problem that people
20 felt what babies are experiencing are superficial reflexes and is
21 not really pain or subjective suffering of any sort and that
22 was because the definition of pain was described in terms of
23 self-report, in terms of linguistic ability. And apart from
24 that self-report sorts of evidence were not admitted into the
25 definition of pain.

1 So, I wrote an editorial which was published in
2 October of 1996 in the journal -- the official journal of the
3 IASP and that was titled new per speck TEUFBZ on the definition
4 of pain where we challenged the definition of pain and we said
5 that it excludes those who do not have linguistic ability paly
6 including premature and full-term newborn infants or young
7 children or people with dementia or people with other
8 neurological problems whereby they're not able to express their
9 pain and convey their pain as -- that they are suffering.

10 Q. When did the society add this note to the definition of
11 pain?

12 A. It was almost five years later in May of 2001 that this
13 note was added.

14 No those five years we made multiple presentations at
15 world Congresses of pain, at meetings of the American pain
16 society and all of these were -- all of these arguments were
17 reviewed by the committee on tax on O my and they finally added
18 that note in 2001.

19 Q. You just referred to the committee on tax on O my, can you
20 explain what that is?

21 A. This is the committee which is part of the IASP which
22 basically helps define all the terms associated with pain.

23 Q. Looking again at the second page of this exhibit, below the
24 note we were just discussing there was another note that
25 mentions that pain is always subjective. Can you explain what

1 that means?

2 A. Certainly.

3 What this means is that pain, like any other
4 experience, has a special importance and significance for the
5 person who is undergoing that experience.

6 So it's a subjective experience pertaining to the
7 subject who is experiencing it.

8 THE COURT: In other words you can't see it, can you,
9 Doctor?

10 THE WITNESS: No, you can't see it.

11 THE COURT: The person who is not experiencing it
12 can't see it.

13 THE WITNESS: Can't say it.

14 THE COURT: See it.

15 THE WITNESS: That's right.

16 THE COURT: With thinks highs.

17 THE WITNESS: Absolutely.

18 THE COURT: I couldn't see it any, but.

19 Others who have vision cannot see pain, isn't that
20 correct?

21 THE WITNESS: That is correct.

22 THE COURT: It's only the person experiencing the pain
23 who can feel it, correct?

24 THE WITNESS: That is absolutely right; and that's why
25 this definition was so important because up until the

1 definition was provided by the IASP, pain was always linked to
2 a proximate stimulus. So unless there is a stimulus people
3 said there cannot be pain where there are many types of pain
4 where you don't need a stimulus, a headache, for example.
5 And so, this definition brought to the forefront the
6 fact that if someone is saying that they are in pain then the
7 definition provides for the fact that they are in pain and that
8 has to be accepted.
9 Q. Doctor, in your opinion can pain be experienced without
10 tissue damage?
11 A. Yes, it can.
12 There are many types of pain that are not associated
13 with tissue damage but they may be described in terms of tissue
14 damages.
15 For example, the pain of angina may not have any
16 tissue damage but there is, you know, a tearing or a pressure
17 kind of pain that people feel during angina.
18 There are other types of pain that can be perceived
19 without tissue damage.
20 Q. If there is no reaction to pain does that mean that a
21 person does not experience pain?
22 A. Well -- and that's where the art of pain medicine comes in,
23 is patients can either magnify their pain or they can suppress
24 their pain response.
25 So, stoic people can completely hide the fact that

1 they may be in pain.
2 Also, you can have the perception of pain but not have
3 a reaction to it under different clinical situations.
4 Q. Doctor, I would like to turn to your opinion in this case,
5 do you have an opinion as to whether a fetus can feel pain?
6 A. Yes. Fetuses that are beyond 20 weeks of gestation can
7 feel pain.
8 Q. And what is the basis for your opinion?
9 A. My opinion is based on multiple lines of evidence looking
10 at the anatomical development of the pain system, the
11 functional correlates of the developing fetus, the
12 physiological responses of fetuses and their behavior.
13 All of these areas contribute to my opinion.
14 Q. Is your opinion to a reasonable degree of medical
15 certainty?
16 A. Yes, it is.
17 Q. Doctor, I would like to take, before we discuss your
18 opinion on fetal pain, I want to ask you, generally about the
19 main anatomical structures responsible in humans for the
20 perception of processing of pain.
21 First, could you define what you mean by anatomical
22 structure?
23 A. These are the receptors and the nerve fibers and the areas
24 in the brain which form the anatomy of the network associated
25 with the pain system.

1 Q. Can you briefly describe the primary an anatomical
2 structures that are required for human being to experience
3 pain?

4 A. Certainly.

5 The pain is initially transTKAOUSed through
6 specialized receptors that may be located on the skin or mucous
7 membranes or may be located on the internal organs or deep in
8 muscle and those receptors get activate and they send their
9 impulses along nerve fibers to the spinal cord.

10 The area within the spinal cord that is associated
11 with processing of pain is called the dorsal horn, it is the
12 back part of the spinal cord, it is a collection of nerve cells
13 within the spinal cord.

14 From the dorsal horn the pain impulse goes up to the
15 supra spinal or centers located above the spinal cord, mainly
16 in the brain stem and the that will PHUS and then finally to
17 the cortex and other sub cortical areas. You.

18 Q. You used a few terms there doctor and I would like to go
19 through a few of them.

20 Could you describe what the dorsal horn is?

21 A. The dorsal horn is basically a collection of nerve cells
22 that is contained within the spinal cord in the posterior, the
23 back part of the spinal cord or the door TAL SART of the spinal
24 cord.

25 (continued on next page) TWM 4/13/04 NAF v. Ashcroft, take 2, can
Anand on

1 direct by Mr. Lane<]

2 Q. What is the thalmus?

3 A. The thalmus is a collection of nerve cells which is the
4 area of pain perception is first registered in the brain. That
5 is located below the cerebral cortex, the gray matter on the
6 surface of the brain.

7 Q. Doctor, you segued to my next question. Can you explain
8 what the cerebral cortex is.

9 A. Yes. The cerebral cortex is the thinking, planning part of
10 the brain where all of these sensory stimuli and their mother
11 responses are coordinated from. So these are about two billion
12 nerve cells that are located on the surface of the brain, and
13 that is called the cerebral could be text.

14 Q. Are there any other apanatomical structures that are
15 involved in the proprocessing of pain?

16 A. Yes. From the spinal cord up to the thalmus, the pain
17 fibers have connections with some collections of nerve cells
18 called nuclei in the brain stem, and these help to coordinate
19 the changes in heart rate or respiratory rate or other things,
20 bodily functions, as a response to pain. And the thalmus is
21 where the final processing is distributed to multiple areas of
22 the brain. These may include the limb PWEUBG steam or the a
23 mig today la where emotional processing occurs, it may include
24 the hypothalmus where a stress response is incorporated, or it
25 may include they po campus which helps formulate the long-term

1 memories associated with pain. So these are the subcortical
2 structures. And then the cortex.
3 Q. Doctor, I would like to go through each one of these main
4 an Tom K-L structures and ask you when they develop in the
5 fetus. What about the receptors for pain?
6 A. The fetus during development has sensory receptors that
7 appear in the area around the mouth at about 7 weeks of
8 gestation. They spread to other parts of the body, covering
9 almost the entire skin surface apart from the top part and the
10 back part of the head by 14 weeks. And by 20 weeks of
11 gestation, all of the skin surfaces and mucus membranes have
12 sensory receptors manifested in them.
13 Q. You had mentioned sensory fibers. When do those develop in
14 the fetus?
15 A. The nerve fibers that connect these receptors with the
16 final cord are developing. You see, the way the reSERPTS
17 develop is that from the final cord these \notify\nerve out
18 growths go towards various parts of the body. They grow into
19 the limbs that are forming and finally reach the target,
20 whether that is the skin or the mucus membrane. Once the nerve
21 fibers have reached their target, that is when the receptors
22 are expressed.
23 So the fact that there are receptors proves the fact
24 that those receptors are connected with the nerve fibers.
25 Q. When does this connection between the fibers and the

1 receptors occur in the fetus?
2 A. As the receptors appear, they are simply the nerve EPBGTSZDZ
3 that have reached their target organ, like the skin, for
4 example.
5 Q. Are these sen ris \notifies\nerves connected to the skin
6 receptors by 20 weeks?
7 A. Yes, certainly they are.
8 Q. Are notify fibers covered with any particular covering in
9 an adult human?
10 A. Yes. In the mature nervous system these nerve fibers are
11 my linated, basically, that means they are covered with a my
12 lin sheath. What this does, it increases the speed of
13 conduction along the nerve fibers. Previously it was thought
14 that babies don't feel pain because their nerve fibers are not
15 mile TPHAEUTd. However, it is now known thatl 80 percent of
16 noxious information is transmitted even in the adult by fibers
17 that are unmile TPHAEUTd. These are called the C fibers, C as
18 in cat, C fibers.
19 Q. Doctor, could you explain when the dorsal horn of the
20 spinal cord develops in the fetus.
21 A. Certainly. The dorsal horn of the final cord starts
22 developing at the end of the first trimester, so about 12 to 13
23 weeks of gestation, and rapidly matures with the
24 differentiation of the nerve cells KWRAGT area with the
25 expression of different chemicals that are used for

1 transmitting impulses in the dorsal horn. By 20 weeks of
2 gestation it is a fairly mature structure.
3 Q. At what gestational week is the thalamus developed in the
4 fetus?
5 A. The thalamus actually starts developing at about 6 to 8
6 weeks of gestation. Then these cells differentiate, and by 16
7 to 18 weeks of gestation the thalamus is a fairly mature
8 structure. Certainly by 20 weeks of gestation the thalamus is
9 well connected both from the spinal fibers from below as well
10 as fibers connecting the thalamus with higher centers.
11 Q. You had mentioned the cerebral cortex. Can you tell us
12 when that develops sufficient to process and perceive pain.
13 A. The cerebral cortex also starts developing, all of these
14 changes in the developing brain occur in parallel. So the core
15 text starts developing at about 8 weeks of gestation. As cells
16 migrate out to the our attention mantle of the surface of the
17 brain, those numbers increase. Those cells are directed to a
18 very precise anatomic location, and even within that location
19 they are placed in particular layers within the cerebral
20 cortex.
21 By 20 weeks of gestation, most of the migration of
22 cells has occurred, they have already formed connections within
23 the cortex, and they have formed connections with incoming
24 fibers from the thalamus.
25 Q. Doctor, are the anatomical structures needed to perceive

1 and process pain present and functional in a fetus by 20 weeks'
2 gestation?
3 A. Yes, they are.
4 Q. Is there anything missing in the system that would perceive
5 and process pain at that gestational age?
6 A. This whole system is undergoing maturational changes at
7 different levels. That maturation proceeds during maturation
8 and even after birth and even during childhood and up through
9 adolescence. That is a pattern called brain PHRAS isty,
10 whereby depending on the precise experiences t pain system
11 changes its structure and function. S- S- but by 20 weeks of
12 gestation, all the components of the pain system are functional
13 and they are connected up together, they have come on line in
14 order to perceive pain.
15 Q. Doctor, you had mentioned anatomical indicators as one of
16 the bases for your opinion, but you had also mentioned
17 functional indicators. Can you explain what you mean by
18 functional indicators.
19 A. Certainly. Starting with the receptors in the skin and
20 mucus membranes, those receptors, when they first appear, they
21 are fully mature in their properties. They have the same
22 threshold of their membrane. They have the same firing
23 frequencies, etc., very similar properties to those of
24 receptors in the adult.
25 The nerve conduction occurs along thinly mile TPHAEUtd

1 and unmyelinated nerve fibers. That is also functional and
2 mature. There is processing within the dorsal horn. There is
3 processing at the thalamus and at the cortical level, and all of
4 those components are functional, as can be seen from nerve
5 conduction studies or from the electrical changes following
6 the stimulation of the sensory testimony. There is something
7 called the somatosensory evoked potential, which shows that
8 there is an electrical impulse that reaches up to the cerebral
9 cortex and you can actually examine the cortical component of
10 an evoked potential, a potential that is evoked by sensory
11 stimulation.

12 Q. Doctor, as a functional matter, does the fetus respond to
13 any other stimuli other than pain?

14 A. Certainly. There is a huge amount of literature showing
15 that the fetus responds to sound and is able to identify and
16 habituate or dishabituate to sound. The fetus is responding to
17 light. The fetus can respond to taste. For example, if you
18 inject a sweet substance, sweet-tasting substance like saccharin,
19 for example, into the amniotic fluid, the swallowing
20 activity of the fetus increases and they respond very actively
21 to the sweet taste within the amniotic fluid. If you inject
22 something that has a bitter taste, then their swallowing
23 activity stops. So those are other modalities that show that
24 the central nervous system is functioning.

25 The other ways in which the developing brain has been

1 studied is by recording fetal EEG, which is
2 electroencephalography. The electrical patterns from the brain.
3 Those electrical signals start up at about 19 to 20 weeks of
4 gestation. That is another area of evidence for the functioning
5 brain in the fetus.

6 Q. Doctor, you had mentioned that the fetus responded to
7 sound. Can you give us an example.

8 A. Certainly. There are numerous studies that have looked at
9 this. I recall an incident where a colleague of mine was
10 pregnant at about 22 weeks of gestation, and she said that her
11 husband was trying to put up a swing set for their older child
12 and dropped one of the pipes onto their driveway, which made a
13 loud clanging noise. She said at that time the fetus had a
14 typical startled response. You see, when new-born infants
15 after birth are exposed to a lot of sound, they will throw out
16 their arms and legs and will have what is called the Moro
17 reflex. She said at 22 weeks of gestation, this fetus had a
18 complete Moro reflex. This is a pediatrician.

19 So there are other studies that have looked at this,
20 whereby people have studied fetal behavior in response to a car
21 horn, for example, within a few feet from a pregnant mother.
22 Those kinds of studies have looked at fetal responses to sound
23 and light and so on. [Moro reflex].

24 MS. WIGMORE: Your Honor, I object and move to strike
25 the doctor's testimony about the conversation with the pregnant woman

1 and being hearsay and lacking personal knowledge.

2 THE COURT: I will grant the motion. S-

3 Q. Doctor, can you tell us about what effect, if any, fetal
4 mother behavior has on your opinion about a fetus's ability to
5 feel pain?

6 A. Certainly. That is another correlate of the functioning,
7 developing brain. The fetus has various types of movements in
8 utero which are initially quite reflexive and Curran domly.
9 But as the fetus develops more and more, these movements become
10 coordinated and have some degree of -- have a much greater
11 degree of coordination, for example. There is an ultrasound
12 study that was done whereby a fetus in its random movements
13 within the amniotic fluid had grasped its umbilical cord, and
14 as a result of that obviously there may have been a decrease in
15 the blood flow to the fetus and the fetus was seen to push away
16 the umbilical cord. So there is some degree of evolving
17 behavior in the fetus that has been studied on ultrasound.

18 There are also studies showing states of wakefulness
19 and sleep in the fetus developing, and those correlate with the
20 patterns of EEG that have been monitored at the same time.

21 Q. Doctor, I would like to turn to the physiological
22 indicators for pain that you mentioned earlier. What did you
23 mean by that?

24 A. There are various responses that occur within the
25 physiology of the body. When a fetus is subjected to a painful

1 stimulus SP-P.

2 Q. What kind of responses are you talking about?

3 A. Those are mainly responses that have been studied in the
4 form of the release of hormones or changes in the circulation
5 within the fetal body.

6 Q. Based on your review of these studies, when do these
7 responses first appear?

8 A. The earliest fetuses that seem to mount a hormonal or a
9 cardiovascular response to painful procedures has been
10 documented at about 16 weeks. But by 20 weeks of gestation
11 those responses become very robust and reproducible.

12 Q. Doctor, I would like to ask you if there is any particular
13 studies performed on the issue of stress hormones.

14 A. Certainly. There is a number of studies that have been
15 done looking at different stress hormones, like bet dean more
16 feign or cat a col means. These are adrenaline and nor
17 adrenaline. These are stress hormones. S- they have been
18 measured in the fetus that is being exposed in an invasive
19 procedure. Cortisol is another hormone that has been measured.
20 There is a series of studies published.

21 Q. Is there any particular study that is particularly relevant
22 to your conclusion about stress hormones?

23 A. There is actually one paper which I feel has had a major
24 impact on my opinion, and that was published by Norman Fisk and
25 his colleagues from imperial College, London, in 2001.

1 Q. Doctor, I would like you to turn to Government Exhibit G8,
2 which is in the binder before you.
3 A. Yes, I have it.
4 Q. What is that document?
5 A. This is a printout of the paper by sir Nicholas Fisk
6 ~~***[STRIKE]***~~ -- sorry, Nicholas Fisk and his colleagues,
7 titled the effect of direct opioid analgesia on hormonal
8 and haemodynamic host response to intrauterine needling.
9 Q. Is this the article you referred to?
10 A. Yes. This is one of the more recent studies from this
11 group.
12 Q. Could you explain what this study looked at.
13 A. Certainly. This is a highly specialized team of
14 researchers. They have studied fetal responses for more than
15 10 years. In this particular study they took fetuses from 20
16 weeks to about 35 weeks of gestation who required a blood
17 transfusion in utero. There are two ways of providing the
18 blood transfusion. Either one can insert a needle through the
19 fetus's abdomen into their peritoneal cavity and then pass
20 through the capsule of the liver into a vein that is located
21 inside the liver, it is called the intrahepatic vein. That is
22 one way of accessing the fetal circulation in order to give a
23 blood transfusion.
24 What they found was that fetuses undergoing this
25 approach of blood transfusion had a very robust hormonal

1 response to the pain associated with the needling. Also, the
2 magnitude of the hormonal response was directly proportional to
3 the duration for which the needling occurred.
4 Another way of providing the blood transfusion is to
5 insert a needle into the umbilical vein, where it comes off of
6 the placenta you can insert a needle over there and give the
7 blood transfusion. The umbilical cord is not innervated, so
8 there is no major pain that can occur as a result of that. And
9 the fetuses did not have a hormonal response.
10 What these researchers did was in fetuses that were
11 undergoing PHREULT blood transfusions, they had the same fetus
12 undergo a transfusion in the intrahepatic vein without any
13 analgesia and it randomly gave for the other transfusion a dose
14 of fentanyl, which is an anesthetic drug, prior to the blood
15 transfusion. They found that the hormonal response was blocked
16 by giving proper analgesia directly to the fetus.
17 So basically this study was a rather complex study,
18 but what it showed is if you have a painful way of giving a
19 blood transfusion, it produces a big hormonal response and what
20 is called a hemodynamic response, a change of circulation. If
21 you have a nonpainful way of giving the blood transfusion,
22 there is no response. If you use the painful way of giving the
23 blood transfusion and give a pain-relieving agent prior to the
24 transfusion, then you can prevent this hormonal and hemody-
25 namic stress response.

