

STATE OF OHIO

CUYAHOGA COUNTY COURT OF COMMON PLEAS

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MATTHEW CHASE WAGONER, et al.

vs.

MARK R. EVANS, M.D., et al.

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* No. 497179

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DEPOSITION of MARCUS C. HERMANSEN, M.D.,

Deposition taken at 22 Gregg Road, Nashua,

New Hampshire, on Tuesday, June 20, 2006,

commencing at 9:35 a.m.

Court Reporter:

Pamela Carle, CCR, RPR

New Hampshire CCR No. 98

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APPEARANCES

For the Plaintiff:

BECKER & MISHKIND CO., L.P.A.

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By: Michael F. Becker, Esq.

For the Defendants:

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By: John T. Bulloch, Esq.

George Moscarino, Esq.

I N D E X

WITNESS: MARCUS C. HERMANSEN, M.D.

EXAMINATION: PAGE
By Mr. Bulloch 4

EXHIBITS FOR IDENTIFICATION:

HERMANSEN	DESCRIPTION	PAGE
1	2/10/06 letter and timelines	13
2	Handwritten list	13
3	Curriculum vitae	13
4	Report	13
5	Article	86

1 MARCUS C. HERMANSEN, M.D.,
2 having been duly sworn,
3 was deposed and testified
4 as follows:

5 EXAMINATION

6 BY MR. BULLOCH:

7 Q. Doctor Hermansen, for the record, my
8 name is John Bullock. I represent Fairview
9 Hospital.

10 I know you've been deposed numerous
11 times in the past, but just as a reminder, the
12 court reporter cannot take down nods of the head,
13 so please verbalize your responses.

14 If you need a break at any time, we'll
15 be happy to break. Also, if you don't understand
16 one of my questions, just ask me to rephrase it,
17 and I'll be happy to try it again, okay?

18 A. Yes, sir.

19 Q. You were kind enough to share with me
20 your records which you have in this case which are
21 fairly substantial, but I just wanted to go over
22 and make sure that we have everything in front of
23 us.

1 There are several depositions you
2 looked at, including depositions of Doctor Saxena,
3 Doctor Alexander, Doctor Evangelista, and
4 Doctor Bachman, correct?

5 A. Correct.

6 Q. Any other depositions besides -- oh,
7 wait a minute. You also reviewed the discovery
8 deposition of Doctor Adler and Doctor Lilien,
9 correct?

10 A. Yes.

11 Q. And would you like to look --

12 A. There may be more.

13 Q. There are.

14 A. There are some on the original.

15 Q. There are some more. And then you
16 reviewed the depositions of Margo Wagoner, Debra
17 Hopp, who was a nurse, I believe, at Parma
18 Community Hospital, and Doctor Mark Evan, who used
19 to be a defendant in this case, correct?

20 A. Correct.

21 Q. Is that the sum of the depositions that
22 you reviewed, sir?

23 A. Yes.

1 Q. And I notice you had several x-ray
2 films, including some from Fairview, as well as
3 some from Parma Community Hospital, correct?

4 A. Correct.

5 Q. Any others that you've looked at?

6 A. No.

7 Q. Any subsequent head films or anything
8 like that?

9 A. No.

10 Q. Now, I understand -- if you could pass
11 over your binder -- that you have some subsequent
12 medical records. I assume -- can I have the
13 entire binder, sir?

14 I assume that if -- if you have the
15 records from North Carolina, if you reviewed
16 anything from any head imaging films it would have
17 been reports, correct?

18 A. Correct.

19 Q. And you did not see any of the films
20 from any of the facilities in North Carolina,
21 correct?

22 A. Correct.

23 Q. The records you reviewed are Parma

1 Hospital, Fairview Hospital?

2 A. Yes.

3 Q. Parma Hospital and Fairview Hospital,
4 correct?

5 A. Yes.

6 Q. Some records from Doctor Evans' office?

7 A. The very last tab.

8 Q. And the only thing that I see from
9 North Carolina is pretty limited, is that correct?

10 A. Yes.

11 Q. One includes an admission to New
12 Hanover Regional Medical Center on 8 -- and August
13 2nd, 2000, correct?

14 A. Yes.

15 Q. It looks like some very limited notes
16 from an admission to University of North Carolina
17 as well, correct?

18 A. Yes.

19 Q. I'm not going to bother counting these,
20 but it appears to be about 20 pages?

21 A. Yes.

22 Q. I'll give those back to you, too, sir.
23 Counsel for plaintiff also provided you with some

1 expert reports, correct?

2 A. Yes.

3 Q. One was from a Doctor Nelson, who is a
4 defense pediatric neuroradiologist, correct?

5 A. Yes.

6 Q. You received, and I assume reviewed,
7 the expert report of Doctor Robert Darnall?

8 A. Yes.

9 Q. I noted you made a couple of notes on
10 Doctor Darnall's report?

11 A. Yes.

12 Q. Doctor Darnall's section entitled
13 interpretation and comments he has a sentence, the
14 course of the symptoms was most consistent with
15 mild to moderate hyaline membrane disease, do you
16 recall that?

17 A. Yes.

18 Q. And you drew what looks to be an
19 unhappy face next to it?

20 A. Yes.

21 Q. And you have circled mild, correct?

22 A. Yes.

23 Q. And then Doctor Darnall has a sentence,

1 the pneumothorases were properly treated and the
2 infant gradually improved, consistent with a
3 typical course of HMD. And you have circled
4 typical course of HMD, correct?

5 A. Yes.

6 Q. And next to that you wrote not in the
7 1990s, correct?

8 A. Correct.

9 Q. Do you remember why you wrote not in
10 the 1990s, what it is that you disagree with
11 Doctor Darnall about?

12 A. Since the introduction of surfactant
13 therapy, that's no longer the typical course. This
14 is the typical course of what we used to see when
15 Doctor Darnall and I both began our careers, but
16 it's not the typical course today, nor was it the
17 typical course at the time Matthew Wagoner was
18 born.

19 Q. So I assume then that you're
20 disagreeing with Doctor Darnall that Matthew
21 followed a typical course?

22 A. Yes.

23 Q. Correct?

1 A. Yes.

2 Q. And then Doctor Darnall has a sentence
3 that reads, at no time during the course of this
4 infant's hospitalization was there any evidence of
5 severe respiratory compromise associated with
6 prolonged or severe lack of oxygen, and he has,
7 parens, hypoxia, and you have underlined prolonged
8 or severe lack of oxygen, hypoxia, and a big
9 question mark next to it, correct?

10 A. Yes.

11 Q. What are you questioning about
12 Doctor Darnall's opinion?

13 A. I'm questioning what his definition of
14 severe hypoxia is. This baby had low oxygen
15 saturations and low blood gas oxygen levels down
16 into the 30s for --

17 Q. Was it prolonged, or was it short
18 bursts of that?

19 A. He was sick pretty much for around five
20 or seven hours. He began getting sick at five
21 o'clock in the afternoon, I believe, and we have a
22 blood gas at midnight that looks bad, and he was
23 pretty sick through that entire period.

1 Q. Doctor, why don't you take the time to
2 look at the blood gasses right now and tell me
3 where you see a prolonged period of hypoxia.

4 A. I'm basing this on more than blood
5 gasses. It's also nursing observations at the
6 bedside.

7 Q. Okay. So tell me where you're seeing
8 prolonged periods of hypoxia?

9 A. This episode begins at 5:40 in the
10 afternoon. Now, my pages have little numbers
11 circled on them, if that helps.

12 MR. BECKER: They wouldn't have that.

13 THE WITNESS: Okay.

14 A. At 5:40 in the afternoon.

15 Q. Of what day, sir?

16 A. On the 25th, the second day of life.

17 Q. Okay.

18 A. The baby has persistent oxygen
19 saturations from 60 into the 70s. It's persistent
20 even after they tried to bag resuscitate the baby,
21 there's no improvement. So this is beginning at
22 5:40.

23 And I'm going to say that the child is

1 sick up until midnight when we have a blood gas.