1 Q. Doctor, when you said a pain-relieving agent delivered to
2 the fetus, how was that delivered for purposes of this study?
3 A. In this study they gave a dose of fentanyl, 12.5 micrograms
4 per kilogram, to the fetus directly into the circulation, the
5 blood circulation of the fetus.
6 Q. Doctor, can you tell us what studies have been performed on
7 circumstance latory response changes in fetuses in response to
8 pain.
9 A. A number of studies have looked at circulation in the fetus
10 and how it changes during painful stimulation, mainly noting an
11 increase in blood flow to the brain as soon as 70 seconds after
12 a painful stimulus occurs.
13 Q. How do we know that pain is causing these responses and not
14 some other stimulus or other factors?
15 A. That is a good question. These are nonspecific responses
16 and would occur in relation to other types of stimulation. For
17 example, blood loss can also lead to a cortisol response or a
18 cata col mean response. However, these responses were
19 documented immediately following a painful stimulus, and they
20 had good controls. They had a control situation whether the
21 lack of the painful stimulus produced no response, a painful
22 stimulus produced a robust response. Based on that temporal
23 sequence, we can relate those responses to the painful
24 stimulation that had occurred.
25 Q. Doctor, I would like to ask you some questions about your

1 opinion as it pertains to the scientific method. Can you tell
2 the Court what is meant by the term "scientific inference.?"
3 A. Certainly. In science, generally we, being a skeptical
4 scientist, we can never prove anything. But we can disprove
5 something. So what we start out with is a hypothesis, and we
6 try to disprove that hypothesis. For example, in this
7 situation the researchers started out saying that the fetus
8 does not respond to pain, but their data showed that there is a
9 significant response, so the null hypothesis was rejected.
10 This way of deductive thinking is called scientific
11 inference, where you look at a pattern of data within a certain
12 experimental paradigm and then figure out what that data means.
13 That is called scientific inference.
14 Q. Is inference, scientific inference, a widely accepted tool
15 in the scientific community for drawing conclusions?
16 A. Certainly.
17 Q. Is it based on scientific evidence?
18 A. It is.
19 Q. Can you explain to the Court how your opinion -- in what
20 ways it is based upon inference.
21 A. Basically, from these multiple lines of evidence that show
22 the fetus actively responding to different types of sensory
23 stimuli, I have inferred that the pain system is developed
24 enough to be functional and that the fetus is experiencing pain
25 as a result of invasive procedures.

1 Q. Doctor, can you explain the term "extrapolation" as it
2 pertains to scientific research.
3 A. Yes. In the scientific method we will try to extrapolate
4 from one experimental paradigm to another experimental
5 paradigm. For example, suppose there is a mechanism of pain
6 that works in adults. If this truly is a mechanism, then there
7 should be other evidence that we can obtain from other age
8 groups or other patient populations by extrapolating that
9 mechanism. So extrapolation is a well-developed tool in the
10 scientific method.
11 Q. Can you explain how your opinion -- how you used
12 extrapolation in your opinion in this case.
13 A. Until the area of fetal pain, I have used extrapolation in
14 two different ways. One is to extrapolate from the studies
15 that were done in animals or experimental animals. Keeping in
16 view that the neurological maturity or the state of mat RAEUGS
17 of the brain and nervous system is corresponding with that of
18 the human fetus, some degree of extrapolation was used in that
19 sense.
20 Also, extrapolation was used simply on the fact that
21 if there is a pattern of fetal response, hormonal or metabolic
22 or circumstance latory response, and if that same pattern of
23 response is seen in older children or adults, who then report
24 that they are experiencing pain, then you can extrapolate to
25 the fact that the fetus would be experiencing pain.

1 Q. Doctor, let me ask you, are there differences between
2 fetuses and an infant born at full term?
3 A. There are certainly huge differences between a fetus at
4 different stages of maturity and the full-term infant, yes.
5 Q. What effect, if any, does that have on your opinion in this
6 case about a fetus's ability to feel pain?
7 A. What we have noted from these multiple lines of evidence is
8 that the pain system has a very low threshold, meaning that the
9 fetus has a much greater sensitivity to pain during the early
10 development of the pain system, and later on that threshold
11 rises or the sensitivity decreases to pain. This is seen
12 throughout development. So in premature neoTPHAEUTS who is 23,
13 24 weeks of gestation, they have a much lower threshold for
14 pain compared to a full-term infant. A full-term infant has a
15 lower threshold to pain as compared to, say, a 1 or 2-year-old
16 child. And during childhood as well there is a progressive
17 increase in the threshold of pain.
18 So my opinion is that between 20 and 30 weeks of
19 gestation there is the greatest sent activity to pain.
20 Q. Doctor, can you explain the scientific reasons why that is
21 so.
22 A. There are many reasons to explain this increased
23 sensitivity to pain. Firstly, there is the early development
24 of the receptors and the density of these receptors is much
25 greater in the fetal skin as compared to an older child or

1 adult. These receptors have connections with the spinal cord
2 whereby the excitatory mechanisms are fully developed by the
3 inhibitory mechanisms or mechanisms that may modulate the
4 incoming no, Judge should say information develop later on,
5 after about 32 to 34 weeks of gestation. So it is that early
6 period there is a greatest sensitivity to pain.

7 Q. You mentioned inhibitory mechanisms. What do you mean by
8 that?

9 A. Inhibitory mechanisms are basically the mechanisms within
10 the spinal cord that help to modulate incoming painful stimuli
11 and also descending inhibitory fibers. These are fibers from
12 the brain stem, from the thalamus, that grow down the spinal
13 cord and make connections with the dorsal horn. The primary
14 functions of these fibers is to block incoming painful
15 information.

16 If I may give an example. Suppose you are shot in the
17 leg and you are trying to escape. The brain will be able to
18 switch off the painful stimuli coming from your leg to allow
19 you to escape from a dangerous situation. But once you are in
20 a safe place, then that pain will return because the block is
21 removed. So there is the ability of the higher centers in the
22 brain to completely block the incoming painful stimuli. That
23 ability develops after 32 to 34 weeks of gestation.

24 Q. Doctor, are there any studies supporting your view about
25 the increased sensitivity to pain in the fetus from 20 to 30

1 weeks?

2 A. Yes, there are. There is multiple lines of evidence that
3 support that.

4 Q. Is there a consensus in the medical community on this
5 issue?

6 A. Yes, there is. The people who are associated with the area
7 of pediatric pain are well aware of the increased sensitivity
8 during the earlier part of gestation.

9 Q. Doctor, can you apply your findings on fetal pain to
10 fetuses who suffer from an anomaly?

11 A. It depends on where the anomaly is located.

12 Q. Can you explain what you mean by that.

13 A. Sure. If a fetus has an anomaly of the brain or the spinal
14 cord or the nerves that participate in pain processing, then
15 these studies would not apply to those fetuses. For example, a
16 fetus may have a condition called congenital hydrocephalus
17 whereby there is an increased amount of fluid within the brain
18 that collects. Those fetuses may have an abnormal way of
19 processing pain.

20 There are other fetuses with chromosomal abnormalities
21 that have an abnormal brain development, and, again, those
22 fetuses would respond differently.

23 Q. Based on your opinion, can you tell us how those fetuses
24 would respond to pain.

25 A. You see, I am somewhat limited in my ability to answer that

1 question, because these fetuses have not been studied in any
2 great degree. For any scientific study what people try to do
3 is to exclude abnormalities of the nervous system in order to
4 see what is the normal response. So there is limited
5 information in that area.

6 There are some studies that have been done in
7 premature infants, premature infants who have documented injury
8 to the brain resulting from bleeding in the brain or a Roth
9 away of white matter, which is the fibers in the brain. And
10 there are some recent studies showing that their pain response
11 was no different from other premature infants with the same
12 neurological maturity who have no damage in their brain.

13 So if we were to extrapolate from these recent studies
14 t it would suggest that there are minimal differences between
15 fetuses who have abnormalities versus those who don't.
16 Although, the information in that area is very limited.

17 Q. Doctor, I would like to ask you a few questions about the
18 term "conscious's" as it relates to pain ~~***[STRIKE]***~~

19 "consciousness" as its RAOETS to pain. What is consciousness?

20 A. Consness is the substratum for every experience -FPT
21 without consciousness, experience is not possible. If I could
22 use a metaphor, we go to the movie theater and we see a story,
23 multiple characters, many events, their interrelationships and
24 interactions, etc. That is all happening as part of the movie.
25 But that is dependent on the screen. If the screen is not

1 there, then the movie doesn't exist.

2 Similarly, consness is the screen on which this
3 three-dimensional movie of all of nature is being played.

4 Q. Is consciousness necessary for the experience of pain?

5 A. Yes, it is.

6 Q. Do you have an opinion as to whether a fetus is conscious,
7 is able to feel pain at 20 weeks' gestation?

8 A. The multiple areas of evidence that we have noted from
9 fetal behavior, from their responses, etc., suggest that the
10 TPAETS is conscious, is able to relate to its environment in a
11 conscious perceiving way.

12 Q. What are those multiple lines of evidence?

13 A. Those are the lines of evidence that we have discussed
14 about fetal responses to sound, to light, to taste, to touch,
15 fetal responses to pain, the KORLTS of the functioning of the
16 brain. Even in the adult human, consciousness is very
17 difficult to measure. All that we know is it is correlated
18 with shifting patterns of activity within the cerebral cortex.

19 Similarly, we have data from fetuses at 19 to 20 weeks
20 of gestation showing the onset of shifting patterns of
21 electrical activity. We have different states of consness:
22 Wakefulness versus sleep, as adult \humans\hums, and similar
23 forms of behavioral states have been noted during fetal life.
24 SP-P.

25 So if you put all of the evidence that favors

1 consciousness on the one hand and suppose you propose the
2 hypothesis that the fetus is unconscious, there is little or no
3 evidence to suggest that the fetus is unconscious.

4 Q. Doctor, let me ask you, does a fetus have any capacity for
5 memory?

6 A. Yes. There are studies that have looked at learning and
7 memory during fetal life. There was a study done whereby women
8 were given a capsule which was tasteless and odorless, but this
9 capsule contained cue minute, an herb. They were asked to take
10 one capsule with their meals for a number of days during their
11 pregnancy. When their babies were born, these babies were able
12 to have an orienting response to the smell of cue PH*EUPB, even
13 though the mother had no taste or smell because these were
14 capsules that they swallowed.

15 So similar studies have been done looking at where the
16 mother was asked to read a certain story during her pregnancy,
17 and when she read the same story after the baby was born, there
18 was an orienting response: The baby calmed down, etc.
19 However, if she read the same story in a different tone, then
20 the baby was able to show a reaction to these changes in the
21 tone of the mother, even though that was the same story.

22 So there is a whole host of studies that have been
23 done in this manner showing that there is learning and memory
24 during fetal life. That, again, cannot occur without
25 consciousness.

1 Q. Is there a consensus in the medical community that
2 consciousness begins at 20 weeks?

3 A. This is a very controversial area. People have come up
4 with markedly different opinions in this regard, as to what
5 does consciousness begin.

6 Q. Within the community that studies the issue of fetal pain,
7 is there a majority view on when consciousness begins in the
8 fetus?

9 A. Yes. Having spoken and having interacted with many
10 scientists who are focused in this area of studying the fetus,
11 there is a consensus that consciousness would begin around mid
12 gestation. So 20 to 22 weeks of gestation. Although, there
13 are divergent opinions.

14 For example, in the mid nineties there was a British
15 parliamentary group that proposed that the fetus is conscious
16 from 6 weeks of gestation, whereas a working part of the royal
17 College of obstetrics and gynecology concluded that a fetus is
18 conscious after 26 weeks. There are many pediatricians who
19 believe that consciousness may manifest at the time of
20 conception, whereas other ped TREUGSs believe consness doesn't
21 occur until a full-term baby is born.

22 So there are divergent views on this topic. I think
23 the truth lies somewhere this the middle, which is what my
24 opinion is. At 2 to 22 weeks is the onset of fetal
25 consciousness.

1 THE COURT: Mr. Lane, is this a convenient time to
2 take our break?
3 MR. LANE: Certainly, your Honor.
4 THE COURT: Court will stand in recess for 10 minutes.
5 (Recess) 4 take 3 continuing direct of Dr. Anand
6 THE COURT: Mr. Lane, you may inquire.
7 MR. LANE: Thank you, your Honor.
8 Dr. Anand, I would like to turn now to your opinion
9 regarding fetal pain and the partial-birth abortion procedure;
10 what is your understanding of the partial-birth abortion
11 procedure?
12 A. A very rudimentary -P one. I am not an obstetrician and I
13 have not performed this procedure before, but from the
14 descriptions I have read of it, the procedure consists of
15 cervical dilation, the surgeon reaches inside the uterus and
16 grasps the lower extremity of the fetus, the fetus is turned
17 around, if it's in the head down position the feet and the
18 lower part of the body are delivered and this incision is
19 made in the back of the head and vacuum suction is applied to
20 suction out the brain of the fetus.
21 Q. You said your understanding is rudimentary, where did you
22 acquire your rudimentary understanding?
23 A. It's basically from descriptions of the procedure on the
24 web and other sources.
25 Q. Doctor you you is said you never performed this procedure,

1 have you ever performed any abortion procedure?
2 A. I have not.
3 Q. Do you have any personal views regarding a woman's right to
4 abortion?
5 A. I feel a woman has an inalienable right to obtain an
6 abortion to protect her health and mental well being with the
7 caveat that it should not cause pain to the fetus.
8 Q. Do you have an opinion as to whether the partial-birth
9 abortion procedure will cause pain to a fetus?
10 A. Yes, it would, if the fetus is beyond 20 weeks of
11 gestation.
12 Q. And could you describe, in your opinion, what kind of pain
13 you would anticipate the fetus would feel?
14 A. Given the increased sensitivity to pain at that period
15 of gestation, the parts of the procedure associated with
16 grasping the lower extremity of the fetus, of manipulation and
17 rotating the fetus within the confines of the uterus, of
18 delivering the fetus through an incompletely dilated cervix as
19 well as the surgical incision made at the back of the head, the
20 puncturing of the intracranial cavity through the occipital
21 PWOEB and through the membranes that covered the brain, all of
22 those parts of the procedure would be associated with prolonged
23 and excruciating pain to the fetus.
24 Q. Doctor, can you describe the membranes covering the brain
25 and how it relates to pain per is acceptance?

1 A. Certainly.
2 There are three membranes that cover the brain,
3 they're called the M M matter the arachnoid matter and the did
4 you remember are a matter. These are basically medical terms
5 that are give to these membranes.
6 The outer most membrane is a fairly tough membrane and
7 is highly innervated so it is likely to cause is severe pain
8 any injury to this membrane.
9 As an example, simply a slight dilation of blood
10 vessels within that membrane leads to a severe throbbing
11 headache. So, by corollary, any injury to that membrane would
12 cause excruciating main pain.
13 Q. Doctor you have used the term innervated a few times here
14 this morning, could you explain what that term means?
15 A. It simply means that that particular structure is supplied
16 with sensory nerves and has the recept I don't understand and
17 the never connections for stimulating the pain system.
18 Q. Doctor, based on your rudimentary understanding of this
19 procedure, does that include a rudimentary understanding of
20 when this procedure is performed?
21 A. I believe this procedure is performed during the second or
22 third trimesters of pregnancy.
23 Q. Doctor, do you have an opinion as to whether the crushing
24 of the fetal skull with forceps would cause pain to the fetus?
25 A. Yes, it would.

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1 Q. Do you have an opinion as to whether puncture of the is
2 skull without suction would cause pain to the fetus?
3 A. Yes, it would.
4 The fetal skull and the bones of the fetal skull are
5 covered by a membrane called the per I OS teeium, and that is
6 also richly innervated with sensory fibers.
7 Q. Doctor, can you is explain for the Court how you are able
8 to render an opinion about this particular procedure despite
9 the fact that you have not performed it?
10 A. This is based on the development of the pain system and
11 what appear to be the man -P EUP HRAEUGSS of the fetus
12 associated with this procedure.
13 Q. Doctor, do you you have any experience in delivering
14 babies?
15 A. Yes. I have delivered several thousand babies.
16 Q. Does that assist you in any way in rendering your opinion
17 in this case?
18 A. Most of the babies that I is have delivered were full-term,
19 this was during my training in India, and it has limited
20 relevance to the period of gestation that we are talking about
21 after 20 weeks.
22 Q. Doctor, do you have an opinion as to whether an abortion
23 procedure in which the fetal body is dismembered as it is being
24 removed would cause pain it the fetus?
25 A. If that procedure occurs after 20 weeks of gestation, it

1 certainly would.

2 Q. Do you have an opinion as to whether cutting the cord would
3 cause pain to the fetus while in utero?

4 A. The umbilical cord is not innervated, it is not supplied
5 with sensory nerves and is so the cutting the umbilical cord
6 would cause minimal, if any pain at all.

7 The reason why I say it could cause minimal pain is
8 because the blood vessels TPH-LTZ umbilical cord are -P
9 innervated, the umbilical arteries and the umbilical vein, so I
10 think the pain would be, you know, mild, if anything at all.

11 Q. Doctor, do you have an opinion as to whether an injection
12 of the feet I sideal agent such as die SKWROFPL gin or KCL into
13 the body would cause pain it the feet?

14 A. It would cause pain with EURBLT duction of the needle into
15 the fetal body yes.

16 Q. Can you describe for the Court how that would compare in
17 your opinion to the partial-birth abortion procedure?

18 A. The application of a needle to the fetal body would cause
19 acute pain which would be transient because it would be
20 followed by the injection of a lethal drug, like digoxin or
21 potassium chloride.

22 And is so, it would be an acute really fairly short
23 lived pain.

24 Q. Doctor, do you have an opinion as to whether labor
25 induction by administration of prostaglandins causes pain to

1 the fetus?

2 A. It would be unlikely to cause pain to the fetus because the
3 prostaglandins are not being injected into the fetal body.

4 However, there is a possibility that prostaglandins
5 may have some effect on the fetus because prostaglandins are
6 part of the chemicals that are released at the time of
7 inflammation and other painful lesions.

8 So, they're also released during acute pain.

9 So, if there is any direct contact of those
10 prostaglandins with the pain nerve endings in the fetal skin,
11 then it could possibly cause pain. But I think the likelihood
12 is fairly small.

13 Q. Doctor, do you have an opinion as to whether vaginal
14 childbirth at full-term causes pain to the fetus?

15 A. There is an is experiential content associated with natural
16 childbirth, although the mechanisms of nature are is such that
17 the, at full-term the baby being born is spared some of the is
18 experiential content.

19 What I mean is that as part of the normal process by
20 which labor is, occurs naturally, there is the release of
21 various hormones, beta indoor fin for example, those levels
22 rise markedly during a normal vaginal delivery.

23 And if you sample the umbilical cord blood from a
24 full-term newborn baby the beta indoor fin levels are about a
25 thousand fold higher than some of the highest levels documented

1 in the blood of adult patients.

2 There are other mediators that may be released as such
3 causing an activation of the indodge I TPHOUS systems of
4 analgesia within the body.

5 Q. Doctor, I want to make sure I have this clear.

6 What, effect, if any, is the release of indoor fins in
7 a full-term vaginal delivery have on a fetal experience of
8 pain?

9 A. Well, it produces -- you see beta indoor fins are drugs --
10 sorry, are mediators, they're not drugs, but beta indoor fins
11 are mediators that attach to the receptors which are the same
12 receptors that bind morphine or TPEPB to nil or some of the
13 opiate drugs, and these are indodge I TPHOUS -P open AOE
14 KWROEUDZ and they result in what's called fetal inhibition is
15 so that there is a generalized inHEUB I Tory tone that is
16 throughout the central nervous system of the baby who is being
17 born.

18 Q. Doctor, do you have an opinion as to whether a is caesarean
19 is section causes pain to the fetus?

20 A. If the fetus is beyond 20 weeks and is delivered through a
21 wide enough surgical incision, it should not cause pain to the
22 fetus.

23 Q. Doctor, I would like to cover one last topic with you,
24 which is the topic of maternal anesthesia. Can you is explain
25 how anesthesia administered to the mother reaches the fetus, if

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1 it does at all?

2 A. Anesthesia that is given to the mother would first have to
3 avoid metabolism by the maternal liver, would have to circulate
4 throughout the maternal blood system and would have to then get
5 transferred across the placenta.

6 See the circulation of the mother and the circulation
7 of the baby are separated by what's called the placental
8 membrane and this membrane is fairly thick to start out with
9 and asthma KHERT occurs this membrane becomes thinner and
10 thinner so that the efficiency of transfer across the membrane
11 in, say, the third trimester is much greater than what would be
12 seen in the is second or first trimester.

13 And so, the blood -- the drugs that are circulating in
14 the mother's blood have to get across this placental membrane
15 and reach sufficient enough concentrations in the fetus' blood
16 in order to then cross the blood brain barrier and have an
17 impact on brain cells in the fetus.

18 For these reasons multiple experts is have noted that
19 the fetus serves as a deep compartment within a pregnant mother
20 is so that this is a compartment that does not equill I bright
21 is very quickly with blood levels that are circulated in the
22 mother.

23 Q. Do you have an opinion as to whether anesthesia
24 administered intravenously to a pregnant woman would prevent
25 the fetus from feeling pain?

1 A. I think there is very low likelihood that it could have any
2 impact on the fetus simply because the concentrations that are
3 generated in the fetal blood would not be effective.

4 Q. We heard testimony here at trial from Dr. Westhoff about
5 the use of the drug TPEPB to nil when administered as an
6 anesthetic agent to a pregnant woman and whether that would
7 result in a fetus feeling pain.

8 Can you tell me whether you think that drug
9 administered to a woman, fentanyl, would result in fetus not
10 feeling pain during a painful procedure?

11 A. Well, that would depend on how the fentanyl is administered
12 and how much fentanyl is given.

13 If I had a patient, say, a pregnant mother with a
14 weight of say 65 or 70 kilograms, I would probably give a dose
15 of, say, 200 micro grams of fentanyl. This 200 micro grams of
16 fentanyl would then be distributed within the mother's blood
17 volume of, which is five liters, giving a fairly low
18 concentration.

19 Also, fentanyl is a drug that is rapidly metabolized
20 by the liver, so within minutes or seconds even after the
21 fentanyl injection the drug is undergoing metabolism.

22 There have been studies that have shown that at
23 full-term, which is when the placental transfer is the most
24 efficient the concentration within the fetal blood is about 30
25 percent that of the maternal blood is so that -P given the

1 higher sensitivity to pain in the fetus, this drug would not
2 accumulate in a sufficient concentration to guarantee that the
3 fetus is does not feel pain.

4 Q. You had is said that the transfer rate -- close to -- at or
5 close to full-term is 30 percent, would you expect that rate to
6 be higher or lower or the same in second trimester?

7 A. I would expect it to be less efficient transfer of fentanyl
8 in the second trimester.

9 So, if anything, we would be overestimating applying
10 these data to the second trimester, we would be overestimating
11 the fetus' exposure to TPEPB TA nil.

12 Q. Doctor, we heard testimony in this case from Dr. Westhoff
13 and Hammond about the use of the TKROG propofol when
14 administered to the woman as anesthesia. In your opinion,
15 would this prevent a fetus from feeling pain?

16 A. I believe there, as stated before, that there would be a
17 very low likelihood that this would have any effect on the
18 fetus.

19 Q. Could you explain your opinion as it pertained to the drug
20 propofol?

21 A. Certainly.

22 There are several studies that have looked at propofol
23 given during the third trimester and one particular study that
24 I can recall is published by TKPW*EUPB, et al, in 1995. They
25 injected a therapeutic dose of propofol into the mother and

1 measured concentration both in mother's blood as well as in
2 baby's blood.

3 What they found was that following a dose of, say, 150
4 milligrams given to a mother the concentration that resulted in
5 mother's blood was about one micro gram per CC of mother's
6 blood. And what they found was that the ratio of the
7 concentration in the baby's blood was about 65 percent that of
8 what was in the mother's blood.

9 Is so, it was about two thirds of what the mother's
10 blood concentration was.

11 And if we postulate that that is the concentration a
12 fetus is exposed to, that would be a very, very low concentration.
13 On the other hand, when propofol anesthesia is given directly
14 to premature neonates, they reach concentrations that are 50 times
15 higher than what a fetus would have been
16 exposed to.

17 So, if we are to ensure that there is a state of fetal
18 anesthesia, we would need to give anywhere from five to 50
19 times the dose of regular doses at the time that is used for
20 the mother.

21 Q. In your opinion, would those doses required be safe for
22 maternal health?

23 A. There certainly would be the, a very high likelihood of
24 toxic side effects in the mother.

25 Q. Doctor, are you aware of any intravenous anesthetic drugs

1 that when safely administered to the mother, would provide
2 complete pain relief to the fetus?

3 A. I'm not aware of any drugs given intravenously into
4 the mother that would provide safe anesthesia to the fetus.

5 These two drugs that you noted, propofol and etomidate
6 (TPHOL), are the drugs that are most lipid soluble, meaning
7 they're soluble in lipids and that's why they are
8 transferred across the placental membranes much more
9 efficiently than other drugs, for example, morphine or meperidine
10 (MEPEPL) they're less lipid soluble so less of those drugs
11 would get across the placenta to reach the fetus.

12 Q. Doctor, let me ask you the same question about an epidural,
13 would an epidural administered to a pregnant woman,
14 would that provide pain relief to the fetus?

15 A. No, it would not.

16 Q. Why is that?

17 A. Because the epidural drugs used for epidural anesthesia or
18 local anesthetics at the time, for example the positive dental
19 never block which is blocking the nerves going to the uterus,
20 these drugs are injected directly in the vicinity of the nerves
21 that are supplying the uterus and surrounding structures. A
22 very small amount of these drugs get absorbed into the mother's
23 blood stream and those concentrations are extremely low in the
24 mother's blood stream which would mean that the fetus hardly
25 sees any of those drugs.

1 Also, these drugs are effective when they're injected
2 in the vicinity of a nerve fiber. So, if you think about it,
3 the concentration in that vicinity of the nerve fiber is
4 thousand fold higher than what it would be in the blood stream.

5 Is so, those drugs would be unlikely to provide any
6 form of anesthesia or pain relief to the fetus.

7 Q. Doctor, I understand your point about the transfer across
8 the PHRA is SEPBA, but can you explain why an amount
9 administered to the mother would not have an effect on the
10 fetus when a fetus is so much smaller than the mother?

11 A. Simply because the drug injected into the mother has to
12 circulate in the mother's blood stream. A lot of the drugs are
13 metabolized by the mother's liver. There is an increased blood
14 flow to the liver during pregnancy.

15 Also, most anesthetic drugs tend to reduce placental
16 blood flow so the PFCF rein is shall flow from the mother to
17 the placenta is reduced by -- that is specific effects of many
18 of these answers at the timic drugs.

19 And is so, the likelihood that sufficient amount of
20 anesthesia gets across to the fetus to have reasonable
21 therapeutic concentration to cross the fetal blood brain
22 barrier and have an impact on the brain of the fetus is very
23 low.

24 Q. Let me ask you about the amount of anesthetic that would be
25 needed to provide relief to the fetus; that the same or higher

1 or lower level than you would find in an ATKULT?

2 A. Well, we can go back to that study that we had referred to
3 earlier, the one by Nicholas TPEUFBG from imperial college
4 London.

5 What they gave directly to the fetus was 12.5
6 micrograms per kilo gram of fentanyl. This was injected
7 directly into the circulation of the fetus. This would produce
8 a blood concentration that would be 25 to 50 times the blood
9 concentration produced by giving an appropriate amount of the
10 same drug to the mother.

11 So, very unlikely to have an anesthetic effect on the
12 fetus.

13 Q. What effect, if any, does it have, in your opinion, as
14 to the amount of anesthetic drug needed as it pertains to the
15 fetus' increased sensitivity to -P pain during the period of
16 is gestation from 20 to 30 weeks?

17 A. We can -- because those studies have not been done all I
18 can do is speculate that a fetus would require higher doses or
19 higher concentrations in the fetal circulation in order to
20 achieve anesthetic concentrations.

21 The only study that I know of direct fetal anesthesia
22 is the one that we refer to from London.

23 MR. LANE: Thank you, Doctor. No further questions,
24 your Honor.

25 THE COURT: Ms. Wigmore, do you want to inquire?

1 MS. WIGMORE: Yes.
2 CROSS-EXAMINATION
3
4 BY MS. WIGMORE:
5 MS. WIGMORE: Your Honor, I have a binder for
6 counsel for the government and for the witness to make it
7 easier should we need to reference prior testimony or the
8 articles that were attached to Dr. Anand's report.
9 May I approach?
10 THE COURT: By all means.
11 CROSS-EXAMINATION
12
13 BY MS. WIGMORE:
14 Q. Good afternoon, TK*PB I'm Amy wig more, we met at your
15 deposition.
16 A. Good afternoon.
17 Q. And just to let you know what I have provided you, I have
18 given you a copy of some prior testimony and your expert
19 report, that's in one binder, and then you have the
20 articles that were attached to your is expert report, as well
21 as one additional article that you provided at your deposition.
22 Dr. Anand, on direct examination you mentioned that
23 you previously -- that you're a pediatric intensivist, correct?
24 A. That's right.
25 Q. And is isn't it true that about 99 percent of your time is

1 devoted to the department of pediatrics?
2 A. The department of pediatrics pays my salary and that's
3 where the major part of my efforts are focused.
4 Although, what I do for the department of pediatrics
5 also helps the departments of answers tease KWROL isgy and
6 pharmacology, etc.
7 Q. Dr. Anand, didn't you testify previously that 99 percent of
8 your time is devoted to the department of pediatrics?
9 A. Yes, it is.
10 Q. And, you mentioned that you had adjunct appointments in
11 anesthesiology, neuro biology and farm aaKOLgy, is that right?
12 A. That is right.
13 Q. Isn't it true that one percent or less of your time is
14 currently is spent on anesthesiology?
15 A. You could characterize it as that or basically like I said
16 before, while I'm working in department of pediatrics, because
17 this is all within the same medical school, I am also
18 simultaneously teaching anesthesia residents or fellows at
19 the time when I am teaching pediatric residents and critical
20 care fellows.
21 Q. Dr. Anand, did you give a deposition in this case?
22 A. Yes, I did.
23 Q. And, what I would like you to do is refer to the transcript
24 of that deposition is which is one of the two binders that I
25 gave to you, and refer, specifically, to page 43. And I want

1 to direct your attention to page 43, line 23. And I want to
2 ask you, did I ask you these questions and did you give
3 these answer:
4 "Q so, based on your indication that about 99 percent
5 of your time is in pediatrics, is it fair to say that one
6 percent or less of your time is currently spent in
7 anesthesiology?
8 "A that's right."
9 Did I ask that question, did you give that answer?
10 A. Yes, I did.
11 Q. And it's true, isn't it, that you are not an answers
12 teeth KWROL gist?
13 A. I'm not board certified in anesthesiology; that's right.
14 Q. In fact, you have never been board certified in that
15 discipline, have you?
16 A. That is correct.
17 Q. And in your current position you are not acting as an
18 answers teeth KWROL is gist, correct is?
19 A. Except when I am on the sedation service, I provide
20 anesthesia to patients who require deep sedation or anesthesia
21 throughout the hospital.
22 Q. Dr. Anand, could you refer, please, to page 42 of the
23 deposition transcript? I want it to direct your attention to
24 line 6 on page 42.
25 "Q is it fair to say that you don't currently

1 act as an answers tease KWROL is gist?
2 "A is absolutely right."
3 Did I ask you that question, did you give that
4 answer?
5 A. That's right. I don't perform all the duties of an
6 anesthesiologist.
7 Q. Thank you.
8 You do not consider yourself an expert in
9 anesthesiology, isn't that right?
10 A. That is right.
11 Q. And you are not trained or certified to deliver anesthesia
12 to any patients, correct?
13 A. That is correct.
14 Q. You don't deliver general anesthesia, do you?
15 A. With the caveat of my practice on the sedation service at
16 Arkansas Children's Hospital, I do not deliver anesthesia in
17 the operating room.
18 Q. And you do not deliver anesthesia to pregnant women, do
19 you?
20 A. No, I do not.
21 Q. In fact, you have never administered anesthesia to a mother
22 is receiving an abortion, right is?
23 A. That is right.
24 Q. You are not a member of any professional organizations
25 relating to anesthesiology, is that right?

- 1 A. That is correct.
2 Q. And you are not an obstetrician, are you is?
3 A. No, I'm not.
4 Q. You are not a gynecologist, correct?
5 A. That is correct.
6 Q. And you have never performed an abortion, have you?
7 A. No.
8 Q. Is it true that you have never written any articles
9 about abortion?
10 A. That is true.
11 Q. And is it true that you have never participated in or
12 conducted any studies of any method of abortion?
13 A. That is correct.
14 Q. Now, your' testifying here today claiming to be an expert
15 in fetal pain, is that correct is?
16 A. That is correct.
17 Q. Is it true that you are not currently conducting any
18 research on pain in the human fetus?
19 A. That is correct.
20 Q. Now, you participated in a study regarding fetal pain
21 when you were at Oxford, is that right?
22 A. That is right.
23 Q. And that was roughly 20 years ago, is that correct?
24 A. Yes, you're right.
25 Q. And that was a study by WA* keen PA*R of, is that

- 1 correct?
2 A. There were actually two studies, one was by Jo akin PA*R of
3 and there was another study published by Joseph SPH*EUT.
4 Both of those are referenced in my C isV.
5 Q. The study involving Dr. PA*R of, that was your rave-your
6 real in that study was to review the study protocol and help in
7 interpreting the result, is that right?
8 A. That is correct.
9 Q. And you are not listed as an AUPBLGon on the publication
10 that emerged from that study, correct is STPH-RPBLGTS
11 absolutely.
12 Q. And you mentioned you participated in one other study
13 involving Dr. Smith, is that correct?
14 A. That is correct.
15 Q. And that study did not conclude that a fetus feels pain,
16 did it?
17 A. It was a study on fetal stress responses as a result of
18 hypoxia and had nothing to do with fetal pain.
19 Q. It wasn't even designed to investigate that question,
20 correct?
21 A. That is correct.
22 Q. Is it true that you haven't participated in any studies
23 directly addressing the question of fetal pain since
24 the 1980s is?
25 A. That is correct.

1 Q. Dr. Anand, your opinions in this case address whether or
2 not a fetus will experience pain as a result of a partial
3 birth abortion, correct?
4 A. Correct.
5 Q. And you include in your expert report a description of five
6 steps that constitute a partial-birth abortion, is that
7 right?
8 A. That is right.
9 Q. Now, before you spoke with counsel for the government in
10 this case you thought that the Partial-Birth Abortion Ban Act
11 covered the vast majority of procedures used for abortion,
12 didn't you?
13 A. Yes, I did.
14 Q. But counsel informed you that the Act applied to only one
15 procedure, is that right?
16 A. That is right.
17 Q. And they told you that procedure was dilation and
18 extraction, or D&X, is that right?
19 A. D&X, that's right.
20 Q. Now, when you use the term partial-birth abortion you
21 equate that with D&X, is that correct?
22 A. To a large extent, yes.
23 Q. The D&X procedure has been given other names as well, is that
24 that right?
25 A. That is right.

1 Q. And one of those names is intact D&E, correct?
2 A. Correct.
3 Q. Now, you have never performed a D&X, correct?
4 A. That is correct.
5 Q. In fact, you have never even observed one, is that right?
6 A. That is right S.
7 Q. And you have testified on direct that your understanding of
8 that procedure is only rudimentary, is that right?
9 A. That is correct.
10 Q. Is so you are not an expert in any way in the D&X, are
11 you?
12 A. On the D&X procedure, no, I'm not an expert. I don't claim
13 to be one.
14 Q. And in fact before you talked with the government's
15 attorneys in this case you did not have much of an
16 understanding of the D&X procedure, did you?
17 A. That is correct.
18 Q. And you are not aware of any studies on fetal pain
19 during the D&X procedure, are you?
20 A. There are no published studies that I'm aware of.
21 Q. And you're not aware of any articles on that subject, are
22 you?
23 THE COURT: Are you suggesting there are, Ms. Wigmore.
24 MS. WIGMORE: No, I'm suggesting that there aren't any
25 and I want the doctor to confirm that.

1 Is that true, Doctor?
2 A. That is true.
3 Q. Dr. Anand, you testified that pain is a subjective is
4 experience, is that correct?
5 A. That is correct.
6 Q. Our inability to communicate with the fetus prevents us
7 from studying pain and its subjective in individual fetuses,
8 isn't that right?
9 A. That is correct.
10 Q. And as of today the technology does not exist to enable us
11 to determine whether the fetus actually experiences pain, is
12 that right?
13 A. That is correct. Direct evidence for fetal experience is
14 not possible at this point.
15 Q. Now pain is both a is sensory and an emotional experience,
16 is that right?
17 A. That is correct.
18 Q. And the emotional aspects of a painful is experience can be
19 very specific for the individual, is that correct?
20 A. That is correct.
21 Q. And at this point we are not smart enough yet to evaluate
22 whether a fetus has the emotional experience of pain, is that
23 correct?
24 A. Direct evidence for the emotional experience of pain is not
25 available to us.

1 Q. In fact, it's technologically impossible at this time to
2 measure emotions in the fetus, correct?
3 A. Yes.
4 All we can do is to measure responses that could be
5 classified as emotional responses of the fetus such as facial
6 expressions or fetal behavior that can be studied under
7 ultrasound.
8 Q. Dr. Anand I am going to ask the question again.
9 Is it technologically impossible at this time to
10 measure emotions in the fetus?? Yes or no.
11 A. No.
12 Q. It's not technologically impossible?
13 A. Sorry. It is. It is I -- yes.
14 Q. Yes, it is technologically impossible to measure emotions
15 in the fetus, correct?
16 A. That is correct.
17 Q. Now, one cannot is experience pain without consciousness,
18 is that right?
19 A. That is right.
20 Q. And currently there are no studies examining the
21 consciousness of the fetus, is that right?
22 A. There is no way that consciousness can be directly measured
23 in the fetus or any other living being.
24 Q. That simply can't be done, correct?
25 A. Absolutely.