2 That's six hours later, we have a blood gas with a
3 PO 2 value of 32. Well, to me those numbers seem
4 prolonged and severe. Evidently to Doctor Darnall,
5 it's not prolonged and severe.

6 Q. So the period that you have at issue
7 goes from 5:40 until midnight of August 25th, is
8 that what I'm understanding you're saying?

9 A. Yes.

10 Q. Okay, we'll revisit that. Doctor, you
11 also have a couple of articles here in your file?

12 A. One article and two abstracts.

13 Q. Okay, and we'll talk about those later,
14 too. You also have a Survanta package insert from
15 May of 2004, correct?

16 A. Correct.

17 Q. And obviously this would not have been
18 a package insert that was in existence at the time
19 of Matthew Wagoner's hospitalization, correct?

20 A. I don't know.

21 Q. Well, certainly this one isn't, because
22 it's dated May of 2004, correct? It might have
23 been the same, but certainly not the one that you

1 have in your file.

2 A. True.

3 MR. BULLOCH: Okay, I'd like to mark
4 this as Exhibit 1, plaintiff's Exhibit 1, please.

5 (Hermansen Exhibit No. 1 was marked for
identification.)

7 (Hermansen Exhibit No. 2 was marked for
identification.)

9 (Hermansen Exhibit No. 3 was marked for
identification.)

11 Q. Doctor, I've marked as exhibits three
12 documents. First is -- and I'll hand it to you
13 and ask you to return it to me -- but first is a
14 letter from Mr. Becker's office that is a prenatal
15 timeline that was prepared by a Judith Gonet,
16 G-O-N-E-T. I assume you received and reviewed
17 that?

18 A. I received it. I don't remember having
19 looked at it.

20 Q. Okay.

21 MR. BULLOCH: Mark this as Exhibit 4,
22 please.

23 (Hermansen Exhibit No. 4 was marked for

2 Q. Exhibit 2 is a handwritten sheet?

3 A. Yes.

4 Q. That's your notes, or what are these
5 from?

6 A. I produced that.

7 Q. Okay, was this at Mr. Becker's request,
8 or was this just something that you had put
9 together?

10 A. It's something I did after reading
11 Doctor Alder's deposition.

12 Q. Okay, did you --

13 A. He was asked a lot about the various
14 causes of cerebral palsy, and I have a publication
15 coming out any day now where I published this table
16 as my thoughts with causes of cerebral palsy.

17 Q. And that would be the one that you're
18 the guest editor of in the Clinics in
19 Perinatology?

20 A. Yes, I wrote one chapter in the book,
21 but I also wrote the preface. And in the preface I
22 give a table for the causes for cerebral palsy, and
23 that comes from the table.

1 Q. We'll get to that, I have a copy of it.

2 Exhibit 3 is your CV?

3 A. Yes.

4 Q. Current CV? And Exhibit 4 is your
5 report, correct? You can have that, I've got it.

6 A. Yes.

7 Q. Doctor, I'm going to hand you back your
8 CV as well so you can refer to that. I had a
9 version of your CV that was dated October 2004,
10 and you were kind enough to give me your current
11 CV this morning, so we don't have a copy of it,
12 but I'll do my best.

13 Have there been any additions since
14 this current CV that we've marked as an exhibit?

15 A. No.

16 Q. You're certified by the American Board
17 of Pediatrics, correct?

18 A. Yes.

19 Q. And you have a sub-board in
20 neonatal-perinatal medicine, correct?

21 A. Yes.

22 Q. And I assume that you do not need
23 recertification?

1 A. Correct.

2 Q. The academic affiliation that you state
3 is associate professor of pediatrics, correct?

4 A. I have two appointments. One in the
5 department of pediatrics, one in the department of
6 OB/GYN.

7 Q. Okay, and those are both with Dartmouth
8 Medical School, correct?

9 A. Correct.

10 Q. Now, is that in Nashua that you hold
11 those appointments, or are those back up in
12 Hanover?

13 A. The school is in Hanover and Lebanon,
14 but most of my teaching takes place in Nashua.

15 Q. Okay. And as such, is that more of a
16 clinical professorship?

17 A. No, it's a full professorship. There
18 are clinical appointments, but this is a full
19 faculty appointment.

20 Q. You lecture residents?

21 A. Yes.

22 Q. What topics do you typically lecture
23 residents on?

1 A. I have two types of lectures, one is I
2 give a series of lectures to the OB/GYN residents.
3 During their year with us they get one lecture a
4 month, so they get 12 a year, of which I give about
5 eight of the 12, and my partners give the other
6 four.

7 I expect each of my partners to give
8 one a year, and I take about eight. These are
9 newborn topics that are appropriate for OB
10 residents. For example, one lecture is on newborn
11 resuscitation. One is on common birth defects.

12 Q. Okay.

13 A. We both have to know and they can learn
14 from our expertise.

15 Q. Are those lectures or presentations
16 listed in your current CV?

17 A. No.

18 Q. Okay.

19 A. And then secondly, I give lectures at
20 Lebanon and throughout New England on behalf of
21 Dartmouth. But those are one here, one there.
22 They invite me to give a lecture at some conference
23 or some meeting, and I do it.

1 Q. Okay, are those at other satellite
2 facilities of Dartmouth-Hitchcock, or is that --

3 A. Sometimes, and sometimes they're at the
4 main hospital.

5 Q. Now, the lectures that you give are
6 predominantly to OB/GYN residents?

7 A. The ones I give here in Nashua, yes,
8 that's true.

9 Q. Do you have any residents in
10 neonatology at Nashua?

11 A. No.

12 Q. So you don't give any lectures here
13 obviously in neonatology?

14 A. I give about one a year to the
15 pediatric department and the medical staff.

16 Q. Do you provide neonatology lectures
17 anywhere else in the Dartmouth system?

18 A. Well, throughout New England. I gave a
19 lecture in Vermont. It was a Dartmouth-sponsored
20 conference, but it was in Vermont.

21 Q. What kind of topic is that that you
22 typically give presentations in neonatology?

23 A. That was on resuscitation issues. How

1 do you train hospital providers to become skilled
2 in resuscitation.

3 Q. I notice in your resume that it seems
4 like you have somewhat of a subspecialty interest
5 in risk management, is that fair?

6 A. I've done extensive medicolegal reviews
7 and testifying, and based upon that experience, I
8 put together a book on that topic.

9 Q. Do you consider yourself to have any
10 other subspecialty in neonatology?

11 A. I'm interested in what we'd call
12 epidemiology, looking at some of the numbers and
13 relationships of cause and effect situations, what
14 causes prematurity, what causes CP, that type of
15 thing.

16 Q. Similar to your upcoming publication in
17 Clinics in Perinatology?

18 A. Somewhat.

19 Q. That type of research? Okay. Doctor,
20 what percentage of time do you spend in the
21 clinical practice of medicine?

22 A. I'm at the hospital as a clinician and
23 to a lesser extent administrator of our program 40

1 to 45 hours a week.

2 Q. Okay.

3 A. I am scheduled to work three 12-hour
4 shifts a week where I'm the clinician in the
5 hospital.

6 We break every day into two 12-hour
7 shifts. I worked this past Sunday for 12 hours. I
8 work tomorrow night for 12 hours, and then I have
9 another 12 hours at the end of the week.

10 So 36 hours a week I'm the clinician in
11 the hospital, and I'm there for one reason or
12 another another five hours.

13 Q. Okay, and what percentage of your time
14 do you spend in teaching responsibilities? Is
15 that just limited to the lectures that you do,
16 eight a year, and then assorted ones throughout
17 the state?

18 A. That's the formal teaching. I teach at
19 the bedside almost every day, but I don't think
20 that's what you're asking about.

21 Q. Well, and if you're at the bedside, I
22 assume you count that as part of your clinical
23 practice, too, correct?

1 A. Commonly clinical, administrative and
2 teaching overlap a lot. You're commonly doing two
3 of those tasks at the same time.

4 Q. Sure.

5 A. So, yes, that's true. I'm teaching
6 during my clinical time.

7 Q. Doctor, you listed in your CV on page
8 3, at least the version that I have, your
9 professional societies that you're a member of,
10 which included American Academy of Pediatrics,
11 section on perinatal pediatrics, Northeast
12 Association, Neonatologists, and New Hampshire
13 Pediatric Society.