1 Q. And there is no consensus in the medical community that
2 consciousness begins at 20 weeks' gestational age for fetuses,
3 isn't that right?
4 A. That is correct.
5 Depending on the medical community that you refer to
6 being the general medical community.
7 Q. And in fact you testified in your deposition that there is
8 no consensus in the medical community that consciousness begins
9 at 20 weeks, isn't that right?
10 A. That is right.
11 Q. You you would agree that it has not been definitively
12 established that there is consciousness starting at 20 weeks '
13 gestational age, isn't that right?
14 A. That is right.
15 The prevalent view of consciousness is that it's not
16 an all or nonphenomenon.
17 Q. Thank you, Doctor, I just want you to answer my questions
18 and I am sure Mr. Lane will is have an opportunity to SKW- you
19 more questions?
20 THE COURT: Please, Ms. Wigmore, you is ask the
21 questions, you don't do the rulings 20.
22 MS. WIGMORE: Your Honor I ask the witness be
23 instructed to answer my questions.
24 THE COURT: Ms. Wigmore, just ask the questions. If
25 you don't mind, I will continue to control the enforcement of

1 this Court.
2 MS. WIGMORE: Thank you, your Honor.
3 Q. Dr. Anand, is is it true that there is no defibtive
4 evidence of when fetal consciousness begins?
5 A. There is no definite evidence to that effect.
6 Q. And it may not be possible to obtain unequivocal evidence
7 of fetal consciousness, isn't that right?
8 A. Yes, ma'am, yes, attorney.
9 Q. Is.
10 Q. Now, are you familiar with the Royal college of obstetrics
11 and STKPWAOEUPB KOLgy in the united kingdom, aren't you?
12 A. Yes I am.
13 Q. Is that organ iszation established a working grab to review
14 materials available regarding fetal consciousness, is that
15 correct is?
16 A. That is correct.
17 Q. And I believe you testified on direct that that group
18 concluded that fetal consciousness did not occur before 26
19 weeks, is that correct?
20 A. That is correct.
21 Q. And in fact it's possible that consciousness does not
22 develop until the moment of birth, is isn't that right?
23 A. Possible, but highly improbable.
24 Q. The process of birth and the demands of independent
25 existence exutero may serve as a trigger for the expression of

1 -P consciousness.
2 Is that right?
3 A. That is a statement of the null hypothesis, yes.
4 Q. And you in fact have stated those exact words in a
5 publication that you relied on in forming your opinions in this
6 case, haven't you is?
7 A. That is correct.
8 Q. And that is the publication behind Tab 5 and item 5 in your
9 expert report, is that correct?
10 A. Yes, that is the manuscript.
11 Q. That's an article that you published in 1999, correct?
12 A. That is correct.
13 Q. Your opinion on fetal con SHOUSZness is based in part on
14 studies of preterm neo TPHA*Eves, is that right?
15 A. That is right.
16 Q. Those are premature babies, is that right?
17 A. That is right.
18 Q. But there are physiological differences between a pre-term
19 neo TPHA*EUT and a fetus, aren't there?
20 A. Yes, there are some physiological differences.
21 Q. And there are radical differences in terms of environmental
22 experiences between a pre-term neo TPHA*EUT and a fetus,
23 correct?
24 A. Correct.
25 Q. Dr. Anand, do you recall testifying about studies relating

1 to fetal memory?
2 A. Yes, I do.
3 Q. Isn't it true that those is studies have been on fetuses in
4 the third trimester of PREG -P Nancy?
5 A. Most of those studies were done in the third trimester of
6 pregnancy.
7 Q. And you are not aware of any studies showing memory of is
8 experience for fetuses of 20 weeks, isn't that right?
9 A. That is right.
10 Q. You agree, don't you, that there is a difference of opinion
11 in the medical community regarding the ability of a fetus to
12 experience pain?
13 A. There are differences of opinion, yes.
14 Q. And there are people in the medical community who have
15 disagreed with your opinion that a fetus can feel pain, isn't
16 that right?
17 A. That is right.
18 Q. And you have reviewed articles that are inconsistent with
19 the opinions you are offering in this case, isn't that right?
20 A. Yes. For due diligence I KHRUTDed those articles in my
21 expert report.
22 Q. Now you mentioned on your direct examination that your
23 opinions involve is some extrapolation, correct?
24 A. That is correct.
25 Q. You have extrapolated from animal models, haven't you is?

1 A. Yes, I have.
2 Q. And you have extrapolated from is studies on premature
3 babies, is that right STPH-RBLGTS that is right.
4 Q. And your opinion is also based on some inferences as you
5 described on your direct examination, correct correct is?
6 A. Yes, that is right.
7 Q. And it's true, isn't it, that your entire opinion in this
8 case is based on inference?
9 A. The opinion is based on multiple lines of evidence. Yes,
10 we infer from those lines of evidence that the fetus feels pain
11 and is consciously aware of that pain.
12 Q. Is it true that your entire opinion in this case is based
13 on inference?
14 A. Yes, that is correct.
15 MS. WIGMORE: Your Honor, I'm about to move to a new
16 subject, I would be happy to continue but if you would like to
17 take the lunch break this would be a good time.
18 THE COURT: Thank you for alerting me to that,
19 Ms. Wigmore and allowing me that judgment.
20 We will take our reiss says at this time and reconvene
21 at 2:00.
22 (will you know -P oon recess) TWM 4/13 take 4.
23 AFTERNOON SESSION
24 2:00 p.m.
25 KANWALJEET ANAND, resumed.

1 THE COURT: Good afternoon.
2 Cross-examination (continued)
3 BY MS. WIGMORE:
4 Q. Dr. Anand, is it true that you are getting paid for your
5 testimony here today?
6 A. Yes, I am.
7 Q. You are being paid at a rate of \$SHAO per hour, is that
8 right?
9 A. That's right.
10 Q. You are also testifying for the government in two other
11 cases on the same subject matter, is that correct?
12 A. That is correct.
13 Q. In those cases are you also receiving \$450 per hour for
14 your work?
15 A. That is correct.
16 Q. Doctor, have you ever had an article rejected by a peer
17 reviewed journal?
18 A. Certainly, many times.
19 Q. In any of those cases has the article that has been
20 rejected by the journal ever been published by another journal?
21 A. That is correct.
22 Q. You rely on surrogate markers as evidence of fetal pain, is
23 that correct?
24 A. That is correct.
25 Q. Those Americas include hormonal stress sponses, is that

1 right?
2 A. That is right.
3 Q. And circulatory responses is another marker that you use,
4 correct?
5 A. That is correct.
6 Q. But isn't it true that hormonal and circulatory responses
7 are not definitive evidence of conscious pain perception?
8 A. That is correct.
9 Q. Just because a fetus experiences a stress response does not
10 necessarily mean that it is experiencing pain, correct?
11 A. Unless the stress response follows a painful stimulus.
12 Q. But you would agree that a stress response does not equate
13 with pain?
14 THE COURT: I think the witness answered your
15 question.
16 A. I don't understand your question, counsel.
17 THE COURT: Let me ask you something, Dr. Anand. Did
18 you already testify in Nebraska and California?
19 TH
20 THE WITNESS: I have testified in Nebraska but not in
21 California yet, your Honor.
22 THE COURT: Are you going to testify in California?
23 THE WITNESS: Yes, sir.
24 THE COURT: When?
25 THE WITNESS: That testimony is scheduled for Thursday

1 of this week.
2 THE COURT: Fine. Thank you.
3 You may continue.
4 Q. Your conclusion that these surrogate markers indicate pain
5 is based on studies of pain in children and adults, is that
6 right?
7 A. Partly, yes.
8 Q. Is it true that there are huge physiological differences
9 between fetuses and adults?
10 A. That is correct.
11 Q. These huge physiological differences impact your ability to
12 extrapolate from studies on adults to fetuses, isn't that
13 right?
14 A. That is right.
15 Q. There are also physiological differences between fetuses
16 and children, correct?
17 A. That is correct.
18 Q. Those differences also impact your ability to definitively
19 conclude that the surrogate markers evidence pain in the fetus,
20 is that correct?
21 A. To some extent, yes, that is correct.
22 Q. Isn't it true that some of the articles that you considered
23 in forming your opinion in this case make clear that the
24 measure of stress response has limitations in determining the
25 existence of fetal pain?

1 A. Yes, that is stated in some of the articles as the null
2 hypothesis.
3 Q. The article by Dr. Smith, which was number 13 listed in
4 your expert report, indicated that stress responses do not
5 necessarily imply pain, correct?
6 A. Can I look at the article you are referring to?
7 Q. Sure. If you would refer, please, I believe it is tab 13
8 in the binder I handed you. That is an article by Richard P.
9 Smith and others, correct?
10 A. That is correct.
11 Q. That is an article from the year 2000, correct?
12 A. That is correct.
13 Q. If you could refer, please, to page 162 of that article,
14 the left-hand column under heading "4. Fetal stress.?"
15 A. Yes.
16 Q. I am just going to read a portion of that paragraph for
17 you. "Because of the obvious difficulties in studying fetal
18 behavior, activation of the hypothalamus will maintain but regulate
19 adrenal access (a stress response) has been proposed as a
20 surrogate indicator of fetal pain. TH-FPLT has limitations.
21 Stress responses do not necessarily imply pain (for example,
22 during exercise) and stress responses do not involve the
23 cortex."
24 That was a statement made by the authors of this
25 article, correct?

1 A. That is correct.
2 Q. Halves something you considered in forming your opinions?
3 A. That is correct. This was one of the articles I had
4 referred to.
5 Q. You also referred to an article by Modi & Glover, that is
6 tab 10, is that correct?
7 A. Yes. This is a chapter in a textbook that I had edited.
8 Q. That is from the year 2000, correct?
9 A. That is also from the year 2000.
10 Q. The authors of that article also noted that there are
11 limitations of using stress responses for determining fetal
12 pain, is that correct?
13 A. Where, may I ask, are you looking?
14 Q. If you could refer, please, to page 221. TKHRS, again, tab
15 10. The first column, third paragraph, first sentence. "The
16 PHAERBLT of stress responses is of value for determining the
17 existence of fetal pain is one is aware of its limitations."
18 That is a statement that was in this article, correct?
19 A. That is correct.
20 Q. These authors acknowledged the limitations of the stress
21 response as evidence of pain, correct?
22 A. Yes, that is correct.
23 THE COURT: Ms. Wigmore, is this a new school of
24 cross-examination where you make a statement and finish every
25 statement with "is that correct"?

1 MS. WIGMORE: Your Honor, I believe I asked the
2 witness about the article in the --
3 THE COURT: I didn't ask about the substance of the
4 question. I just asked if it is a school of cross-examination.
5 I notice the style over and over again. Go ahead.
6 Q. There are situations that are not painful in which an
7 increase in hormone levels will occur, isn't that right?
8 A. That is right.
9 Q. For example, if I were to clap loudly in front of Mr. Hut's
10 face, he might experience a stress response, is that right?
11 A. Very unlikely.
12 Q. Would his den lin levels possibly increase if he were taken
13 by surprise?
14 A. Very unlikely that there would be a measurable increase in
15 the adrenaline level.
16 Q. Held experience a physiological response to that, wouldn't
17 he?
18 A. Yes. You may see that as a transient increase in his heart
19 rate. But unlikely to have any hormonal response to that.
20 Q. A transient increase in his heart rate, is that a
21 circulatory response?
22 A. Yes, you could call it that.
23 Q. That circulatory response wouldn't indicate pain that was
24 being felt, would it?
25 A. Unless the level of noise was noxious, it would not mean

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1 pain.
2 Q. You considered a study by January in a col plows and others
3 in forming your opinion, is that right?
4 A. Which one would you be referring to?
5 Q. Tab 20 in the binders.
6 A. Yes, I did refer to this article.
7 Q. Is it true that that article indicates that since the
8 mechanisms involved in pain perception are not fully
9 understood, it is not possible to conclude that the fetus
10 experiences pain? I am referring to page 80. Specifically to
11 the right-hand column, the second paragraph from the bottom.
12 A. Yes, that is stated in this article.
13 Q. This is one of the articles you relied on in forming your
14 opinions?
15 A. Yes. This is published in 1994, and I did rely on this
16 article.
17 Q. You also relied on an article by van Hut hal lo and knew
18 about enHAOUPBZ, is that correct?
19 A. Which article?
20 Q. This is at tab 2 2.
21 A. Yes. This is one of the articles I included as part of due
22 diligence in my expert report.
23 Q. The title of the article is fetal pain question mark, is
24 that right?
25 A. That is right.

1 Q. I am directing your attention to the very last paragraph on
2 the first page of this article, which states, "Being purely
3 subjective, pain is a difficult parameter to measure. While
4 measurements of pain with cooperate are aative subjects are
5 based on subjective scales of pain intensity, these methods are
6 not KPHREUBGable to neonates or premature babies. Therefore, a
7 number of indirect methods have been developed to assess
8 clinically their possible painfulness. These methods are based
9 on change in either behavior, eg quality of cry or something
10 patterns (or unknow about I can PRABG testimony, e TKW-RBGS
11 pulse rate or blood pressure. They are still being developed.
12 None is yet suitable for its assessing the pain in fetuses."
13 Did I read that correctly?

14 A. That is correct.

15 Q. This article was published in 2000, is that right?

16 A. Yes. This was published before the definition of pain was
17 changed.

18 Q. The definition of pain you are referring to is a definition
19 by the international association for the study of pain, is that
20 correct?

21 A. Yes, that is correct.

22 Q. That is a document you reviewed with Mr. Lane during your
23 direct examination, correct?

24 A. Certainly.

25 Q. Now, are you familiar with a term "no se ception"?

1 A. Yes, I am familiar with that term.

2 Q. What does that refer to?

3 A. It refers to the measured activity within no, sir se
4 acceptors, which are the receptors responsible for transducing
5 noxious stimuli as well as the nerve fibers and the central
6 nervous system tracts and areas associated with the processing
7 of noxious stimuli.

8 Q. Would it be fair to say that no sus ception is the wiring
9 that TPHAEUBLGS one to experience a stress response? Is that
10 accurate?

11 A. No, that's not.

12 Q. Is it the ability, the receptors that you were referring
13 to, is that part of the no as you SEPTive pathway?

14 A. That is correct.

15 Q. Isn't it true that the pain definition you were referring
16 to indicates that activity induced in the no sus accept tore,
17 in the no susceptible pathways by a noxious stimulus is not
18 pain?

19 A. Where are you referring to?

20 Q. If you could refer to tab 2 in the binder that I handed to
21 you. This is the IASP pain terminology exhibit you reviewed
22 with Mr. Lane during direct examination, correct?

23 A. It is tab 1.

24 Q. Tab 1. I apologize. Do you have that?

25 A. I am just getting to it. Where are you looking in this

1 document?
2 Q. Before I move to that, this is the IASP definition that you
3 reviewed on direct examination, correct?
4 A. Yes, I believe, yes.
5 Q. If you could refer, please, to the second page of this
6 document, the very last paragraph, the last sentence states,
7 "Activity induced in the no sus accept tore and no susceptible
8 pathways by a noxious stimulus is not pain, which is always a
9 psychological state, even though we may well appreciate that
10 pain most often has a proximate physical cause, is that
11 correct?
12 A. Yes, that is what it says.
13 Q. Under the IASP definition of pain, activity induced in the
14 no sus accept tore and no susceptible PAG WAEUGTS is not by
15 itself pain, is that right?
16 A. That is what the committee PHRAOEFS yes,.
17 Q. You disagree with that, is that correct?
18 A. To some extent I do, yes.
19 Q. So in your mind a TKWEUPGS of pain should be equivalent of
20 no susceptible activity, is that right?
21 A. Not really. The converse may not be true.
22 Q. Is it true that you have advocated for a change in the
23 definition of pain respect?
24 A. Yes, counsel I have.
25 Q. The change you have suggested with respect to no sus

1 ception has not been accepted, is that right?
2 A. That was not a change that I had recommended. I had
3 recommended that the complete definition of pain be rewritten
4 whereas the committee bass not prepared to take that radical a
5 step. They simply compromised. This is decisions by
6 committee. Compromised in adding a note below the traditional
7 definition of pain.
8 Q. Your definition that you proposed was not completely
9 adopted by the committee, is that fair?
10 A. I had not proposed a definition at all, counsel. I had
11 recommended that a multidisciplinary working task force be put
12 together in order to rewrite the definition of pain, because
13 this one had out lived its usefulness.
14 Q. Is this the definition that you read on direct examination
15 the current definition of pain?
16 A. It is. That is the current definition of pain, counsel.
17 Q. You testified during direct examination that you believe a
18 fetus experiences pain at 20 weeks' gestation, is that correct?
19 A. That is correct.
20 Q. Is that 20 weeks after the last menstrual period? Is that
21 how you are measuring gestation?
22 A. That is 20 weeks after conception.
23 Q. So that would be approximately 2 weeks later than 20 weeks
24 LMP, correct?
25 A. That is correct.

1 Q. That would be 22 weeks LMP?
2 A. If LMP is the beginning of the pregnancy, if you measure it
3 from there, yes, it would be 22 weeks LMP.
4 Q. It is not your opinion that a fetus of 20 weeks gestation,
5 and I will use your 20 weeks, which is 20 weeks
6 post-conception, it is not your opinion that a fetus at that
7 age will definitely experience pain, correct?
8 A. Based on the multiple lines of evidence provided, there is
9 a greater likelihood than not that the fetus will experience
10 pain after 20 weeks post-conception.
11 Q. So it is your opinion that a fetus likely will experience
12 pain at 20 weeks, is that right?
13 A. To a great, high degree of probability, yes.
14 Q. It is possible that a fetus at 20 weeks will not experience
15 pain, correct?
16 A. A much lower probability. I have to retain my scientific
17 skepticism and be ready to challenge my pet hypotheses as well.
18 Q. So it is not a medical fact that a fetus between 20 and 24
19 weeks' gestation will feel pain, is that right?
20 A. It depends on how you define medical fact.
21 Q. It is not 100 percent certain, is it?
22 THE COURT: Are you talking about moral cert. sued,
23 Ms. Wigmore?
24 MS. WIGMORE: I am actually asking whether it is a
25 medical fact.

1 THE COURT: He tells you he doesn't understand what
2 that term means. You don't want to use "moral" I guess. OK.
3 Rephrase the question.
4 Q. Didn't you say in your deposition in this case that beyond
5 20 weeks of gestation there is a greater than 80 percent
6 probability that a fetus will experience pain when subjected to
7 tissue injury?
8 A. That is correct. That is what I meant by a very high
9 degree of likelihood that a fetus will experience pain.
10 Q. So there is a 20 percent chance that it won't, correct?
11 A. That is correct.
12 Q. Now let's talk about how you came up with an 80 percent
13 probability. You didn't do any mathematical calculations to
14 arrive at that number, did you?
15 A. No, I did not. I summarized the evidence and its value. I
16 tried to simplify the summary of my opinion in saying that
17 there is an 80 percent or greater probability that a fetus will
18 experience pain after 20 weeks post-conception. That doesn't
19 mean that out of 100 fetuses 80 or more will will pain and 20
20 and perhaps less will not experience pain. What it means is is
21 that for every fetus that is exposed to tissue injury, there is
22 a high degree of likelihood that they are experiencing pain [>
23 will will should be will experience<]
24 Q. That is based on your subjective review of the materials,
25 correct -FRPLTS that is correct.

1 Q. That is not based on a mathematical calculation, right?
2 A. It is based on what might be characterized as complex
3 multivariety analysis and assessment of the published data.
4 Q. Did you use any multiplication to get to that number?
5 A. No, I did not.
6 Q. Did you use any addition or subtraction?
7 A. No, I did not [> multivariousiat<]
8 Q. No studies have been done to determine definitively whether
9 a fetus could experience pain at 20 weeks, is that right?
10 A. It depends on what is the level of certainty that you want
11 to define "definitively" by. If you give me the level of
12 certainty that you desire, I could perhaps give you evidence
13 that may support that level of certainty or perhaps design an
14 experiment that could possibly come up with the level of
15 certainty that you desire.
16 Q. Doctor Anand, I am asking you, were any studies done up to
17 this point indicating definitively whether a fetus could
18 experience pain at 20 weeks?
19 A. I beg to submit that it depends on what you consider is
20 definitive evidence. If you tell me what is definitive
21 evidence, I can produce that before you.
22 Q. Can you please refer to page 183 of your deposition
23 transcript in the binder I provided to you.
24 A. Certainly. 183?
25 Q. 183. I want to direct your attention to line 14 on that

1 page. Do you have that? Do you have that, Dr. Anand?
2 A. Yes, I do.
3 Q. The question you were asked was, do you know what studies
4 would need to be done to determine definitively whether a fetus
5 could experience pain at 20 weeks?
6 "A. Yes. I think some of the studies that could be done would
7 be the neuroimaging studies, studies such as using functional
8 MRI scans or studies using PET scans -- PET stands for pos I
9 tron E mission tomography scans. And those scans would, if
10 done immediately after a painful procedure is performed to the
11 fetus, would show activity in areas of the brain that are
12 associated with pain processing. That really would be very
13 strong evidence, almost definite evidence, in view of pain
14 processing in the fetus. It would also be the kind of evidence
15 that could explore the patterns of pain processing in fetuses
16 before 20 weeks, if there is any, and could, you know, could
17 explore this hypothesis: Is there any pain perception before
18 that.
19 "Q. But these studies have not to date been conducted?
20 "A. No, they haven't."
21 Were you asked those questions and did you give those
22 answers?
23 A. Yes, I did.
24 Q. It is true, isn't it, that there are fairly huge technical
25 difficulties associated with attempting to perform such

1 studies?
2 A. That fact.
3 Q. You talked during your direct examination about the
4 anatomical structures in the fetus. Do you recall that?
5 A. That's right.
6 Q. Is it true that the anatomical structures in the fetus
7 continue to develop after 20 weeks' gestation?
8 A. They certainly do.
9 Q. You talked about the cortex during your direct examination.
10 Do you recall that?
11 A. Certainly.
12 Q. Isn't it true that the cortex continues to develop after 20
13 weeks' gestation?
14 A. Certainly. And even after birth and during the entire life
15 of the individual the cortex continues to develop.
16 Q. You offered the opinion that a fetus would be subject to
17 intense pain from the abortion procedures described in the
18 Partial-Birth Abortion Ban Act of 2003. Do you recall that?
19 A. Yes, I do.
20 Q. You cannot state with a degree of certainty that all
21 partial-birth abortions will result in pain to the fetus, can
22 you?
23 A. I can state my opinion to a degree of medical certainty
24 that all fetuses beyond 20 weeks of gestational age will
25 experience severe pain by the partial-birth abortion procedure.

1 Q. And you have an 80 percent degree of certainty about that
2 opinion, is that right?
3 A. 80 percent or greater, counsel.
4 Q. You believe that the procedures covered by the Act can
5 include abortions earlier than 16 weeks, don't you?
6 MR. LANE: Objection, your Honor.
7 THE COURT: What is the objection?
8 MR. LANE: Beyond the scope of direct, beyond the
9 scope of this expert's opinion. It is not proffered on the Act
10 or interpretation of the Act or abortion procedures.
11 MS. WIGMORE: Your Honor, I disagree respectfully.
12 His opinion says that the procedures covered by the Act cause
13 pain to the fetus. So I think that I should be entitled to
14 inquire into this area.
15 THE COURT: Sustained.
16 Q. You cannot state with a reasonable degree of certainty that
17 fetuses experience pain before 20 weeks, is that right?
18 THE COURT: Sustained. Coming through the back door
19 what you weren't allowed to do in the front door, Ms. Wigmore,
20 doesn't work. Next question.
21 Q. Dr. Anand, your opinions regarding fetal pain focus on D&X,
22 is that right?
23 A. That is right.
24 Q. You told us earlier you have never performed a D&X, is that
25 right?

1 A. That is correct.
2 Q. But you would estimate that it takes somewhere between 5
3 and as a minutes, is that correct [> 5 and 15<]
4 A. Yes, that is correct. I had offered that opinion during my
5 deposition.
6 Q. The last two steps in your description of a D&X are
7 surgical incision and vacuum suctioning, is that right?
8 A. That is right.
9 Q. You would agree with me that those two steps will take a
10 short time, is that correct?
11 A. Yes. If I were to imagine myself doing a procedure, then
12 in my sort of estimation that would take a few minutes, yes.
13 Q. Once those steps are completed, the surgical incision and
14 the vacuum suctioning, the fetus will have no brain function,
15 is that correct?
16 A. That is correct.
17 Q. Isn't it true that an integral aspect of the experience of
18 pain is that it must be processed and registered in the brain?
19 A. That is correct.
20 Q. Wouldn't you agree that to the extent a fetus has the
21 ability to experience pain during an abortion procedure, the
22 D&X is not the only abortion procedure that would cause pain?
23 A. I have not offered an opinion about any other procedure.
24 Q. Doctor, didn't you testify on direct about dismemberment
25 D&E's?

1 A. That's right. I apologize. There are other procedures
2 that could cause pain to the fetus after 20 weeks of gestation.
3 Q. I'm sorry to interrupt you. Are you familiar with the
4 dismemberment D&E?
5 A. I am familiar with it to the extent that I have read about
6 the procedure. I have not performed any of those procedures.
7 Q. In a dismemberment D&E, it is your opinion, isn't it, that
8 at 20 weeks of gestation a fetus undergoing that procedure
9 would experience severe pain?
10 A. That is correct.
11 Q. Isn't it true, Doctor, that assuming the same gestational
12 age, a D&E procedure involving dismemberment would be more
13 painful to a fetus than a D&X procedure?
14 A. That is possible, yes.
15 Q. It is possible or it is a fact?
16 A. It is possible inso PHUFPS I have to use my imagination and
17 speculate as to what is involved in a D&E procedure as opposed
18 to an intact D&E procedure or a D&X procedure as you classify
19 it.
20 Q. Having previously testified under oath that a D&E involving
21 dismemberment would be more painful to a fetus than a D&X
22 procedure at 20 PWEBGS?
23 A. Could you refer me to the statement? [> check the
24 question<]
25 Q. Sure. Why don't we start with your ne ask a trial

1 transcript, which is also in the binder that I put before you.
2 I want to direct your attention to page 1081.
3 A. I have it.
4 Q. Do you have that page, Doctor? I want to call your
5 attention specifically to line 4, 1081 SRRB line 4.
6 "Q. And in fact isn't it true that it's your opinion that a
7 fetus undergoing a dismemberment procedure could experience
8 greater pain as a result of the fact that there is more somatic
9 injury?
10 "A. That is correct, yes."
11 Did you give that testimony, Doctor?
12 A. Yes, I did.
13 Q. If you could refer now to your deposition from this case,
14 page 118. I direct your attention to line 22 on that page.
15 "Q. Do you have any opinion as to whether in a D&E procedure
16 involving dismemberment, whether that's more or less painful to
17 a fetus than one which is aborted using the D&X procedure?
18 "A. Not having performed these procedures myself, I would
19 imagine that the D&E procedure, because it is associated with
20 much greater somatic injury" --
21 THE COURT: Ms. Wigmore, slow down, would you please?
22 MS. WIGMORE: I will read the question and answer
23 again.
24 Q. "Q. Do you have any opinion as to whether in a D&E
25 procedure involving dismemberment, whether that's more or less

1 painful to a fetus than one which is aborted using the D&X
2 procedure?
3 "A. Not having performed these procedures myself, I would
4 imagine that the D&E procedure, because it is associated with
5 much greater somatic injury to the fetus, would cause more pain
6 than the D&X procedure."
7 Did I read that correctly?
8 A. Yes, that is correct.
9 Q. Did you give that answer at your deposition?
10 A. I certainly did.
11 Q. You spoke on your direct examination about induction
12 abortions. Do you recall that?
13 A. Yes, I do.
14 Q. You haven't been involving in performing an induction
15 abortion, have you?
16 A. No, I have not.
17 Q. [> haven't been involved<]
18 Q. You have described the various steps of the D&X procedure
19 in your expert report, correct?
20 A. Yes, I have.
21 Q. The first three steps of that procedure as you have
22 described it are (a) grasping the lower extremity of the fetus
23 with a forceps or other surgical instrument, (b) manipulating
24 or rotating the fetal position within the uterus, and (c)
25 forcible extraction of the fetal legs and lower body through

1 the uterine cervix," is that correct?
2 A. That is correct.
3 Q. Those steps may occur in the induction abortion procedure,
4 correct?
5 A. Not having performed either of these procedures myself, I
6 am not aware of what the induction procedure would involve.
7 Q. Do you understand that the induction procedure involves
8 taking the fetus out of the woman's uterus?
9 A. Certainly, yes. My understanding is that prostaglandin PES
10 as a ris are inserted, which causes the onset of labor S-, and
11 the rotation of the fetus may or may not occur. The fetus may
12 be delivered head-first or by footling or breech in the
13 induction procedure. So part of those steps that I imagined
14 would occur in the D&X procedure probably would not occur in
15 the induction procedure.
16 Q. But there would be some extraction of the fetus in the
17 induction procedure, correct?
18 A. Certainly, yes. With the onset of labor, there will be the
19 expiration of the placenta and very likely fetal demise may
20 follow before the fetus is extracted.
21 Q. You testified with respect to the D&X that all steps of the
22 D&X would cause pain to the fetus, correct?
23 A. That is correct.
24 Q. To the extent there is extraction in the induction
25 abortion, that would also cause pain to a fetus at 20 weeks, is

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1 that your view?
2 THE COURT: You mean if the fetus is already dead as
3 well?
4 Q. No. I am asking you to assume that the fetus is still
5 living. If that is the case, would there be pain associated
6 with extracting the fetus from the uterus?
7 A. Yes. If placental expiration has not occurred and the
8 fetus is receiving adequate oxygen nutrition, etc., to keep it
9 alive, then as a result of the induction procedure there would
10 be pain associated to the fetus. Again, if the prostaglandins
11 have resulted in the dilation of the cervix, it may not be to
12 the same extent that dilation occurs in the D&X procedure.
13 Q. Do you have any knowledge as to how much dilation there is
14 in a D&X procedure?
15 A. I'm sorry, I am totally ignorant in this area.
16 Q. Let me ask you to assume that the fetus suffixiates in an
17 induction procedure. If that is the case, would there be any
18 pain with the asphyxiation, assuming the fetus is 20 weeks or
19 older?
20 A. I believe there would not be a feeling of pain associated
21 with asphyxiation.
22 Q. You testified, you were asked questions about whether there
23 was pain associated with birth at full term. Do you remember
24 that?
25 A. Yes, I do.

1 Q. You commented at birth at that point there have been some
2 inhibitory mechanisms that have been developed, was that your
3 opinion?
4 A. That is my opinion.
5 Q. Those inhibitory mechanisms in your opinion are not
6 developed as of 20 weeks, is that correct?
7 A. That is correct.
8 Q. So to the extent a fetus is alive and goes through the
9 process of induction [> check the second prior question<] at 20
10 weeks, that would result in pain to the fetus, wouldn't it?
11 A. If the fetus is alive, that's right.
12 Q. You also testified on direct that cutting the umbilical
13 cord would not be painful. Do you recall that?
14 A. I do.
15 Q. Can't cutting the umbilical cord result in a stress
16 response to the fetus?
17 A. The stress response probably would not occur because fetal
18 demise occurs before the hormonal or hemodynamic response can
19 be generated.
20 Q. Is it your view that the stress response would not occur if
21 the umbilical cord were cut?
22 A. True.
23 Q. Could you refer, please, to your Nebraska trial transcript,
24 page 1080. I direct your attention to line 1 on 1080.
25 "Q. And you were asked about whether cutting the umbilical cord

1 would cause pain to the fetus, and I believe your testimony was
2 probably not or very little, is that correct?
3 "A. That is correct.
4 "Q. Are you offering an opinion as to whether it's always
5 possible to cut the umbilical cord prior to removal of the
6 fetus during a surgical abortion?
7 "A. I'm simply offering an opinion on the degree of pain that
8 would be caused.
9 "Q. And a fetus in that situation would, however, likely mount
10 a stress response, is that correct?
11 "A. If that fetus were to survive long enough, yes, there would
12 be a stress response to the blood loss associated with cutting
13 the umbilical cord."
14 Were you asked those questions and did you give those
15 answers?
16 A. I certainly did.
17 THE COURT: Are you suggesting that is inconsistent?
18 MS. WIGMORE: Yes, I am, your Honor. He testified --
19 THE COURT: You are suggesting it?
20 MS. WIGMORE: I am.
21 THE COURT: The Court doesn't necessarily agree.
22 Q. To the extent there is a stress response associated with
23 cutting the umbilical cord, that is not evidence of pain, is
24 it?
25 A. Like I said, if the fetus were to survive enough to mount a

1 stress response, that stress response, if it could be measured,
2 would not be associated with pain but would be associated with
3 acute blood loss that has occurred.
4 Q. So no all stress responses are equated with pain, is that
5 correct?
6 A. That is correct.
7 Q. [> so not all<]
8 Q. I believe you testified on direct examination that to the
9 extent the fetus is able to feel pain, injecting a needle into
10 the fetus's heart will cause pain, is that correct?
11 A. That is correct.
12 Q. So an injection of digoxin or potassium chloride into the
13 fetal body will certainly cause pain to the fetus in your view,
14 is that right?
15 A. Yes, from the point of entry of the needle into the fetal
16 body to the point when fetal demise occurs as a result of
17 cardiac arrest.
18 Q. To the extent that the injection causes poisoning of the
19 fetus, could that cause pain to the fetus?
20 A. My opinion is that it would simply be the pain associated
21 with the needle puncture. I am not aware that the digoxin
22 injection or potassium chloride injection per se could cause
23 pain. If they were delivered into the fetal circulation
24 without the need for an invasive procedure SP-P.
25 Q. Just so I am clear, are you saying that you don't know one

1 way or the other whether the actual chemical potassium
2 SKHROEURD or digoxin would itself cause pain?
3 A. That is correct.
4 Q. Your conclusions on anatomical development do not hold true
5 for fetuses with brain anomalies, is that right?
6 A. That is right. Those fetuses have not been scud studied to
7 any great extent.
8 Q. And you can't extrapolate your findings about pain to
9 fetuses with brain anomalies, is that right?
10 A. Yes, that is correct.
11 Q. So if a fetus had a condition known as anencephaly, that
12 fetus would not experience pain, is that right?
13 A. It would depend on the degree of anencephaly that the fetus
14 has. If there are fetuses that have an advanced development of
15 their thalamus, hippo campus, limbic system, there would be a
16 subcortical substrate for experiencing pain in those fetuses.
17 S- there are fetuses that are completely enas a TPALic where
18 there is not even the fusion of the spinal cord or the brain
19 stem. In those fetuses I presume there would not be any
20 perception because the thalamus has not developed.
21 Q. So you cannot testify with a reasonable degree of certainty
22 that fetuses with brain anomalies at 20 weeks will experience
23 pain, is that right?
24 A. I do agree with you.
25 Q. You personally have not conducted any studies measuring the

1 effects of anesthesia on fetuses, is that right?
2 A. That is correct.
3 Q. You do not personally administer anesthesia to pregnant
4 women, is that right?
5 A. No, I don't, although I serve as a reviewer for several
6 journals when these articles are submitted for publication.
7 Q. Perhaps I confused. I was speaking of your experience in
8 actually administering anesthesia. You don't do that to
9 pregnant women, do you?
10 A. I don't.
11 Q. There is a huge amount and variability in terms of how an
12 anesthetic is administered and the dosage amount, is that
13 right?
14 A. Yes. That depends on the practice patterns of the STHAOES
15 KWROLGT.
16 Q. You are not aware of any studies on anesthetic agents
17 administered to a woman during a D&X procedure, are you?
18 A. No, there have been no published studies as far as I am
19 aware.
20 Q. Isn't it true that there is limited information addressing
21 the delivery and efficacy of anesthesia to fetuses during the
22 second trimester?
23 A. That is correct. This is a relatively underinvestigated
24 area.
25 Q. And there have been no conclusive studies looking at the

1 amount of maternal anesthesia that would be needed to curb pain
2 in a fetus, isn't that right?
3 A. I beg to submit that those studies would be unethical,
4 because they would expose the mother to toxic doses of
5 anesthetic drugs. So no IRB committee would approve those
6 protocols.
7 Q. You say that those studies would expose the mother to toxic
8 amounts of drugs. But isn't it true that there has been no
9 study about the amount that is needed?
10 THE COURT: Your question suggests that doctors should
11 do something unethical in order to study it?
12 MS. WIGMORE: No. Actually I am just asking how old
13 one know how much SHAOERB is needed without having done the
14 study.
15 THE COURT: Rephrase the question.
16 Q. Doctor Anand, has any study been done to determine how much
17 anesthetic is needed to transmit anesthesia from the mother to
18 the fetus?
19 A. There are no direct studies of producing an anesthetic
20 fetal anesthetic state because of the toxicity that would
21 involve to the mother. S- however, anesthesia has been
22 administered to premature babies who are from 23 weeks of
23 gestation and upwards. And we do have some idea of the blood
24 levels that would be required to produce a state of anesthesia
25 in those babies. So we would extrapolate from premature

1 infacts of the same neurological maturity to fetuses of that
2 neurological maturity about the production of a fetal
3 anesthetic state.
4 Q. You have already testified that there are physiological
5 differences between preterm neonates and fetuses, is that
6 right?
7 A. That is correct, although there are little, if any,
8 neurobiological differences between fetuses and preterm
9 neonates of the same neurological maturity.
10 Q. Doctor Anand, you would agree, wouldn't you, that there is
11 a huge gap in our knowledge regarding the effects of maternal
12 anesthesia on the fetus in the second trimester of pregnancy?
13 A. That is correct.
14 Q. In forming your opinions regarding the effects of
15 anesthesia, you assumed that methods of local anesthesia are
16 routinely applied for D&X, didn't you?
17 A. Yes, that is my estimation.
18 Q. You don't actually administer anesthesia for D&X, right?
19 A. No, I don't.
20 Q. You don't observe as anesthesia is administered for D&X,
21 correct?
22 A. That is correct.
23 Q. And you haven't read any studies or articles about what
24 type of anesthesia is administered for D&X, have you?
25 A. There are none in the literature that I could locate.