14 Do you hold any type of administrative
15 or office in any of those societies?

16 A. No.

17 Q. Under institutional committees and
18 appointments, you have several. It's Southern
19 New Hampshire Medical Center. And I believe
20 Southern New Hampshire Medical Center is a
21 facility in Nashua, correct?

22 A. Correct.

23 Q. Are you still currently director of

1 neonatology?

2 A. Yes.

3 Q. What do your responsibilities as
4 director of neonatology entail?

5 A. I make sure things work right. I put
6 the team together. I schedule the team, I look at
7 the quality of care, I make sure that my team of
8 providers is working well with nurses and nursing
9 issues.

10 I'm the interface between the doctors
11 and the nurses. I serve on various hospital
12 committees on behalf of the neonatal program.

13 Q. How many doctors are in your group?

14 A. Four doctors, one nurse practitioner.

15 Q. And they're all neonatologists?

16 A. Actually -- yes, it's four and one.

17 No.

18 Q. They are not. What are the other
19 others --

20 A. The others are pediatric hospitalists.

21 Q. So you're the only neonatologist?

22 A. No, there are two neonatologists, and
23 two and a half pediatric hospitalists, but we all

1 do the same job.

2 Q. I assume that the neonatologists cover
3 the NICU at Nashua, correct?

4 A. We all do. Everyone in the group does.

5 Q. Including the pediatric intensivists?

6 A. Yes. We have pediatric hospitalists
7 who are doing some neonatology, and we have
8 neonatologists like myself doing some hospital
9 pediatrics.

10 Q. Because of that, I assume you're a
11 level 2 nursery here in Nashua?

12 A. Yes.

13 Q. How does the state of New Hampshire
14 license NICUs?

15 A. This is the Live Free or Die state, and
16 so they don't. They have no regulations, and you
17 can call yourself whatever you want.

18 Q. Okay.

19 A. And we've talked about it internally,
20 the nurse manager and I, about if we should call
21 ourselves a level 3, when should we, is there any
22 reason to. It's totally arbitrary, and it's
23 self-designated in this state.

1 Q. There's no licensing by any entity,
2 Department of Health or any other entities similar
3 to that in the state?

4 A. Well, they licensed our beds. We're
5 licensed for a 17 bed neonatal unit, but they don't
6 classify the unit, we do.

7 Q. Is there any outside entities that have
8 input into the classification of your NICU, like
9 insurance companies or Joint Commission of
10 Accreditation of Healthcare Organizations?

11 A. About a year ago the American Academy
12 of Pediatrics came out with a new classification
13 system. And in that system there's level 1, 2, 3A,
14 3B, 3C and 3D, and we're a 3A according to the
15 American Academy of Pediatrics.

16 The hospital in Manchester can do more
17 than we can, they're a 3B. Dartmouth can do more
18 than that, they're a 3C, and Boston Children's
19 Hospital is at the top, they're at 3D. So the
20 American Academy does have a system that would put
21 us as a 3A.

22 Q. What's the difference between, as far
23 as you understand it in this AAP accreditation

1 process or -- probably a wrong -- characterization
2 process is probably a better term, correct,
3 because it's not an accreditation process?

4 A. You're right.

5 Q. But as characterization of levels, what
6 would be the difference between your facility and
7 the facility up in Lebanon?

8 A. They can take care of premature babies
9 as small as you can get. We put a limit. We don't
10 go below 27 weeks, or 1,000 grams birthweight.

11 Q. Okay.

12 A. They go down to 500 grams and 23 weeks.
13 So they take care of more smaller premies, and,
14 secondly, they have more subspecialty. They can do
15 pediatric surgery; we can't. They've got pediatric
16 cardiologists; we don't. They have a lot of
17 pediatric subspecialists that we don't have.

18 Q. How many beds is your NICU here?

19 A. Today it's 17. We just got a CON
20 approved, because we're going to rebuild it. And
21 although it's going to be bigger floor-space wise,
22 we're cutting to 14 in the future.

23 Q. So you'll be a 14 bed or Isolette unit?

1 A. Yes.

2 Q. I assume from what you said, then the
3 state of New Hampshire does have certificate of
4 needs process?

5 A. Yes.

6 Q. Which you have to apply for and get
7 approval. You can't just willy-nilly build beds
8 in New Hampshire?

9 A. We were approved two or three months
10 ago for a new construction.

11 Q. Okay. Assume you know Doctor Robert
12 Darnall, correct?

13 A. I know him well.

14 Q. Does he have any subspecialty in
15 neonatology that you're aware of?

16 A. Yes.

17 Q. Do you know what his subspecialty is?

18 A. I know his primary research interest,
19 and probably clinical interest as well, relates to
20 regulation of breathing and apnea.

21 Q. So if you were going to characterize
22 his practice, if there is such a thing, it's a
23 pulmonary neonatologist, would that be fair?

1 A. I don't know how much he does with
2 pulmonary physiology, it's more breathing control
3 and apnea. He's working a lot in the lab now.
4 He's, I think, cut back to half time clinical and
5 half time research, and I think it relates to
6 breathing control, not pulmonary mechanics, and I
7 may be wrong on that.

8 Q. Okay. I assume you recognize him as an
9 expert?

10 A. Yes, I have much respect for
11 Doctor Darnall.

12 Q. Have you ever referred patients to him
13 or to his institution?

14 A. Yes.

15 Q. Do you refer patients to him that --
16 babies that need surfactant and long-term
17 ventilation support?

18 A. If they're under a thousand grams or
19 less than 27 weeks, I would call Dartmouth. As I
20 say, they've got, I believe, five attendings there.
21 So you might say, well, there's one chance in five
22 he'd answer the phone, but I think he's only
23 working half time clinical, so there's about one

1 chance in ten that he would be the accepting
2 doctor.

3 Q. Okay, so if you have a particularly
4 difficult case, you would tend to call up to
5 Lebanon, is that fair?

6 A. That happens on occasion. Five to ten
7 times a year.

8 Q. Sure. You would defer to the doctors,
9 then, in Lebanon and Doctor Darnall on issues
10 that -- difficult issues that relate to pulmonary
11 conditions of the neonate?

12 A. I'm not sure I understand what you're
13 getting at.

14 Q. Let me try to rephrase it. If you,
15 again, had a particularly difficult case and you
16 called up to Lebanon and talked to Doctor Darnall
17 or one of his partners up there, and they
18 suggested that you do a certain course of
19 treatment or that you send the baby up to Lebanon,
20 I assume that you would defer to their judgment on
21 those issues?

22 A. Any case I can keep here, I don't need
23 their advice. Once I turn the case over to them,

1 it's their case; they can do what they want.

2 Q. Okay.

3 A. I send a lot of these cases to
4 Dartmouth not because I don't have the skills and
5 expertise, but some of it's because I'm using
6 pediatric hospitalists and nurses that haven't
7 taken care of those small babies, and I'm doing it
8 to -- for that reason. But I don't think that
9 their neonatologist have more or less expertise
10 than I do.

11 Q. All right. Yet, you will send patients
12 up to Lebanon that are particularly difficult,
13 correct?

14 A. These are cases that are going to be
15 difficult for my team and my nurses, not
16 necessarily me.

17 Q. All right.

18 A. And it's not in their best interest to
19 be here for that reason.

20 Q. But you also told me that you would
21 contact the doctors up at Hanover or talk to
22 Doctor Darnall for some advice on some occasions,
23 correct?

1 A. Not necessarily to get advice, but to
2 accept the patient in transfer.

3 Q. Doctor, back to your institutional
4 committees and appointments. As a director of
5 neonatology at Southern New Hampshire Medical
6 Center, what percentage of your professional time
7 is spent in that capacity?

8 A. In the administrative role, five hours
9 a week. Sometimes more or sometimes less. If
10 we're recruiting, I'm going through an active
11 recruiting process, I might put in ten or 15 hours
12 doing those weeks.