1 Q. So you don't have a basis for knowing what methods of
2 anesthesia are routinely applied for D&X, do you?
3 A. To the extent that one relies on discussions with
4 colleagues who are in the anesthesia department and summarizes
5 their opinion, that is how my impression was formed.
6 Q. In forming your opinions, did you talk to doctors who
7 perform D&X?
8 A. I haven't spoken with any doctors who performed a D&X, but
9 I have spoken with STHAOES KWROLGTS who are practicing
10 obstetric anesthesia, and I have also read through some of the
11 testimony that was heard by Congress several years ago, as well
12 as read articles on the technique of anesthesia used for
13 termination of pregnancy, the studies that are published,
14 although they don't pertain to the D&X procedure, they pertain
15 to the induction of labor procedure.
16 So from all those sources of information, I feel --
17 and I am not an expert in this area -- I feel that there may be
18 a lot of procedures that are performed using local or epidural
19 anesthesia.
20 THE COURT: Doctor, have you already read some of the
21 expert statements of plaintiffs and their experts in this case?
22 THE WITNESS: I have not seen any expert testimony,
23 your Honor.
24 THE COURT: Or expert statements?
25 THE WITNESS: There was one expert statement which was

1 a rebuttal statement that was sent to me. It was about half a
2 page in length, back in the second week of February. That was
3 all I was able to read. That was a statement from Dr. Creinin,
4 I believe.
5 THE COURT: All right.
6 Q. Doctor, would it surprise you to learn that there are
7 doctors who use general anesthesia for D&X?
8 A. No, it would not surprise me.
9 Q. General anesthetics readily cross the placental barrier and
10 fetal blood-brain barrier, isn't that right?
11 A. It depends on what drugs are being used for general
12 anesthesia.
13 Q. I want to direct your attention, Doctor, to page 10 of your
14 expert report, which is in the binder before you.
15 A. Which tab is that?
16 Q. It should be the very first document in the testimony
17 binder. Is it there?
18 A. I have it here.
19 Q. I want to refer you specifically on page 10 to the fourth
20 sentence. Did you say in your expert report that general
21 anesthetics (inhalational anesthetics and certain opiates such
22 as if I understand it nil and SUF if I understand it nil) can
23 provide some degree of pain relief to the fetus because they
24 readily cross the placental barrier and fetal blood-brain
25 barrier"? Is that a statement that you made?

1 A. Yes.
2 Q. Isn't it true that during open fetal surgery under maternal
3 general anesthesia, inhalation agents are considered to provide
4 adequate fetal anesthesia?
5 A. Where are you referring to?
6 Q. I am not referring to anywhere particularly right now. Is
7 that a true statement or not a true statement?
8 A. I don't perform fetal surgery, so I am not qualified to
9 comment on that.
10 Q. So you don't know one way or the other whether general
11 anesthesia delivered to the mother will affect the fetus?
12 A. I believe there are inhalation an anesthetics. These are
13 the anesthetic gases such as hal oh that I know,
14 isofluoroAEUPB, ceive oh fluoroAEUPB and similar anesthetic
15 gases, would E qui liberate fairly quickly across the something
16 barrier and would produce some level of anesthesia in the fetus
17 S- [> placental barrier?<]
18 Q. Doctor, do you recall testifying on direct examination that
19 the article by Fisk had a particular impact on your opinion?
20 A. Yes, I do.
21 Q. If you could turn, please -- actually, it may be easier to
22 use the binder that Mr. Lane gave you. This is Government
23 Exhibit G8.
24 A. I don't have that binder.
25 Q. I will pass it up to you.

1 MS. WIGMORE: Your Honor, may I approach?
2 THE COURT: You may.
3 Q. Again, it is tab G1.
4 A. Yes, I have it.
5 Q. This is an article that had impact on your opinions, is
6 that correct?
7 A. It certainly did.
8 Q. Could you refer, please, to page 14 of that article. I
9 want to direct your attention to the third paragraph on that
10 page, first sentence, which says, "Although the human fetus in
11 the last half of gestation has the necessary neuro connections
12 for no sus ception, it is not known whether the human fetus
13 experiences pain." That is a statement that you relied on in
14 forming your opinions, isn't it?
15 A. I certainly did.
16 Q. If you could refer to the last sentence of that same
17 paragraph, which states, "Because the relation between stress
18 responses and pain is not clear, it is not possible from our
19 data to conclude that the human fetus experiences pain in
20 utero." Did I read that correctly?
21 A. Yes, you did.
22 Q. That is part of the article that had a significant impact
23 on your opinions in this case, Doctor?
24 A. It certainly did.
25 Q. You testified at your deposition that the MRC report, the

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1 expert group on fetal pain, was another document that supported
2 or confirmed your opinion, is that right?
3 A. Yes, that is correct.
4 Q. If you would turn to the very last tab of your binder, the
5 article binder, I believe you will find that article.
6 A. Yes, I have it.
7 Q. I want to direct your attention specifically to page 4,
8 section 4.1.
9 A. Yes.
10 Q. Specifically, to the third and fourth sentences of that
11 section, which begins, "Much public concern." Do you have
12 that?
13 A. Yes.
14 Q. Those sentences read, "Much public concern has arisen
15 because of the possibility that a fetus may feel pain. Such a
16 possibility is founded on the existence of reflex movements and
17 neural activity produced by sensory stimulation. Despite the
18 usefulness of such reflexes in understanding spinal cord and
19 PRAEUPB stem pain processing, it would be a mistake to equate
20 them with true pain experience, which must involve the cortex
21 and develop post natally looping with memory, anxiety, and
22 other cognitive brain functions." That is a statement that was
23 in another article you relied op in forming your opinions in
24 this case, Doctor?
25 A. This was not part of the articles I relied on in making my

1 expert report. It was offered to me on the day before my
2 deposition, and I actually read it on the day of the
3 deposition.
4 Q. You testified, didn't you, that this document confirms your
5 opinion?
6 A. It certainly does. The conclusions of this MRC working
7 group were very similar to the conclusions I have stated in my
8 expert report.
9 Q. Let's turn to the conclusions, Doctor. If you could turn
10 to page 11 of the article, at the top of that page there is a
11 heading that says "9. Conclusions.?"
12 A. Yes, I have it.
13 Q. In the second sentence, that reads, "the basic molecular
14 and cell ular mechanisms of fetal and neonatal pain are still
15 poorly understood as are the effects of anesthetics or
16 analgesics." Did I read that correctly?
17 A. That is correct.
18 Q. And that is consistent with the opinions you have to offer
19 here today?
20 A. Yes, to some extent.
21 MS. WIGMORE: Thank you, Doctor. I have no further
22 questions at this time.
23 THE COURT: Mr. Lane, do you have any redirect?
24 MR. LANE: Yes, just briefly, your Honor.
25 REDIRECT EXAMINATION

1
2 BY MR. LANE:
3 Q. Good afternoon, Dr. Anand.
4 A. Good afternoon, counsel.
5 Q. During the course of your cross-examination you had made a
6 statement that consciousness is not an all or nothing thing.
7 Could you explain what you meant by that.
8 A. Certainly. The prevalent notion in the medical community
9 is that consciousness is not an all or none phenomenon. It is
10 not like a switch that is off one minute and on the next. It
11 is more like a dimmer, where there is a gradual increase in the
12 levels of consciousness.
13 Perhaps to use a common metaphor, when one arises from
14 a deep, dreamless state of sleep, there are stages of
15 TKROUSiness coming to full consciousness. That is how it is
16 believed that KOPGSness develops in utero, that early levels of
17 consciousness gradually increase to the level of consciousness
18 that can be noted in fetuses.
19 Q. Doctor, I would like to turn your attention to what was
20 called your witness binder and your deposition testimony on
21 page 42.
22 A. Sir, 42 of the deposition?
23 Q. Page 42 of your deposition, yes. During your
24 cross-examination, lines 6 through 8 were read in of your
25 deposition, saying "It was fair to say you weren't an STHAOES

1 KWROLGT." For purposes of completeness, I would like you to
2 read in lines 9 through 18 of your deposition, the question and
3 answer, if you could.

4 A. The question is, "Have you ever acted as an STHAOES
5 KWROLGT?"

6 The answer is, "In the course of my practice as a
7 pediatric intensivist, there are lines of activity that would
8 classify what I have done as acting as an STHAOES KWROLGT. So
9 there are procedures performed on infants and children where I
10 have provided anesthesia, and in that sense you could say that
11 I'm acting as an STHAOES KWROLGT. But those functions that are
12 common to the fields of STHAOESiology and critical care
13 medicine."

14 Q. Thank you, Doctor.

15 Doctor, you were asked about a number of articles
16 during your cross-examination. I am not going to make you
17 revisit all of them, but I would like to ask you a few
18 questions. The first was about an article found at tab 5.

19 A. I have it.

20 Q. You were read in cross-examination a portion of that
21 article talking about birth as a trigger to consciousness. I
22 would like to ask you if you could provide a context for that
23 statement in the course of this article of which you are an
24 author.

25 A. Certainly. The style of writing in a number of these

1 articles is to first state the opposite view and then to
2 examine that view in the light of the evidence that has been
3 generated. In this article where we discuss the development of
4 consciousness in the term neonate, in the preterm neonate, and
5 in the fetus, at one point we say that perhaps an argument
6 could be that the process of birth serves as a trigger for
7 consciousness.

8 However, if consciousness is triggered by birth, there
9 is insufficient data to recommend that that is actually the
10 case. And we go through those lists of arguments within this
11 article.

12 We finally conclude in this article, as I have in my
13 report submitted, that consciousness likely develops around mid
14 gestation in the human fetus.

15 Q. Doctor, this methodology of stating the opposite conclusion
16 and then looking at evidence, does this apply to any of the
17 other articles that were mentioned on cross-examination?

18 A. It certainly does. Medical writers will frequently state
19 the opposite viewpoint and then plug away at it until they are
20 able to TREPBT hypothesis. That is integral to the scientific
21 method.

22 Q. Doctor, during cross-examination it was discussed that you
23 had not recently studied the issue of fetal pain in utero.
24 Have you done any recent work on fetal pain with premature
25 infants and neonates?

1 A. I have worked extensively on the pain experienced by
2 premature infants.
3 Q. What gestational ages have you worked with?
4 A. The smallest preterm infant was of 22 or 23 weeks [>
5 invent<]
6 Q. Doctor, finally I would like to ask you a question about
7 this MRC report which was just discussed. You had said it, in
8 your view, confirms your views and opinions in this case.
9 Could you explain that for the Court. That is found at the
10 last tab I believe of the article binder.
11 A. Certainly. This was a view that was put together in August
12 of 2001. Like some of the other evidence that has been
13 examined was relevant to the information available in 2001.
14 And this MRC expert group was charged with the task of defining
15 areas that the MRC, which is the medical research council,
16 should invest in the future to advance this field. The
17 conclusions of this task force were very similar to the
18 conclusions that I have made. They relied on the same
19 neuroanatomical and neurophysiological maturations. They came
20 up with very similar conclusions in terms of the effects of
21 analgesic drugs on stress responses in animals and human
22 fetuses, and they put together the potential areas for future
23 research. Many of those areas since then have been
24 investigated, and data since 2001 has been available. As you
25 might recall, the article by Nicholas Fisk and colleagues was

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1 published after this report came out. And there are other
2 articles that have been published since then.
3 MR. LANE: No further questions, your Honor.
4 MS. WIGMORE: Just a few.
5 THE COURT: Any recross, Ms. Wigmore?
6 MS. WIGMORE: Just a few questions, your Honor.
7 THE COURT: Go right ahead.
8 RECCROSS-EXAMINATION
9
10 BY MS. WIGMORE:
11 Q. Dr. Anand, Mr. Lane asked you about your article at tab 5
12 in the articles binder. Isn't it true that you stated in that
13 article that it may be impossible to obtain unequivocal
14 evidence for fetal consciousness?
15 THE COURT: May I have that question read, please.
16 (Question read)
17 THE COURT: I don't think it is his article, is it?
18 MS. WIGMORE: Yes, your Honor.
19 THE COURT: It was authored by the witness?
20 MS. WIGMORE: That's correct.
21 THE COURT: All right.
22 THE WITNESS: Yes, I am the first author on this
23 article, your Honor.
24 THE COURT: All right.
25 A. You are absolutely right, counsel. The consciousness is

1 the substratum to all experience. Consciousness, by
2 definition, is beyond thought. In order to measure something
3 beyond thought is impossible, even in the adult human. So how
4 can I tell that the people in this room are conscious? From
5 their behavior. If their eyes are opened, if they are
6 occasionally moving if they show understanding on their face, if
7 they have any emotional reactions to the proceedings in this
8 room, I would surmise that they are conscious. But can I
9 measure their consciousness? I can't. Can they measure their
10 consciousness? No, they can't.

11 It can be experienced. It is like silence. Can you
12 measure silence? It can't be measured but it can be
13 experienced. And that is what applies to the human fetus as
14 well.

15 Q. Doctor Anand, the people in this room can tell you about
16 their feelings, can't they?

17 A. They certainly can. And if a fetus could talk, they could
18 tell us about their feelings.

19 Q. Doctor, in that same article, tab 5, if you could refer to
20 page 68, the bottom paragraph in the first column, I want to
21 refer you to the last two sentences, which begin, "All the
22 lines of evidence reviewed above suggest the presence of
23 consciousness from about 20 to 22 weeks of fetal life.
24 Accumulating data may confirm or refute these tentative
25 projections, although current practice should incorporate the

1 use of effective pain management for all premature and
2 full-term new PWORPBS." Did I read that correctly?

3 A. Yes, you did.

4 Q. So in this article you referred to the presence of
5 consciousness from about 20 to 22 weeks as tentative
6 projections, is that right?

7 A. Yes. This article was submitted for publication in June of
8 1998 and was published in '99.

9 Q. You mentioned on your redirect the theory that many authors
10 will set forth contrary conclusions in their articles and then
11 come around and reject those. Did I get that right?

12 A. That is correct.

13 Q. Let's refer back briefly to a few of the articles that you
14 and I talked about, starting with tab 10. That is the article
15 by Modi and Glover for the chapter of the book by those
16 individuals?

17 A. That is correct [> or the<]

18 Q. Those individuals do not conclude anywhere in this article
19 that fetuses experience consciousness at 20 weeks, do they?

20 A. They certainly propose the likelihood of consciousness.
21 Again, this chapter was published in the year 2000, and the
22 evidence has accumulated since then. STKPWHRL TKR*L --

23 A. As it applies --

24 THE COURT: Please don't interrupt again, Ms. Wigmore.

25 You may finish your answer, Doctor.

1 THE WITNESS: Thank you, your Honor.
2 A. As I had stated for the review article that I had authored
3 in '99, those were tentative conclusions. But since then there
4 has been accumulating data and there is a lot more known about
5 fetal behavior and the various studies confirming previous
6 tentative conclusions.
7 Q. Thank you, Doctor. What I want to know is did these
8 authors in this document state that fetuses experience
9 consciousness at 20 weeks?
10 A. I would have to read the article, counsel. If you would
11 give me a minute.
12 Q. Sure.
13 A. May I read to you from this article?
14 Q. Sure.
15 A. This is on page 218, the right-hand side column. It is the
16 first paragraph from the top. "We do not know when, if at all,
17 consciousness begins during fetal life. Even in the adult, the
18 physical basis of consciousness is far from understood. Most
19 evidence suggests that consciousness is associated with
20 activity in the cerebral cortex. Greenfield has emphasized
21 that one should not think of consciousness as an all or none
22 phenomenon but rather like a dimmer switch. This concept of
23 evolving consciousness might well apply to the developing fetus
24 in whom conscious experience is both unlikely to be the same or
25 have the same physical basis as in the adult. Frogs, for

page 126

1 example, do not have a developed verbal cortex, lacking layers
2 4 to 6. If they are conscious at all, their experience may be
3 associated with activity in a much less complex neuronal
4 network, possibly more analogous to the fetal subplate zone."
5 Q. Are you finished, Doctor S-
6 A. I think that should suffice, yes.
7 Q. It is true, isn't it, that these authors concluded that we
8 do not know when, if at all, consciousness begins during fetal
9 life? Is that right?
10 A. That was the first sentence in that paragraph STKHR then if
11 you refer to page 219, second paragraph, last sentence, they
12 concluded, "This evidence supports clinical observations that
13 the no sus SEPT EUFR system is functional at 24 weeks but
14 provides little assistance in establishing the existence of
15 pain perception at earlier gestations," is that right.
16 A. That is correct. That was the state-of-the-art in the year
17 2000.
18 MS. WIGMORE: Thank you, Doctor. I have no further
19 questions.
20 THE COURT: Doctor, thank you very much. You may step
21 down.
22 (Witness excused)
23 THE COURT: Is the next portion of the government's
24 case the reading of the deposition transcript?
25 MS. GOWAN: No, it is not, your Honor. The government

1 proposes to call Laureen Tews.
2 THE COURT: Do you want to start? We are about ten
3 minutes away from our usual afternoon break. Which would you
4 prefer?
5 MS. GOWAN: At the convenience of the Court.
6 THE COURT: My, my. Wonderful. Why don't we get Ms.
7 Tews in here and get her started.
8 MS. GOWAN: Defendant calls Ms. Laureen Tews.
9 LAUREEN TEWS,
10 called as a witness by the defendant,
11 having been duly sworn, testified as follows:
12 THE CLERK: Please state and spell your full name
13 slowly for the record.
14 THE WITNESS: It is Lauren Ann Tews, LAUREEN ANN TEWS.
15 THE CLERK: Thank you. Please be seated.
16 THE COURT: Ma'am, how SPOU spell your last name?
17 THE WITNESS: TEWS.
18 THE COURT: Thank you, ma'am. You may inquire.
19 DIRECT EXAMINATION S-
20
21 BY MS. GOWAN:
22 Q. Ms. Tews, where are you employed?
23 A. I work for the National Abortion Federation.
24 Q. How long have you worked for the National Abortion
25 Federation?

1 A. Since August of 1997.
2 Q. What is your title?
3 A. I'm the medical abortion initiative director.
4 Q. What are your duties and responsibilities as the medical
5 abortion initiative director?
6 A. I am responsible for directing our educational programming
7 for healthcare providers in terms of the safe and effective use
8 of early medical abortion.
9 Q. Were you designated by the National Abortion Federation in
10 accord with Rule 30(b)(6) of the Federal Rules of Civil
11 Procedure to testify in this case?
12 A. Yes, I was.
13 Q. Were the areas that you were designated to testify about
14 the purpose of and the development of content for NAF policies
15 concerning NAF annual meetings, NAF risk management meetings,
16 conferences, and seminars?
17 A. Yes.
18 Q. Were you also designated by NAF to testify concerning NAF
19 policies about medical education programs that are presented,
20 supported, sponsored, endorsed by or in any way affiliated with
21 NAF?
22 A. Yes.
23 Q. Were you also designated to testify about NAF's creation
24 and maintenance of documents, audiotapes, videotapes, and other
25 information in recorded form relating to the presentation by

1 Dr. Martin Haskell entitled "Dilation and extraction for late
2 second trimester abortions" at the NAF risk management seminar
3 in September of 1992?
4 A. Yes.
5 Q. Were you also designated by NAF to testify in connection
6 with that NAF risk management seminar, the recording on
7 audiotape and/or videotape of Dr. Martin Haskell's
8 presentation, including identification of the voice of Dr.
9 Martin Haskell?
10 A. Yes.
11 (Continued on next page) 4/13/04 Judge Casey direct of Law reason
TAO*Z, by Ms. Gowan.
12 BY MS. GOWAN:
13 Q. And did you testify on those subjects at a deposition in
14 this case?
15 A. Yes, I did.
16 Q. And were you authorized to do so?
17 A. Yes.
18 Q. Is NAF accredited to provide continuing medical education
19 by the accreditation counsel council to continue medical
20 education?
21 A. Yes.
22 Q. Is that counsel known as A C C M E?
23 A. Yes, it is.
24 Q. When was NAF accredited by A C C M E to provide continuing
25 medical education?

1 A. In June of 1981.
2 Q. Has it held its accreditation status since that time?
3 A. Yes, it has.
4 Q. As far as you are aware, are physician's required to
5 accumulate a certain amount of continuing medical education in
6 order to satisfy their state licensure requirements?
7 A. Yes. It's dependent on the state.
8 Q. And does NAF provide physicians with continuing medical
9 education credit if the physicians attend NAF-sponsored annual
10 meetings?
11 A. Yes.
12 Q. And does NAF provide physicians with continuing medical
13 education credit if the physicians attend risk management
14 meetings?
15 A. Yes.
16 Q. Is that would be NAF risk management meetings?
17 A. Yes.
18 Q. And does NAF distribute certificates after ten days to
19 attendees upon request to verify attendance, that I remember
20 ten days for purposes of obtaining credit?
21 THE FOREPERSON: They turn in to us an evaluation
22 form, yes.
23 Q. And is NAF required by the ACC M E to keep records relating
24 to the meetings for which credit is given?
25 A. Yes.

1 Q. And as part of its aFred I TAEUGS, is NAF required to have
2 a committee that oversees the general medical education
3 activities that NAF provides?
4 A. The ACCME requires that there be an -- that we report on
5 the organizational framework in which our CME program exists
6 and at NAF we do have a medical education committee of our
7 board of directors.
8 Q. And does that committee oversee the general medical
9 education ABG TEUFPTS that NAF provides?
10 A. Yes.
11 Q. And does NAF hold risk management meetings or seminars?
12 A. Yes.
13 Q. And at least since the mid-1990s are those meetings held
14 once a year?
15 A. Yes.
16 Q. Prior to that time were they held in two smaller meetings
17 each year?
18 A. I believe I recall that from the documents that I reviewed
19 in preparation for my deposition.
20 Q. And these meetings are held at various locations within the
21 United States and Canada, correct?
22 A. Yes, that's correct.
23 Q. And the meetings are attended by NAF members, physicians,
24 nurses, nurse practitioners, clinical staff and administrators,
25 right?

1 A. Yes.
2 Q. And are the meetings planned by the NAF medical education
3 committee?
4 A. The planning process begins during the medical education
5 committee meetings.
6 Q. And does the NAF medical education committee discuss
7 various topics for the purposes of determining what might be
8 presented at a risk management meeting?
9 A. Yes.
10 Q. Does the medical education committee responsible for
11 setting the overall goals for the medical education program
12 that NAF sponsors for its members?
13 A. Yes.
14 Q. Are those goals then implemented by NAF staff members who
15 help plan the risk management meetings, the chair of each
16 particular program and the factual TEU for the program?
17 A. Right, that combination of folks help to implement the
18 manning of the risk management meetings.
19 Q. And does the factual TEU for the program include the
20 presenters of various papers or presentations at any given risk
21 management meeting?
22 A. Yes.
23 Q. And do the faculty members or presenters sometimes provide
24 to NAF the material that the factual TEU member would be
25 presenting in advance of the meeting?

1 A. Sometimes that's the case.
2 Q. And when the faculty members provide such information to
3 NAF, does NAF then compile that information in a syllabus for
4 distribution at the meeting?
5 A. Yes. The faculty submit occasionally submit certain
6 material for us to include in the syllabus, which is a
7 compilation of faculty material.
8 Q. And then you bind that material into one syllabus, correct?
9 A. Yes, that's correct.
10 Q. And is that then Xeroxed and paid for by NAF?
11 A. Yes.
12 Q. And then those materials are sent for distribution at the
13 meeting, is that correct?
14 A. Yes, that's correct.
15 Q. Distribution to the attendees?
16 A. Exactly.
17 Q. And from time to time NAF pays faculty members, as is
18 typical at many medical meetings and hon or rarea for
19 participation in the meeting, correct?
20 A. That's not the typical practice but we do occasionally
21 provide an honoraria for certain faculty from from time to
22 time.
23 A. Correct.
24 Q. From time to time does NAF pay for the hotel charges for
25 the faculty members who attend the risk management meeting?

1 A. Again, that's not typical but from time to time we do
2 occasionally do that for certain TPABG um TEU.
3 Q. And NAF charges a registration fee for its members had they
4 attend risk management meetings, correct?
5 A. Yes.
6 Q. For a two-day meeting currently that fee might be in the
7 REUPBth of \$300 to \$ 500 is that right?
8 A. That's the estimate that I gave during my deposition.
9 Q. And T-GS not unusual, is it for NAF to offer a faculty
10 member free registration for the day of the meeting at which
11 the faculty member is speaking, is that right?
12 A. Yes, that's correct.
13 Q. And at the risk management meeting NAF staff are onsite to
14 facilitate the running of the meeting, correct?
15 A. Yes, a group of staff members would be.
16 Q. Assist by staffing registration desk, checking in the
17 REPBLGS STRAPBTS, helping the faculty with audiovisual problems
18 and answering logistical questions and the like, right?
19 A. Yes.
20 Q. And the purpose of the risk management meetings is to meet
21 the continuing educational needs of NAF members, is that right?
22 A. Yes.
23 Q. And the focus of the risk management meetings is clinical
24 in nature, right?
25 A. Generally there is a clinical focus track. Occasionally

1 there might be another track that is not clinically focused.
2 Q. And at the clinically focused track there is information
3 that's presented about and discussed concerning medical issues,
4 is that right?
5 A. Yes.
6 Q. And is it fair to say that the various aspects of providing
7 abortion care are the general subject matter discussed at these
8 meetings?
9 A. That's correct.
10 Q. Now, NAF also holds annual meetings, correct?
11 A. Yes.
12 Q. And, again, one of the goals of the annual meetings is to
13 ensure that the educational needs of NAF members are met,
14 correct?
15 A. Yes.
16 Q. The annual meetings tend to be a little more comprehensive
17 than the risk management seminars, is that right?
18 A. Yes; there is more content.
19 Q. And continuing medical education credit is also given to
20 attendees who wish to attend the annual meetings, correct?
21 A. The medical track of the annual meeting is accredited for
22 CMES.
23 Q. Now, prior to 2000 it was typical for portions of the risk
24 management meetings and the annual meetings to be audio taped,
25 is that right?

1 A. For portions, yes.
2 Q. And prior to 2000 NAF would contract with outside local
3 vendors who would make the tapes of the presentations at the
4 risk management meetings and the annual meetings, correct?
5 A. Yes.
6 Q. As I understand the the meetings are no longer taped and
7 that's for security RBZ, is that right?
8 A. You could characterize it in general as security reasons.
9 Q. But prior to 2000 aspects of the meetings were taped on a
10 regular basis by NAF, correct?
11 A. Yes.
12 Q. And NAF's practice was to get bids from three vendors for
13 the taping project, is that right?
14 A. Our current practice when we work with vendors is to get
15 three bids, correct.
16 Q. And was that the practice prior to 2000, if you know?
17 A. I don't know that specifically. It wouldn't surprise me.
18 Q. And NAF's practice, prior to 2000, was to enter into
19 contracts that would specify vendor responsibilities and NAF
20 responsibilities, is that right?
21 A. With regard to the audio taping?
22 Q. Yes.
23 A. Yes.
24 Q. And under the contract the vendors were permitted to sell
25 the tapes onsite to attendees at the meetings, is that right?

1 A. That's my understanding.
2 Q. And NAF would receive copies of the tapes, correct?
3 A. Again, that's my understanding.
4 Q. And the tapes were available for order through NAF by
5 members, correct?
6 A. Yes.
7 Q. And NAF would charge the members who, subsequent to the
8 meetings, would order copies of the tapes, correct?
9 A. I believe so, yes.
10 Q. And NAF keeps a copy of the tapes of its meetings in the
11 NAF resource center in its offices in Washington, D.C.,
12 correct?
13 A. We have copies of some of the previous meetings' audio
14 tapes, those sets of audio tapes are not necessarily complete.
15 Q. And the copies that you do have, the sets are kept in a
16 case with a cover on the case that has a sleeve for each tape,
17 is that right?
18 A. The sleeve is -- the sleeves that I have seen relate to the
19 set of tapes, not to each individual tape.
20 Q. And the sleeves identify the particular annual meeting or
21 seminar at which the underlying tape is related, is that right?
22 A. That's correct.
23 Q. For example, the cover of the sleeve in which a tape is
24 kept might say NAF annual meeting 1998, or NAF risk seminar
25 management 1992 and the like, correct?

1 A. Yes.
2 MS. GOWAN: Your Honor, may I approach the witness?
3 THE COURT: Surely.
4 BY MS. GOWAN:
5 Q. Ms. Tews, I'm showing you what was marked as Gover Government
6 Exhibit 66A at your deposition in this case, it is a document
7 entitled second trimester abortion from every angle, fall risk
8 management seminar, September 13th-14th, 1992, Dallas, Texas.
9 Presentations, bibliography and related materials.
10 What is the logo that appears at the bottom of the
11 first page of this document?
12 A. That is NAF logo.
13 Q. And is that the copyright symbol right below that logo?
14 A. It appears to be, yes.
15 Q. And is this the cover sheet for the syllabus from the fall
16 risk management program that was presented in Dallas in
17 September of 1992?
18 A. I believe it is.
19 Q. And if you take a look at the second page, could you tell
20 us what that is?
21 A. That's the table of contents.
22 Q. And if you count down six entries, could you tell us what
23 the paper is that's listed there in the table of contents?
24 A. Second trimester D&X, 20 weeks and beyond, Martin Haskell,
25 M. D., D&E for late second trimester abortion, Haskell, page

1 27.
2 Q. And then if you turn to the next page, is that the Haskell
3 article appearing on page 27?
4 A. Yes, it appears to be.
5 Q.
6 MS. GOWAN: Your Honor, I will represent to the Court
7 and I have discussed this with plaintiff's counsel, that
8 Government Exhibit 66A appears in the Congressional record for
9 the Partial-Birth Abortion Ban Act of 1995 hearing before the
10 committee on SKWRAOD issue reUnited States Senate 10 4th
11 Congress first session on HR-1833, November 17th, 1995. That
12 appears at pages 1 through 12 of the Congressional record.
13 May I approach, your Honor?
14 THE COURT: You may.
15 Q. Ms. Tews, I'm showing you an audio cassette that has been
16 marked as Government Exhibit E; could you please read for us
17 the title that's marked on that audio tape?
18 A. It's a second trimester sec TPHAOEBGZ D&E 14 to 19 weeks,
19 20 weeks and beyond. NAF-2/september 13-14th, 1992, Dallas,
20 Texas.
21 Q. Is there a Bates Number on that audio cassette?, a NAF
22 number?
23 A. On the backside, yes.
24 Q. Could you read that for us, please?
25 A. It says 1NAF 028907.

1 Q. And as far as you know, was that videotape produced to the
2 government by the national abortion federation in discovery in
3 this case?
4 A. This audio tape?
5 Q. Yes. Sorry. Thank you?
6 A. Because it has a NAF Bates label I'm assuming it was, yes.
7 Q. And did you listen to portions of the audio tape that we've
8 TPHARBGD as Government Exhibit E?
9 A. I listened to portions of an audio tape from this meeting
10 on this topic.
11 Q. And were the portions that you listened to relating to
12 Dr. Haskell's presentations?
13 A. Yes.
14 Q. And did you listen to the entire presentation made by
15 Dr. Haskell as reflected on that tape?
16 A. Yes, in preparation for my deposition.
17 Q. And do you know, Dr. Haskell?
18 A. Yes, I do.
19 Q. And have you ever spoken with him?
20 A. Yes.
21 Q. And is Dr. Haskell a NAF member?
22 A. Yes, he is.
23 Q. And was Dr. Haskell a NAF member at the time of the 1992
24 risk management seminar in Dallas, Texas?
25 A. I believe, he was.

1 Q. And when you listened to Government Exhibit E, did you
2 recognize Dr. Haskell's voice on the tape?
3 A. I did not.
4 Q. And in preparation for your 30B 6 testimony which called
5 for identification by NAF of the speakers on the tape, did you
6 take steps to determine whether Dr. Haskell's voice was on the
7 audio tape?
8 A. Yes, I did.
9 Q. And you spoke with colleagues who identified his voice,
10 correct?
11 A. That's correct.
12 Q. And as NAF's Rule 30B 6 witness you testified that
13 Dr. Haskell is a speaker whose voice is captured on the audio
14 tape, correct?
15 A. Yes, that's correct.
16 Q. And is that your testimony here today, Ms. Tews?
17 A. Yes, it is.
18 Q. And you were also able to identify Dr. Michael Burnhill as
19 the person who introduces Dr. Haskell at the meeting and whose
20 voice is also captured on the tape, correct?
21 A. Yes, I believe that's correct.
22 Q. Now this audio tape was made under contract between NAF and
23 an outside vendor, is that right?
24 A. Yes, it was.
25 Q. And does this audio tape fairly and accurately represent

1 the presentation that Dr. Haskell made at the September 13th
2 through 14th, 1992 risk management conference in Dallas, Texas?
3 A. I was not in attendance at that meeting but I have no
4 reason to believe that it doesn't accurately represent what
5 Dr. Haskell said at that meeting.
6 Q. And at the time in 1992, as the regular practice of NAF to
7 contract with a vendor to make tapes of presentations such as
8 Dr. Haskell's presentation at NAF risk management seminars,
9 correct?
10 A. I'm sorry, it was the regular practice of NAF to --
11 Q. It was the practice for NAF at the time to contract with
12 vendors to make tapes of the presentations of faculty members
13 such as Dr. Haskell who present papers at Avenue annual
14 seminars and meetings, right?
15 A. At least some of the presentations, correct.
16 Q. And the tape were kept in the course of NAF's business and
17 in its resource center, right?
18 A. That's where the tapes that we have currently are kept.
19 Q. And copies of the tapes would be provided to members, NAF
20 members on request for a charge, is that right?
21 A. Yes.
22 MS. GOWAN: May I approach, your Honor?
23 THE COURT: You may.
24 Q. Ms. Tews, I'm showing you what has been marked as
25 government's trial Exhibit K-1. If you would take a moment to

1 look at that.
2 This is a two-page document, is that the national
3 abortion federation logo at the top of the first page of this
4 document?
5 A. That is a logo that we've used in the past.
6 Q. And this is, document is dated June 18th, 1993; if you turn
7 to page 2 do you see it is signed by Barbara? And Barbara is
8 identified as Barbara RA*Dford, executive director?
9 A. Yes, I see that.
10 Q. Do you know Barbara? Barbara radford?
11 A. I don't know her penal TKPWHREU.
12 Q. Do you know whether she was the executive director of NAF
13 in or about June of 1993?
14 A. I'm assuming she was from the title underneath her name.
15 MS. GOWAN: Your Honor, the government would move into
16 evidence Government Exhibit K-1. There was no objection
17 interposed by plaintiffs in the pretrial order.
18 THE COURT: Any objection.
19 MS. PARKER: No objection, your Honor.
20 THE COURT: It will be received.
21 Q. Ms. Tews, could you please read for us the salutation of
22 this letter and the first two paragraphs? And I would urge you
23 to please read slowly.
24 A. Dear NAF member:
25 Many of you have called in recent weeks regarding the

1 distribution of an inflammatory anti abortion flyer depicting
2 graphic details of the dilation and extraction (D&X) abortion
3 procedure. We would like to provide you with some accurate
4 information on this procedure, as well as some guidelines in
5 discussing this and any other abortion procedure with the press
6 and public.
7 First of all, the D&X procedure was presented as part
8 of the NAF second trimester abortion from every angle risk
9 management seminar held in September of 1992. The surgical
10 method described differs from a classic dilation and evacuation
11 (D&E) procedure only in that it does not rely upon
12 dismemberment to remove the fetus. Rather, the surgeon grasps
13 and removes a nearly intact fetus through an adequately dilated
14 cervix. This is a small variation in a basic surgical
15 procedure that has been performed for over 15 years.
16 The workbooks and TAEUPDZ from this meeting are
17 available to all NAF members who would like additional
18 details."
19 MS. GOWAN: Your Honor, pursuant to Federal Rule of
20 evidence 803.6, the government moves the admission of
21 Government Exhibit E as a record or data compilation in tape
22 form of events at a regularly scheduled NAF meeting organized
23 for the purpose of providing continuing medical education as
24 NAF is accredited to do and was and is the practice of NAF to
25 hold.