13 Q. How often do you recruit doctors?

14 A. Whenever someone leaves the group. I
15 have someone leaving at the end of this month, and
16 I have a new person starting at the end of July. I
17 end up recruiting one new person every couple of
18 years. I have been here nine years and probably
19 have done four or five recruitments.

20 Q. How long -- how many weeks span does it
21 take you to recruit a doctor, typically?

22 A. The biggest holdup in the process is
23 getting the state license. We tell these people

1 after we reach an agreement for a contract they
2 aren't going to start working for about four more
3 months.

4 So our recruitment may take anywhere
5 from one to six months, and then it's three or four
6 waiting for license after that. So it's between
7 six to 12 months.

8 MR. BULLOCH: Off the record.

9 (Discussion off the record.)

10 Q. Doctor, I also show that you are the
11 chairman of the neonatal intensive care unit
12 committee?

13 A. Yes.

14 Q. What does that entail?

15 A. It's a meeting every month or two with
16 our providers and the nurse managers just talking
17 about issues that are going on, clinical issues.

18 Q. Okay, and that's -- by providers you
19 mean the doctors and the nurses?

20 A. Yes.

21 Q. And as the chairman, do you spend some
22 time getting this organized and spending some time
23 doing those type of things?

1 A. A little time, not much.

2 Q. How much -- how many hours a week would
3 you estimate you spend in that role?

4 A. One or two hours a month. If a meeting
5 lasts an hour, I probably put in another hour
6 before the meeting and after the meeting.

7 Q. Okay. You're also on the Maternal
8 Child Health Council?

9 A. Yes.

10 Q. What is the Maternal Child Health
11 Council, Doctor?

12 A. That's a monthly meeting, the last
13 Tuesday of the month. It has pediatricians,
14 obstetricians, one anesthesiologist, one family
15 practitioner, nurses from pediatrics, labor and
16 delivery, the neonatal unit and a couple of
17 administrators.

18 It's a group of about 15 people that
19 get together. It's an opportunity for -- for
20 example, the pediatricians to bring up an issue
21 with the obstetricians, and then they take it back
22 to their department, or the obstetricians at the
23 last meeting were upset with the anesthesiologists

1 and came to that forum.

2 Q. Who covers resuscitation in your
3 hospital when resuscitation's necessary?

4 A. My group, one of the five of us. We're
5 there 24 hours a day, and we go into about half the
6 deliveries now.

7 Q. And does the anesthesiologist have any
8 role in neonatal resuscitation?

9 A. No.

10 Q. Now, the Maternal Child Health Council,
11 what would you estimate you spend per week doing
12 that?

13 A. Just going to the meetings, and that's
14 one and a half hours once a month, and maybe I have
15 one follow-up issue to work on after the meeting,
16 maybe a memo to send out, couple of phone calls to
17 make, not much.

18 Q. And does that happen on a monthly basis
19 since you're the head of the neonatology group?

20 A. Yes, I probably put in two hours a
21 month on activities related to that --

22 Q. In addition to the hour and a half?

23 A. No, that counts the hour and a half.

1 Q. The next thing I'm showing is principal
2 investigator for the Boston University birth
3 defect study at Southern New Hampshire Medical
4 Center.

5 A. That's still going on, but it doesn't
6 take much time.

7 Q. What are you investigating? Obviously
8 birth defects, but can you give me some specifics
9 on what that's all about?

10 A. They look at cases that are born with
11 birth defects at our hospital, and five other
12 normal babies born that same week, and they
13 interview the family of the birth defect baby and
14 the five control families, and they're looking for
15 causes of birth defects. We're one of about 30 or
16 40 hospitals in this study.

17 Q. I see. So you're looking to see if the
18 mother was exposed to some noxious chemical, for
19 example?

20 A. Yes.

21 Q. Now, how often --

22 A. Now, I know one of their emphases is
23 looking at asthma medicines, and -- because so many

1 pregnant women have asthma and take a lot of
2 medicine, they're trying to see if those medicines
3 cause birth defects. That's one specific focus
4 that's going on now.

5 Q. Does the study of children with birth
6 defects include children that are diagnosed with
7 cerebral palsy?

8 A. No.

9 Q. Even though that cerebral palsy could
10 be caused by some prenatal event?

11 A. We're looking at babies diagnosed as
12 having a birth defect in the newborn period, and
13 cerebral palsy isn't diagnosed in the newborn
14 period.

15 Q. Okay, fair enough. As a principal
16 investigator with that group, how much time do you
17 spend in that?

18 A. Maybe that's misleading. I'm only
19 principal investigator at our hospital.

20 Q. I understand.

21 A. I'm the liaison for our hospital. In a
22 year's time, the entire year, two or three hours.
23 I have to go to annual meeting of the institutional

1 review board and present the activities of the
2 project. That's about it.

3 Q. Okay. Continuing medical education
4 committee, I think I know what that is, I won't
5 have you explain that to me, but how much time do
6 you spend on that a month or a week?

7 A. Two hours a year. We have one meeting
8 a year.

9 Q. Okay.

10 A. And we plan the program for the next
11 year.

12 Q. Ethics committee, how much time do you
13 spend on the ethics committee?

14 A. I go to one meeting a month.

15 Q. So, hour or two a month?

16 A. Yes.

17 Q. Credentials committee?

18 A. That's a good committee. That's two
19 hours a month. It's a very good committee.

20 Q. Are there any other institutional
21 committees that you're involved in that I'm not
22 showing on my somewhat outdated CV?

23 A. No. This one says I'm on the

1 breastfeeding practice committee.

2 Q. Okay.

3 A. They meet once a month, but I commonly
4 do not attend. If I'm working that day, I attend.
5 If I'm not working, I send someone else for me.

6 Q. Anything else?

7 A. No.

8 Q. Okay, noninstitutional committees. You
9 have the neonatal resuscitation program provider.
10 Can you explain to me what that is?

11 A. That just means that I hold a card that
12 says I'm approved, I've taken the program. The
13 neonatal resuscitation program, I've taken it and
14 passed.

15 Q. You have no administrative
16 responsibility in that organization that certifies
17 doctors as being neonatal resuscitators?

18 A. Not at this time. I used to, but I
19 stopped in 1996.

20 Q. Okay. And then I've got PALS
21 providers, is that the same type of thing?

22 A. Yes.

23 Q. And you hold no administrative capacity

1 with Pediatric Advanced Life Support program?

2 A. Correct. On the bottom.

3 Q. Oh, yeah, you have a few more. There's
4 the New Hampshire state birth defects surveillance
5 system advisory committee that I list as well.
6 Are you still involved in that?

7 A. No, I don't think that group is active
8 right now.

9 Q. Okay, and then you were kind enough to
10 hand me portions of your current CV, and what is
11 NICU NET moderator?

12 A. It's an Internet forum for NICU
13 conversations and discussions from all over the
14 world, and there are five people who serve as
15 moderators, so that we take two week blocks. So
16 every two weeks I approve, disapprove, or edit
17 postings before they get posted.

18 Q. Okay. Now, you do that -- there's five
19 of you, so you do that ten times a year, correct?
20 Would that -- my math is terrible. You do it ten
21 weeks out of the year?

22 A. Correct.

23 Q. How much time does that take you when

1 you're actively serving as the moderator?

2 A. Ten or 15 minutes a day. Once a day I
3 sit down and look at the postings and deal with
4 them.

5 Q. Okay, the Nashua Police Department
6 consultant. What capacity is that, sir?

7 A. We're looking at drug use in pregnancy
8 and its effect on the babies. And there have been
9 two women so far prosecuted in the last year for
10 cocaine use and its effect on the baby.

11 Q. So you're actively involved in the
12 criminal justice system with this, or are you
13 offering testimony?

14 A. No, I'm working with the -- both the
15 police and the prosecutor's office in developing
16 their strategy and their approach to this.

17 For example, one of the first meetings
18 I had a lot of input in deciding what drugs we
19 would focus on. What drugs we could say are
20 harmful to a baby, and what ones we couldn't say
21 that.

22 Q. Okay, the two cases that were
23 prosecuted was where the mother was abusing some

1 harmful substance?

2 A. They were both cocaine. I feel
3 strongly about cocaine.

4 Q. When the prosecutor brought charges
5 against those two mothers, were you called to
6 testify at trial about the harmful effects of
7 cocaine on the developing fetus?

8 A. On one of them they reached a
9 settlement, a plea bargain, just before it went to
10 trial, but I would have testified as an expert for
11 the state.

12 Q. Okay, and the other one?

13 A. One of my partners happened to be the
14 treating physician and testified.

15 Q. Okay. And this has been going on
16 since -- roughly two years or year and a half?

17 A. Yes.

18 Q. Last year did you testify at all in any
19 criminal matter?

20 A. No.

21 Q. How much time are you spending in that
22 role per month?

23 A. Most months, none. It's as needed.

1 Q. The two cases that you dealt with this
2 past year, how much time total did you spend on it
3 in that capacity?

4 A. Five hours.

5 Q. And you're also the physician volunteer
6 for New Hampshire medical malpractice screening
7 panels, correct?

8 A. Yes. That's a new program in the
9 state, and I haven't had a panel to sit on yet, but
10 I'm -- if you called the board, they'd tell you I'm
11 available. If one comes up in my field, they may
12 call on me.

13 Q. Were you involved in setting this
14 program up through the legislature or anything?

15 A. Not at all.

16 Q. So New Hampshire now has an
17 arbitration -- well, let's back up. It probably
18 has -- before you can file a claim in
19 New Hampshire it has to be approved as malpractice
20 by this panel, is that it?

21 A. No, it's after it's filed the case goes
22 to a panel of three people. I know one's a lawyer,
23 one's a doctor, and I'm not sure about the third

1 one.

2 And if the panel reaches a unanimous
3 decision, then that can be presented to the jury.
4 If it's not unanimous among the panel, nothing
5 happens.

6 Q. So you have no ability to find for the
7 plaintiff or the defendant, but if it's unanimous,
8 it goes to -- the defense or the plaintiff can
9 bring that up to the jury's attention, correct?

10 A. That it went to this panel, and that's
11 what the panel found, yes.

12 Q. And how much time do you spend -- oh,
13 you told me, you haven't spent any time yet,
14 correct?

15 A. Right.

16 Q. I've got manuscript reviewer for
17 numerous publications. Are you still involved in
18 any of those?

19 A. I probably did more last year than any
20 other year in my career.

21 Q. Are you reviewing for -- I've got two,
22 four, six, eight -- nine journals, or are there
23 additional ones?

1 A. I have 11 now.

2 Q. Okay, what are the two new ones?

3 A. I don't know.

4 Q. Oh, alphabetical. And I assume you
5 review articles as requested?

6 A. Yes.

7 Q. You don't review every article that
8 goes into these publications?

9 A. Oh, no.

10 Q. How much time a month are you spending
11 in this role?

12 A. It would average about one hour a
13 month.

14 Q. For all of these journals or each
15 journal?

16 A. Total. That is I probably get one
17 article to look at every other month, and I
18 probably put in about two hours on it.

19 Q. And you've got some very interesting
20 additional clinical experience. Can you give me
21 some idea how you became a neonatologist and a
22 veterinarian at the same time?

23 A. I worked with gorillas at the

1 Cincinnati Zoo. I was in Cincinnati, and we
2 consulted with them when they had babies. That was
3 my first experience. We brought them in, they were
4 small, and the mother wasn't taking care of them,
5 and we put them in incubators, and fed them, and I
6 got to do that.

7 In Kentucky we worked closely with the
8 thoroughbred industry. Their vets would round in
9 our neonatal unit, and we would round on their
10 farms.

11 I took care of Bongo antelopes at the
12 Pittsburgh Zoo. There was an infant born that got
13 very sick, went into liver failure. They had
14 called on me because I had helped them earlier with
15 the delivery and resuscitation of giraffes.

16 I read where a mother had a giraffe at
17 the zoo that died during birth -- the calf died.
18 And I called them up and said next time you have
19 one of these births, give me a call, I'll come and
20 see if I can help, and they did.

21 So I resuscitated a giraffe, and then
22 after that they called me for the bongo. The
23 alpacas we have here on our farm.

1 Q. Doctor, you have also provided us a
2 list of various articles that you have published
3 over the years. What I'd like to ask you is which
4 ones do you feel are directly related to the
5 issues in this case?

6 A. I don't know if any of them are. None
7 of them come to mind.

8 Q. Are there any articles specifically
9 related to hyaline membrane disease?

10 A. No, I don't think so.

11 Q. What about articles related to using
12 ventilators in newborns?

13 A. No.

14 Q. And I did not see any articles related
15 to use of surfactants, is that fair?

16 A. That's fair.

17 Q. Okay. Article No. 15, and yours might
18 be different, Doctor. I'll just give you the
19 title of it, it was Hermansen and Hasan, An
20 Evaluation of a Computer Program to Predict the
21 Outcome of Hyaline Membrane Disease, published in
22 the American Journal of Perinatology. I assume
23 you recall that article?

1 A. That was an interesting one, yes.

2 People at Vanderbilt wrote a computer program, and
3 it looked at blood gasses for the first six or 12
4 hours, I don't remember, and it would predict
5 whether the outcome was -- the outcome was mild,
6 moderate or severe, and they published that.

7 Well, I took babies and took the blood
8 gasses from the same time period, six hours, and I
9 showed them to my two colleagues, two
10 neonatologists, and asked them to predict the
11 outcome, and they beat the computer.

12 So even though the Vanderbilt people
13 published that their computer was good, I showed
14 that the neonatologist was still better.

15 Q. So the neonatologist was better at
16 predicting whether or not a child would develop
17 hyaline membrane disease than this computer
18 program?

19 A. No, the outcome of hyaline membrane
20 disease.

21 Q. Okay.

22 A. I think it was things like time on the
23 ventilator, time on oxygen.

1 Q. Severity of the hyaline membrane
2 disease?

3 A. Yes. They had to classify it as mild,
4 moderate or severe, and the computer class made a
5 prediction, and the neonatologist made a
6 prediction, and the neonatologists were better.

7 Q. Part of what I noticed in there was
8 your reasoning that you felt the neonatologists
9 did better was that the computer didn't really
10 consider birthweights. It had to be inputted, but
11 it didn't consider birthweights in making a
12 projection on what happened.

13 A. I don't remember that.

14 Q. I'll give you --

15 A. I haven't looked at that article for a
16 long time.

17 Q. If I can find it here. Doctor, I
18 apologize, but the study I'm going to give you has
19 my highlighting on it. But if you want to take a
20 look at the abstract or any portion of the article
21 you want to review, feel free to take time to do
22 that. Very interesting article.

23 A. I haven't looked at it in a long time,

1 but I was right. It says, although the computer
2 program predicted the outcomes with moderate
3 success, it was less accurate than the
4 neonatologists.

5 Q. Read on. It's an interesting study.

6 A. Where do you want me to start reading?
7 That was the last sentence of the abstract.

8 Q. The question, and I think I highlighted
9 it there, was that the reason that you found that
10 the neonatologists did better was that the
11 computer did not consider birthweights.

12 A. I don't remember that. If that's what
13 it says, that's what it says.

14 Q. Let me point it out to you.

15 A. Fine.

16 Q. Let me read what you've got here under
17 discussion. Surprisingly, we found that the
18 neonatologists predicted the outcome of
19 uncomplicated HMD, hyaline membrane disease, more
20 accurately than the model.

21 Additionally, physicians with the
22 greatest experience may be the best predictors.
23 The physician with the greatest clinical

1 experience, 12 years, was the best predictor at 24
2 correct, followed by the physician with 11 years
3 of experience, 22 correct, and then the physician
4 with two years of experience, 18 correct.

5 A. That was probably me.

6 Q. So certainly experience in this field
7 pays off, correct?

8 A. Right.

9 Q. Okay, go on here. Here. It is
10 noteworthy that the three groups of patients
11 selected by the physicians were of significantly
12 different birthweight and gestational age. It is
13 likely that the physicians used birthweight to
14 help differentiate among the three outcomes.