1 The government asserts as an alternative ground for
2 the admission of this exhibit --
3 THE COURT: Go ahead, put it on the record.
4 MS. GOWAN: 801(d)(2)(B) as a party admission by
5 a deposition.
6 MS. PARKER: Your Honor, the plaintiffs object.
7 THE COURT: All right, you're overruled. It will be
8 received.
9 MS. PARKER: Your Honor, may I be heard?
10 THE COURT: Sure.
11 MS. PARKER: For my grounds?
12 This tape is a compilation, it's therefore hearsay
13 within hearsay. Even if the government has established the
14 first level of hearsay, these compilations contain statements
15 on them, embedded statements for which there needs to be a
16 separate hearsay exception and none is available.
17 Rule 101D 2 does not provide a basis. There is no
18 evidence that NAF adopted the statements made by outsiders on
19 these tapes or relied on them for any purpose and there is
20 therefore no separate basis for the embedded statements
21 contained in the compilation.
22 THE COURT: They were sold, were they not, provided at
23 a price to members who requested them?
24 MS. PARKER: Your Honor, they were sold but they
25 contained numerous statements on them and there has to be a

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1 separate hearsay basis for each statement contained on their
2 tape and none is available.
3 THE COURT: Overruled.
4 Is this an appropriate time, Ms. Gowan.
5 MS. GOWAN: Yes, your Honor.
6 THE COURT: To take our break?
7 MS. GOWAN: Yes.
8 THE COURT: All right.
9 (recess).
10 THE COURT: Ms. Gowan, you may inquire.
11 MS. GOWAN: Your Honor, at this time we would like to
12 play for the Court Government Exhibit E in evidence.
13 What I would like to do is just play the first few
14 seconds of the tape, which identifies the meeting and then fast
15 forward to Mr. , excuse me Dr. Haskell's presentation.
16 THE COURT: Do whatever you want. The other side,
17 plaintiffs wishes to play something I'm sure they well counter
18 your presentation, it seems to be the standard course here.
19 Go ahead.
20 (audiotape played).
21 MS. GOWAN: Your Honor, while we are using the counter
22 to get the space, I had the office, using its outside
23 contractor, prepare a transcript of the tape. I'm not going to
24 offer the transcript into evidence but I did want to provide a
25 copy to counsel to the extent they would like it as the tape is

1 being played, and also to the Court.
2 THE COURT: Fine. All the time and need in a jury
3 where you have one, in this case it is assistance to the Court.
4 If plaintiffs have any objection as to the accuracy of the
5 transcript they can object but I'm sure they're going to follow
6 along now.
7 MS. GOWAN: We are just about there. We will be
8 hearing the end of side one and then we will flip the tape and
9 hear considerable portion of side 2.
10 THE COURT: Very well.
11 (audiotape played).
12 MS. GOWAN: I think, your Honor, it will flip
13 automatically.
14 THE COURT: Very well.
15 MS. GOWAN: Your Honor, I apologize, it is just the
16 lead in here. It is coming. My expertise is not tech technology.
17 That's all right, the Court is technically challenged too.
18 MS. GOWAN: Then you have sympathy for me.
19 Here we go.
20 (audiotape played).
21 (continued on next page) TWM 4/13 take 6 Lawrence Tews on direct<]
22 MS. GOWAN: Thank you, your Honor. I have no further
23 questions for Ms. Tews.
24 MS. PARKER: Your Honor, just a couple .
25 CROSS-EXAMINATION S-

1
2 BY MS. PARKER:
3 Q. Good afternoon, Ms. Tews. Ms. Tews, at NAF presentations,
4 does NAF make an effort to present different views on the same
5 topic at its meetings?
6 A. Yes. It is typical that a range of techniques or
7 therapeutic options would be presented at our meetings.
8 Q. And do these opposing views ever appear on the NAF tapes of
9 the meetings?
10 A. Sure.
11 Q. Are and a half tapes with opposing views ever sold?
12 A. Yes.
13 Q. Who are the NAF tapes sold to?
14 A. Generally, they would have been sold to our members or
15 attendees of the meeting.
16 Q. Are they sold to the general public?
17 A. No, they are not.
18 Q. Do the views presented at the meetings by the faculty at
19 the meetings, reflect NAF's organizational views?
20 A. No, they do not.
21 Q. Does NAF authorize its members to speak on NAF's behalf at
22 NAF meetings?
23 A. No.
24 Q. You said that the medical education committee assists with
25 meeting planning when Ms. Gowan was asking you questions, is

1 that correct?
2 A. Yes.
3 Q. Does the medical education committee assist the faculty in
4 preparing their presentations for NAF meetings?
5 A. No. The faculty generally are preparing their
6 presentations in TKPEPBLT. We choose our faculty because they
7 are experts.
8 Q. Does the medical education committee assist in preparing
9 materials in advance of the presentations at the meetings?
10 A. No, they do not.
11 Q. Do faculties submit their materials to any NAF staff for
12 comment prior to presenting them at the NAF meetings?
13 A. Generally, no.
14 Q. Do the faculties submit their materials to the medical
15 education committee for comment prior to presenting them at NAF
16 meetings?
17 A. As far as I know, that's not the case.
18 THE COURT: Ma'am, you weren't here for this program
19 that was played on the audiotapes, so you don't know whether
20 that one was submitted in advance by Dr. Haskell or not?
21 THE WITNESS: Your Honor, in preparation for my
22 deposition, I reviewed a file of materials related to the
23 planning of that meeting, including correspondence with faculty
24 who were presenting at that particular meeting, and there was
25 no indication that there was advanced review, or there was not

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1 a copy.
2 THE COURT: You weren't there?
3 THE WITNESS: No, I was not. I reviewed the meeting
4 files.
5 THE COURT: So there was no way you could know one way
6 or another?
7 THE WITNESS: Again, your Honor, in preparation for my
8 deposition, I reviewed the files.
9 THE COURT: Ma'am, were you there?
10 THE WITNESS: I was not.
11 THE COURT: So you have no idea of whether he walked
12 in and gave them an advance outline or full text or the sound
13 or videotape, because you weren't there, right?
14 THE WITNESS: Again, your Honor, I reviewed the
15 files --
16 THE COURT: You read some correspondence. You keep
17 telling me that, ma'am. You weren't there, right?
18 THE WITNESS: I was not at the meeting in 1992, no, I
19 was not.
20 THE COURT: Were you working for NAF at the time?
21 THE WITNESS: No, I was not.
22 THE COURT: Go right ahead with your questioning.
23 Q. Ms. Tews, did you give a deposition in this case?
24 A. Yes, I did.
25 Q. Was that deposition conducted pursuant to Rule 30(b)(6) of

1 the Federal Rules of Civil Procedure?
2 A. Yes, it was.
3 Q. Are you familiar with a deposition notice that was served
4 by the government for your deposition?
5 A. Yes, I am.
6 Q. In conjunction with the deposition, did you have a duty to
7 prepare on the topics listed in that deposition notice?
8 A. Yes, I did.
9 Q. Was among those topics the typical procedures for NAF's in
10 meetings and the presentations made at those meetings?
11 A. Yes, it was.
12 Q. Can you describe for the Court what you did to prepare for
13 that deposition. -FPLGTS sure. In preparation for my
14 deposition, I reviewed documents related to our meeting
15 planning since the early 1990's, I spoke with current and
16 former colleagues, I listened to audiotapes that were outlined
17 in the notice of deposition. I spoke with my lawyers and also
18 watched a videotape.
19 Q. During the course of that preparation, did you learn any
20 evidence to indicate that the particular presentation by Dr.
21 Haskell was reviewed in advance by anyone on the NAF staff?
22 A. No, I did not.
23 Q. Did you find any evidence that the presentation by Dr.
24 Haskell was reviewed in advance by anyone on the NAF medical
25 education committee?

1 A. No, I did not.
2 Q. Is it typical for the NAF staff to review in advance
3 presentations made at NAF meetings?
4 A. No, that is not typical.
5 THE COURT: How could this witness testify, as she
6 wasn't there, what happened in 1992?
7 MS. PARKER: Your Honor, she was a Rule 60(b) 1K3W4R6
8 witness with a need to gather that information.
9 THE COURT: But you haven't established that she knew
10 anybody that was there.
11 MS. PARKER: Your Honor n conjunction with Rule
12 30(b) (6) --
13 THE COURT: I heard you, ma'am. I am blind, yes, but
14 not deaf. It still doesn't give her knowledge of a time that
15 she wasn't there. Go ahead.
16 Q. Ms. Tews, in the clinical context, as far as you are aware,
17 once a faculty person has been chosen to give a presentation,
18 has NAF ever refused to allow that presentation to go forward
19 at a meeting?
20 A. No, not that I am aware of.
21 Q. Does NAF check the accuracy of the views presented by
22 faculty at its meetings?
23 A. Again, we don't typically review the content of the
24 presentation of faculty members given ahead of time.
25 Q. Who is responsible for developing the content of the

1 faculty presenters' presentations at NAF meetings?
2 A. The faculty.
3 Q. You said that you sometimes pay honoraria to the faculty
4 who present at NAF meetings?
5 A. It is not typical, but we sometimes do.
6 Q. In 1992 what was NAF's practice with respect to honoraria?
7 A. It was very uncommon to provide honoraria for faculty, and
8 that was especially the case of TPH-F AF members who were
9 presenting at the meeting, members who would have been in
10 attendance at the meeting regardless of their faculty status.
11 Q. I'm sorry. Did you say it was common or uncommon?
12 A. It was uncommon.
13 Q. Do the same faculty present at every NAF meeting?
14 A. No, they do not.
15 Q. Are there some faculty who only make a single presentation
16 at a NAF meeting and then never present again?
17 A. Yes.
18 Q. Is Dr. Haskell an employee of NAF?
19 A. No, he is not.
20 Q. Was he an employee of NAF at the time he spoke at the 1992
21 risk management meeting?
22 A. No, he was not.
23 THE COURT: Did he get paid?
24 THE WITNESS: I don't know the answer to that
25 question, but it would be very atypical.

1 THE COURT: I didn't ask you that, ma'am. I asked you
2 a very simple question. You have talked a lot and your counsel
3 brought you through it about all the 30(b)(6) preparation you
4 made. The question is, did you go back and check to find out
5 whether or not he was paid?
6 THE WITNESS: I did review the records that we had in
7 preparation for my deposition --
8 THE COURT: Ma'am.
9 THE WITNESS: And I did not see any indication that he
10 was paid in those files.
11 THE COURT: Did you go through all the checks?
12 THE WITNESS: No, I didn't see any checks --
13 THE COURT: Did you go through the hotel bill to see
14 if it was paid?
15 THE WITNESS: No, I did not.
16 THE COURT: Next question.
17 Q. Other than presentations by faculty, what, if any, other
18 things are contained on the tapes of the NAF conferences?
19 A. The tapes that I have heard include questions and answers.
20 So, for instance, the attendees of the meeting generally have
21 an opportunity to ask questions of the faculty after a
22 presentation, and that might appear on an audiotape. You heard
23 on that audiotape that there is introductory material from the
24 facilitators of a panel.
25 Q. On the tape that we have just listened to, do you know

1 whether there were any questions and answers taped on that same
2 tape?
3 A. In preparation for my deposition, I did listen to that
4 audiotape, and I believe that there were questions that
5 followed Dr. Haskell's presentation.
6 Q. Were there any other presentations on that same tape?
7 A. Yes, I believe there were.
8 Q. On the tape we just listened to, it sounded as if Dr.
9 Haskell made reference to a videotape that he was showing at
10 the same time?
11 A. Yes, it did sound like that.
12 Q. Does NAF have that videotape?
13 A. No.
14 Q. Would it have been typical for NAF to review such a
15 videotape in advance of a prese
16 presentation?
17 A. No, not at all.
18 THE COURT: She wasn't there. You could only extend
19 this witness so far, ma'am.
20 MS. PARKER: I asked her whether it was typical.
21 THE COURT: She wasn't there. She wouldn't know.
22 What is typical now may be one thing, but she wasn't there.
23 MS. PARKER: Your Honor, given that the government has
24 introduced these as business records, the typical practice of
25 the organization --

1 THE COURT: But she can't testify to it.
2 MS. PARKER: -- it relevant. She testified on direct
3 to the practice of the organization for purposes of --
4 THE COURT: She went back and found files. She can't
5 testify as to the program if she wasn't in attendance.
6 MS. PARKER: Your Honor, he information formed the
7 basis for the government's admission of these tapes as business
8 records.
9 THE COURT: Records and files that were under her
10 control but not what practice was when she was not an employee
11 and wasn't present.
12 MS. PARKER: I believe on direct the government
13 elicited much about the practice of the organization.
14 THE COURT: Next question. I am not going tot argue
15 with you. I have ruled.
16 MS. PARKER: I have no further questions, your Honor.
17 THE COURT: Any redirect?
18 MS. GOWAN: Just very briefly, your Honor.
19 REDIRECT EXAMINATION
20
21 BY MS. GOWAN:
22 Q. Ms. Tews, the risk management meetings have initial
23 planning that occurs with the medical education committee
24 meetings with the board of directors, correct?
25 A. Yes, that's true.

1 Q. At those meetings, various topics are discussed by the
2 medical education committee about what might be an appropriate
3 topic to cover at a risk management meeting, correct?
4 A. Yes, that's correct.
5 Q. There was some discussion about who might be appropriate to
6 chair the risk management meeting, correct?
7 A. Yes.
8 Q. Then based on those conversations, the planning of the
9 meeting moves forward, correct?
10 A. Yes, that's correct.
11 Q. The majority of the members on the committee are
12 physicians, aren'tr they?
13 A. Yes, that is correct.
14 Q. They might also include members of the board who have
15 particular expertise or interest in education or medical
16 education, right?
17 A. Uh-huh.
18 Q. It is the general function of the medical education
19 committee to review evaluations from attendees that have been
20 submitted at prior annual meetings and risk management meetings
21 to generate topics that might be appropriate for subsequent
22 meetings, correct?
23 A. The medical education committee generally reviews summaries
24 of evaluations from previous meetings.
25 Q. That the for the purpose of helping to decide what topics

1 might be generated for the following meetings, correct?
2 A. That is one of the purposes for reviewing summary
3 evaluations.
4 Q. In addition to reviewing the evaluations from prior
5 meetings, the committee might review some past topics that have
6 been covered to assess if there are topics that have not been
7 covered that might be relevant for a number of years and that
8 might be renewed at NAF meetings, correct?
9 A. Yes, that's correct.
10 Q. The committee might be aware of research that is being done
11 or emerging technologies that would be appropriate topics for a
12 risk management meeting, correct?
13 A. Yes, that's correct.
14 Q. The committee then votes on a meeting top *EUG for the
15 seminar, correct?
16 A. As I discussed in my deposition, I am not sure if there is
17 a formal vote, but there is a decision that is made by the
18 medical education committee about moving forward with the
19 planning of the risk management meeting.
20 Q. Directing your attention to page 21/line1 of your
21 deposition in this case:
22 "Q. Does the -- does a committee then vote on a meeting topic?
23 "A. Does the med Ed committee then vote?
24 "Q. Yes.
25 "A. It's my discussion that there's some discussion whether

1 there is a formal show of hands. I'm not clear on that,
2 whether there is a formal vote.
3 "Q. Is it as you say the med Ed committee, are they the ones
4 that make a decision about what the topic for the risk
5 management meeting would be?
6 "A. Generally, I believe that's how the topics are determined."
7 Were you asked those questions and did you give those
8 answers at your deposition in this case?
9 A. Yes, I did.
10 Q. During the course of the discussion about the topics, there
11 would also be discussion about who would be appropriate to
12 chair that meeting, correct?
13 A. Yes.
14 Q. You reference some documents or correspondence that you
15 uncovered in connection with your 30(b)(6) preparation in this
16 case, and one of those was a letter from Dr. Martin Haskell to
17 Dr. Burnhill at NAF, correct?
18 A. I believe it was addressed to Dr. Burnhill, not at the NAF
19 address but at his professional address.
20 Q. Correct. And Dr. Burnhill was the one who introduced Dr.
21 Haskell at the seminar that we just listened to, isn't that
22 right?
23 A. Yes, that's correct.
24 Q. In that letter didn't Dr. Haskell outline what it is that
25 Dr. Haskell proposed to present at a NAF annual meeting or

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1 seminar?
2 A. Yes, that's correct.
3 Q. Didn't he outline that his proposal was to describe a
4 technique for late second trimester/early trimester abortions
5 intact extraction without induction?
6 A. I can't recall the specific wording of that letter, but
7 that is the general content, I believe.
8 Q. Wasn't the idea, as expressed by Dr. Haskell to Dr.
9 Burnhill, that the purpose would be to educate providers about
10 D&E techniques for late second trimester and third trimester
11 abortion?
12 A. Again, that is the general content that I remember.
13 Q. And relate Dr. Haskell's experience with the method and
14 describe the method and selection criteria?
15 A. That's right the general content that I recall.
16 Q. Didn't Dr. Haskell propose to present a case study, present
17 a video of intraoperative ultrasound techniques, propose to
18 speak on cervical preparation and operative sequence?
19 A. I can't recall if that was in the letter. It appears as if
20 you are reading it from the letter. And that is the general
21 content of the letter.
22 MS. GOWAN: Thank you. I have no further questions.
23 THE COURT: Any recross?
24 MS. PARKER: No, your Honor, nothing further.
25 THE COURT: Ma'am, you may step down.

1 (Witness excused)

2 THE COURT: According to my watch, it is about 6
3 minutes of 5:00. I guess we will recess for the day and
4 reconvene at 9:30 tomorrow.

5 Before we go, Ms. Gowan, I understand my clerks have
6 been talking to you Mr. Hut. Is it your understanding, which
7 surprised me, that you will only be addressing the stay? I
8 thought also there was a mandamus before the Second Circuit on
9 next Tuesday.

10 MS. GOWAN: Your Honor, yes, the mandamus petition
11 will be before the Second Circuit on Tuesday, as well as the
12 stay issue.

13 THE COURT: I thought they agreed to address, as the
14 Seventh Circuit did, the substance of this production respect.
15 Otherwise, we would be faced with a dilemma that I don't even
16 wish to entertain.

17 MS. GOWAN: Your Honor, that's right. The posture,
18 because of the nature of the Presbyterian appeal, is a little
19 unclear, and it is possible that the Second Circuit could still
20 have a merits briefing that would be different than a stay
21 briefing and a mandamus briefing.

22 Certainly it is the government's view that the
23 mandamus petition by New York Presbyterian Hospital was just a
24 merits appeal that had been repackaged, if you will, in the
25 form of a mandamus petition. And we did, in our opposition to

1 the mandamus petition, address the merits.

2 THE COURT: You did?

3 MS. GOWAN: We did.

4 THE COURT: All right. I can't imagine, although it
5 would never cease to amaze me, but I can't imagine
6 understanding that a trial was about to commence, that --

7 MS. GOWAN: We certainly did, your Honor. We put in
8 our full briefing on that. We also argued lack of appellate
9 jurisdiction in our papers.

10 THE COURT: All right. That is for the ladies and
11 gentlemen of the 17th floor.

12 MS. GOWAN: In our view, the decision by the ladies
13 and gentlemen of the 17th floor will certainly be tantamount to
14 a decision on the merits, and we certainly hope the circuit
15 will agree with that view, and we intend to urge that view on
16 the Court on April 20th.

17 THE COURT: All right. Might I ask, are we doing
18 transcripts in the morning? Not that I wish to pry, but just
19 serve a general idea. Or do we have a witness starting off
20 tomorrow?

21 MS. GOWAN: Your Honor, saved by the bell. ACOG
22 today, and we will start with a live witness tomorrow, and
23 hopefully we will have ACOG in the afternoon [> no ACOG today,
24 I guess<]

25 THE COURT: I suppose that all points of view, whether

1 it is best late in the day or early in the day. All right,
2 fine. That is what we are doing. We have a live witness in
3 the morning and the deposition will be later in the day.

4 MS. GOWAN: Thank you, your Honor.

5 THE COURT: I wish you all a lovely evening. Court
6 will stand in recess.

7 (Adjourned to 9:30 a.m., April 14, 2004)