15 Okay?

16 And then in the abstract you've got,
17 while actual severity was linked to birthweight,
18 the model did not utilize birthweight in its
19 predicted algorithm.

20 And I'll be happy to share that with
21 you again if you want to review it. But the
22 question is, do you believe that -- or at least
23 the time that you wrote that -- that the reason

1 that the physicians did better was, one, based on
2 experience, and, two, based on the fact that they
3 took birthweight into consideration?

4 A. It appears so.

5 Q. So I guess the question that I have for
6 you, then, Doctor, is why? What is the relation
7 between birthweight and severity of hyaline
8 membrane disease?

9 A. Statistically as a group, the smaller
10 you are, the worse the outcome.

11 Q. Okay.

12 A. But that isn't true for every baby, or
13 that would be the only factor to consider.

14 Q. Certainly.

15 A. There are some very tiny babies that
16 have good outcomes and some very big babies that
17 have bad outcomes, but it's one factor that is
18 worth considering.

19 If you're going to predict outcomes,
20 I'd predict a thousand gram baby would have a worse
21 outcome than 2,000 gram babies, and I'll be right a
22 reasonable amount of the time, but not always.

23 Q. Sure. So just to summarize then,

1 birthweight is important in predicting outcome?

2 A. Yes.

3 Q. Correct?

4 A. Yes.

5 Q. The bigger the baby, the more likely a
6 better outcome, correct?

7 A. Sure.

8 Q. Article 18 is an article captioned
9 Diminished Splenic Function in Asphyxiated Term
10 Infants. Do you recall that study?

11 A. Yes.

12 Q. I believe the study demonstrated that
13 hypoxic events cause changes in the function of
14 the spleen, correct?

15 A. Yes.

16 Q. And was the change that you found
17 related to decreased blood cell counts?

18 A. No. There are specific types of cells,
19 the red blood cells, and when they get old or
20 damaged, they get pits in them, and they were
21 called pitted or pocked cells.

22 Q. Okay.

23 A. And the spleen should take those out of

1 circulation. And if you look with a special stain,
2 the pathologists can look and count pitted cells,
3 and if you have too many of them, if the number is
4 high, that means the spleen is not doing its job.
5 And we looked at babies with low Apgar scores and
6 showed they had higher pit counts.

7 Q. Do you currently use pit counts in your
8 practice?

9 A. No. Virtually no one does.

10 Q. Is this related, in your mind, to the
11 hypoxic events on the bone marrow as well, where
12 you see thrombocytopenia?

13 A. No. This is the spleen not removing
14 cells that it should remove.

15 Q. And the reason that it's not removing
16 cells the way it should, because it took a hit, if
17 you would, hypoxic hit, correct?

18 A. Yes.

19 Q. So in that regard it is related to what
20 you expect to see other effects on the body with
21 hypoxic hits, correct?

22 A. With asphyxia, yeah, I -- this is an
23 asphyxial event.

1 Q. So much like the spleen, you expect to
2 see some effect on kidney function, for example?

3 A. With asphyxia, you commonly see kidney
4 dysfunction.

5 Q. Liver dysfunction?

6 A. Very common.

7 Q. Bone marrow dysfunction?

8 A. Very common.

9 Q. You tend to see damage or injury of
10 some sort to the intestinal tract?

11 A. Not as common, but you can.

12 Q. Compression of bone marrow resulting in
13 thrombocytopenia?

14 A. Well, they get thrombocytopenia, but
15 I'm not sure if that's always the cause of it. It
16 may be that the platelets are being consumed or
17 destroyed in the circulation.

18 Q. Okay.

19 A. You get low platelets either from lack
20 of production or increased destruction, and it may
21 be as much of a destruction problem.

22 Q. So, for example, if you have a bleed in
23 the head as a result of a hypoxic ischemic event,

1 you might be losing platelets in that regard?

2 A. I'm thinking if you go into a process
3 what we call DIC, where they begin using up their
4 clotting studies, they drop their platelets.

5 Q. Now, you mention this occurring in an
6 asphyxial state, correct?

7 A. Yes.

8 Q. Is asphyxial state the same in your
9 mind as a hypoxic ischemic event?

10 A. Sometimes.

11 Q. What about a prolonged hypoxic ischemic
12 event?

13 A. Sometimes it could give you your
14 classical asphyxia picture, sometimes it won't.

15 Q. When does it, and when doesn't it?

16 A. Asphyxia is interference with
17 respiration resulting in hypoxia and acidosis.
18 Now, you can have hypoxia and ischemia taking place
19 without it -- without the asphyxia process, without
20 it -- the primary process being interference with
21 respiration, without it being a placenta problem,
22 without it being an airway issue, that's asphyxia.
23 Asphyxia gives you a systemic response.

1 The body goes into certain compensatory mechanisms,
2 and then the body fails, the heart fails, the
3 organs all suffer. That's an asphyxia picture.

4 But a baby could be hypoxic and have
5 poor blood flow and not go into the asphyxia
6 picture. Just get, you know, a hypoxic insult, but
7 not build up the acid, not go into multi-organ
8 failure, not begin having seizures, not -- it's
9 more of a hypoxic problem than an asphyxial
10 problem.

11 Q. All right, you mentioned the acidemia,
12 correct?

13 A. Yes.

14 Q. And actually, the acidemia, you've
15 written, is protective at times, correct?

16 A. Yes.

17 Q. So the absence -- you have found that
18 the absence of acidemia is sometimes more damaging
19 to the body organs than not having acidemia,
20 correct -- or having acidemia?

21 A. Yes. With your asphyxiated babies,
22 with birth asphyxia, and they come out and they
23 look asphyxiated and they act asphyxiated.

1 If they do not develop an acidemia, I'm
2 especially worried about them. I think that's a
3 very high-risk situation. If they get a mild or
4 even moderate acidemia, they tend to do pretty
5 well.

6 Q. Now, you're describing -- also you
7 describe shunting that occurs normally when you go
8 into a hypoxic situation, correct?

9 A. Well, with the asphyxia sequence, that
10 happens.

11 Q. Well, it happens in prolonged
12 hypoxemia, too, doesn't it, Doctor?

13 A. No, usually not.

14 Q. How do you explain then low oxygen
15 levels in babies with congenital heart disease who
16 don't sustain brain damage?

17 A. Okay, that's a good example. Those
18 kids with congenital heart disease may be hypoxic,
19 but they're not asphyxiated, they're not shunting,
20 their kidneys aren't hurt.

21 You know, they're not showing asphyxia,
22 they don't get acidotic, they don't drop their
23 blood pressure, they don't go into kidney or liver

1 failure, but there's hypoxia and there's the risk
2 of hypoxia to them, but they're not shunting.

3 Q. And yet no neurological damage in those
4 children?

5 A. If it's severe and prolonged, they can.
6 That's why we operated on them. That's why we try
7 to get their oxygens optimized.

8 Q. Those are certainly very low PO 2
9 levels in many of those children, correct?

10 A. Yes.

11 Q. And very prolonged periods of time,
12 obviously.

13 A. Variable, yes.

14 Q. Do those children tend to show any
15 damage to any other organ if they have damage to
16 the brain?

17 A. No. And that's a very good example of
18 significant hypoxia not looking like asphyxia.

19 Q. All right, so I'm confused. Are you
20 saying that those children can have damage to
21 their brain but not show any multisystem organ
22 failure?

23 A. Yes.

1 Q. Do you know of any articles that I
2 can -- that you can send to me that I can review
3 that supports your position?

4 A. I could find them. I'm willing to
5 spend an hour coming up with articles that would
6 show cyanotic heart disease resulting in brain
7 damage without multi-organ dysfunction or failure.
8 I know I could find articles that show that.

9 That's -- that's what happened. That's
10 why cyanotic heart disease is bad and why we have
11 to fix it, but they don't go into shock and
12 multi-organ failure. That's a different process.
13 Asphyxia is different than hypoxia.

14 Q. And your contention is that you don't
15 get the multisystem organ failure unless you have
16 asphyxia?

17 A. That's an asphyxial phenomena.

18 Q. And, again, explain to me the
19 difference between asphyxia and a hypoxic
20 condition. In your mind, what are the
21 similarities and what are the differences?

22 A. I'm going to talk about acute asphyxia,
23 it's a sudden event, hypoxia is part of it. They

1 get hypoxic, they build up acid, the heart tries to
2 compensate, it ultimately gives out, the organs
3 suffer, the brain gets hurt primarily from
4 ischemia.

5 With asphyxia the brain damage is
6 usually from the heart giving out and having
7 ischemia to the brain. It's not even from the
8 hypoxia.

9 Which is a whole different process than
10 just having severe hypoxia for a prolonged period
11 of time. Their hearts don't give out, they may not
12 become acidotic, they don't go into kidney failure,
13 they don't seize, but their brain cells can suffer
14 a lack of oxygen.

15 Q. And then what do you typically see by
16 way of blood pressure in the latter group?

17 A. Normal.

18 Q. Throughout?

19 A. Yes.

20 Q. And what do you typically see as far as
21 pulse oximetry readings or PO 2 levels?

22 A. Low.

23 Q. How low?

1 A. Well, that's -- that's actually a good
2 question, how low of hypoxia does it take to cause
3 brain damage. I think it's a function both of how
4 low and how long.

5 You could drop very low and if it's
6 only for a minute or two or three or probably five,
7 it's not going to hurt you. But if you drop
8 moderately low for many hours, that can hurt you.
9 So it's not just a function of how low causes
10 damage, but how long and how low.

11 Q. And is there any difference in the
12 incidence of multisystem organ failure in those
13 two groups, one that's dropping very low for a
14 shorter period of time, and one that's dropping
15 low, not quite as low, for a more prolonged period
16 of time? Neither would exhibit multisystem organ
17 failure, in your opinion?

18 A. Because blood flow is being maintained,
19 that's right.

20 Q. To those organs?

21 A. To the organs of the body.

22 Q. But not the brain?

23 A. Blood flow is maintained, it's not just

1 good blood. It's hypoxic blood. It's blood
2 without enough oxygen in it, but there's blood flow
3 going on.

4 Q. I'm sorry, it's hypoxic blood going to
5 the other organs, too; why don't you see damage to
6 those organs?

7 A. Their damage comes primarily from
8 ischemia.

9 Q. Whose damage?

10 MR. BECKER: The other organs.

11 MR. BULLOCH: He's testifying, Mike.
12 He's doing fine.

13 A. We're talking about these other organs,
14 the kidneys. The kidneys can withstand the
15 hypoxia. Ischemia is what really causes them to go
16 into failure. The liver, it's generally not hurt
17 from hypoxic, it's the ischemic component.

18 Q. The heart?

19 A. The same.

20 Q. The bone marrow?

21 A. Chronic hypoxia -- chronic turns on the
22 bone marrow.

23 Q. How long does it take to turn on the

1 bone marrow?

2 A. I don't know the answer to that.

3 Q. Well, do you have any opinion? Is it
4 days, is it weeks, is it hours?

5 A. I'd be speculating.

6 Q. Okay.

7 A. It would be total speculation, and I'm
8 not going to do that today.

9 Q. And, Doctor, I don't want you to do
10 that today. What I understand you're saying,
11 though, unless you have an acute, sudden, dramatic
12 asphyxial event, you're likely not to have
13 multisystem organ failure, correct?

14 A. Correct.

15 Q. Then how do you explain the appearance
16 of multisystem organ failure when you have
17 uteroplacental insufficiency? Explain that to me,
18 Doctor, because isn't that a prolonged, low level
19 hypoxemia?

20 A. There is a prolonged, low level
21 hypoxia. They don't go into multi-organ failure.

22 Q. You're saying that women that exhibit
23 uteroplacental insufficiency do not show any hits

1 to the bone marrow, the kidney, the heart, the
2 intestines, is that what your testimony is today?

3 A. No. We're talking about longstanding.
4 Their bone marrow does get stimulated. The babies
5 doesn't grow well. If their head keeps growing,
6 their brain's fine, their kidney's are fine, their
7 liver's fine, the heart's fine.

8 It's true that the bone marrow gets
9 turned on; it doesn't fail. The bone marrow
10 doesn't go into bone marrow failure or poor
11 function, it gets turned on, it works overtime.

12 I don't think chronic placental
13 insufficiency causes organ failure of any organ
14 that I can think of.

15 Q. You've never seen that or you've never
16 seen anything written on multisystem organ
17 failure -- you know what I'm meaning by
18 multisystem organ failure, is what we've been
19 talking about, correct?

20 A. Yes.

21 Q. It's a hit to the kidneys, the liver,
22 the heart, the intestine?

23 A. Right, we're talking about the same

1 process.

2 Q. I want to make sure we're talking about
3 the same thing. You're not aware of any articles,
4 or in your experience you've never seen any
5 multisystem organ failure in a uteroplacental
6 insufficiency, right?

7 A. If we're talking about chronic
8 insufficiency, that's right. Now, if --

9 Q. And chronic you mean -- and I'm sorry
10 to interrupt you -- but chronic you mean because
11 the placenta is small for the baby as opposed to
12 an abruption, correct?

13 A. Or the blood flow through the placenta
14 isn't good, perhaps mom has hypertension or severe
15 diabetes, or it's a bad placenta for some reason,
16 or mom has chronic hypoxia. That can lead to
17 chronic placental insufficiency to the fetus, and
18 they do not go into multi-organ failure.

19 Q. Okay. And you also don't see it when
20 the placenta is small for the baby's size,
21 correct?

22 A. Correct.

23 Q. All right. Doctor, article 22 is

1 listed in your CV, it's a 1995 article, it's
2 entitled incidence, timing and follow-up of
3 periventricular neuromalacia. I believe you
4 authored that article with your wife, correct?

5 A. She was the first author.

6 Q. You're well aware of that article,
7 correct?

8 A. Yes.

9 Q. One of the findings was that cystic PVL
10 occurs between 17 and 104 days of age, correct?

11 A. Yes.

12 Q. And you also found a strong correlation
13 between PVL and spastic diplegia, correct?

14 A. Yes.

15 Q. What causes PVL?

16 A. I think there are two main causes, and
17 they're totally different. One cause is poor blood
18 flow to the brain of these small premies -- and
19 it's nearly always in small premies, not big
20 premies -- but poor blood flow to the brain
21 commonly from low blood pressure or very low carbon
22 dioxide levels.

23 In the last ten years we've recognized

1 a second cause, and I don't even think we knew
2 about it in 1995, and I don't think we mentioned it
3 in there, and that's maternal infections,
4 intrauterine infections in these small premies.

5 It can release some chemicals that can
6 hurt the brain. So there may be maintenance of
7 blood flow to that area, but these chemicals go to
8 that part of the brain and destroy it.

9 Q. And the chemicals and the syndrome that
10 you're referring to is fetal inflammatory
11 syndrome, and it releases cytokines and things
12 like tissue necrosis factor, correct?

13 A. Yes. And I don't think that was talked
14 about in 1995. I know we didn't mentioned it in
15 that article. That's relatively new knowledge.

16 Q. In your study -- I'm sorry, those were
17 the only two things that caused PVL as far as you
18 are aware of, correct?

19 A. Yes.

20 Q. In your study, none of the children --
21 none of your findings show any association between
22 PVL and athetosis or athetoid movements, correct?
23 I've got the article if you want to take a look at

1 it.

2 A. I think that's right. Usually it's in
3 the area of the brain that would cause spasticity
4 to the legs.

5 Q. Right. So you're going to say
6 spasticity but not athetosis, correct, related to
7 PVL?

8 A. Yes.

9 Q. What causes athetosis in a baby?

10 A. Damage to some deeper matters of the
11 brain, I believe in the gray matter around the
12 basal ganglia, a different area of the brain
13 getting hurt.

14 I'm not going to be able to go too far
15 into this conversation, because you're beginning to
16 take me into neurology, but I'll try my best.

17 THE WITNESS: I'm going to take 30
18 seconds.

19 MR. BULLOCH: Oh, sure.

20 (Recess taken.)

21 Q. Doctor, in your newer CV that you don't
22 have a copy of that's in front of you, could you
23 take a look at page 11?

1 A. They're totally different. What's the
2 article?

3 Q. It's actually numbered, number 49.

4 A. Okay.

5 Q. What is that article titled?

6 A. Cerebral palsy.

7 Q. Do you recall what that article was
8 about?

9 A. That's the preface to the book. It's
10 about two or three pages long.

11 Q. This is the current book that is coming
12 out?

13 A. Any day now. I was hoping I would have
14 it for today.

15 Q. And article 50, what is article 50?

16 A. Perinatal infections and cerebral
17 palsy. It's a chapter in that book.

18 Q. And does that -- you authored that
19 chapter?

20 A. Yes.

21 Q. With your wife?

22 A. Yes.

23 Q. Okay. And does that deal with

1 primarily fetal inflammatory syndrome that we've
2 discussed?

3 A. Not primarily, but extensively.

4 Q. Okay.

5 A. It's -- that concept and the cytokine
6 concept are discussed, and I do say that that's a
7 cause of cerebral palsy in small preterm infants.

8 Q. And that particular chapter deals with
9 other types of infections like viral infections,
10 for example?

11 A. Yes.

12 Q. What viral infections can cause
13 cerebral palsy?

14 A. The most common one would probably be
15 what we call CMV. Others, herpes, but they're
16 relatively uncommon.

17 Q. Did you see any evidence that Margo
18 Wagoner had a CMV infection?

19 A. No.

20 Q. You reviewed the prenatal records?

21 A. I don't remember seeing those.

22 Q. Does the CMV infection have to be
23 active at the time of the delivery, or can it be a

1 longstanding infection or process?

2 A. No, it's a longstanding. Many women
3 don't know they have it.

4 Q. So if there's evidence in the record --
5 and I don't quite honestly know if there is or
6 there isn't -- but if there's evidence in the
7 record that Margo Wagoner had a CMV infection,
8 would that be important to your analysis in this
9 case?

10 A. No.

11 Q. Does CMV cause PVL?

12 A. No.

13 Q. How does CMV manifest in a baby?

14 A. 90 percent of the time it's
15 asymptomatic and it has no effect on the baby,
16 they're fine, nine babies out of ten that are born
17 with CMV.

18 Q. Obviously I'm not interested in the
19 nine out of ten, I'm interested in the one that
20 affects the baby. What happens in that baby?

21 A. To that brain specifically, some of
22 them are born with what we call microcephaly, which
23 is a very small head, very small. Others are born

1 with calcifications in their brain. When you see
2 this pattern of calcium deposits, that's not good.

3 They have hearing problems and vision
4 problems. Well, if you take away your hearing or
5 your vision sense, you don't develop very well
6 either. So that's another reason they don't
7 develop very well.

8 Q. Are there any other infections that
9 cause cerebral palsy that mom can transmit to the
10 baby?

11 A. There are many, many ways that maternal
12 infections can cause cerebral palsy. We mentioned
13 some of these viruses. Related to that would be
14 things like rubella, toxoplasmosis, syphilis.

15 I think what's relatively common is for
16 women to get infected -- get an intrauterine
17 infection late in pregnancy, and it causes the
18 placenta not to work well, and those babies are
19 born asphyxiated. They're asphyxiated babies from
20 bad placenta functioning, so that's harmful to the
21 baby.

22 Q. And, again, can it be a longstanding
23 infection that causes some uteroplacental

1 insufficiency, and that could cause damage to the
2 baby's brain without causing a recognized
3 asphyxial event?

4 A. I don't think so. I don't think when
5 we're talking about longstanding placental
6 insufficiency that we consider chronic infections
7 to be a cause of that. Usually it's a placental,
8 or blood flow or blood vessel issue, but not an
9 infection.

10 Q. Doctor -- I'm sorry, I didn't mean to
11 interrupt you. Are you done?

12 A. Yes.

13 Q. On your presentations, and I didn't
14 write down the page, but there's one numbered
15 51 --

16 A. Yes.

17 Q. -- on the CV that's been marked as an
18 exhibit. Can you tell me the title of that
19 presentation?

20 A. Common resuscitation errors, strategies
21 for improving your likelihood of success.

22 Q. Does that have anything to do with
23 surfactant rescue, or is that more birth

1 resuscitation?

2 A. Birth issues.

3 Q. Doctor, I wanted to explore with you a
4 little bit your testimonial history. How many
5 cases do you review on behalf of a lawyer during a
6 typical year?

7 A. Fifty.

8 Q. And that number has been pretty
9 constant for a fair number of years, correct?

10 A. Yes.

11 Q. I understand that you've been doing 50
12 cases a year since the 1980s, correct?

13 A. The late '80s.

14 Q. So if my math is correct -- and I don't
15 vouch for my math, it's not one of my strong
16 suits -- but you've reviewed over a thousand cases
17 in your career, correct?

18 A. That's about right.

19 Q. What percentage has been on behalf of
20 the plaintiff as opposed to on behalf of the
21 defendant?

22 A. 80 percent.

23 Q. 80 percent on behalf of the plaintiff?

1 A. Yes.

2 Q. 20 percent on behalf of the defendant?

3 Have you reviewed any cases recently for any
4 defendants?

5 A. Yes.

6 Q. Do you recall --

7 A. I have two from the last month that I
8 can think of.

9 Q. Do any of those deal with surfactant?

10 A. No.

11 Q. Have you reviewed any cases on behalf
12 of a defendant that dealt with the administration
13 of surfactant?

14 A. I've looked at a thousand cases over 20
15 years, and I don't remember one for either defense
16 or plaintiff having to do with surfactant. I do
17 not remember this being an issue in any other case.

18 Q. So it's very rare from a medicolegal
19 standpoint, correct?

20 A. It's rare clinically not to give
21 surfactant in a case like this. People give it,
22 that's why it's rare.

23 Q. All right, we'll get to that. I didn't

1 see in your file, Doctor, anything on billing
2 statements, but do you have any idea how much time
3 you have spent in reviewing this case on behalf of
4 the Becker, Mishkind law firm? I'm not -- you
5 know, I understand you do a lot of cases, so a
6 very rough estimate based on the volume of
7 material here?

8 A. Before the -- if we exclude this
9 deposition and preparation, I would have said
10 around five hours.

11 Q. Is that pretty typical for a case that
12 you review?

13 A. Commonly it's two to three hours.

14 Q. Okay.

15 A. And then not much happens until the
16 deposition. This was a little more because there
17 were reports coming in and x-rays coming in and
18 phone calls going on.

19 Q. How many depositions do you sit for a
20 year?

21 A. Twenty.

22 Q. And how much time do you typically
23 spend preparing for a deposition?

1 A. It's highly variable. Anywhere from
2 one hour to ten hours.

3 Q. What would you say it averages?

4 A. Three or four.

5 Q. And how many times have you appeared at
6 trial?

7 A. Two or three.

8 Q. This year?

9 A. Oh, I thought you meant in a typical
10 year. Well, actually this year I think it's been
11 three. I don't know why, but there were a lot all
12 at once in like February and March.

13 Q. Okay, but two to three per year is
14 typical?

15 A. Yes.

16 Q. You spend some time preparing for
17 trial, I presume?

18 A. Yes.

19 Q. About how many hours do you typically
20 spend preparing for trial?

21 A. Two or three on the records, and then I
22 always read my own deposition, and then anything
23 else that I think is important, such as other

1 depositions or -- so four to six hours.

2 Q. And then I don't assume many cases are
3 up here in Nashua, correct?

4 A. I don't know what you mean.

5 Q. Well, let me ask it this way. You've
6 represented plaintiffs -- you've served as an
7 expert on behalf of plaintiffs and defendants all
8 over the country, correct? There's not very many
9 states you haven't served as an expert witness in?

10 A. I have served as an expert in between
11 30 to 35 states.

12 Q. All right.

13 A. And just yesterday I got a call from
14 Hawaii.

15 Q. Lucky you.

16 A. I've never been to Hawaii.

17 Q. Never been to Hawaii; you're going to
18 enjoy it.

19 A. I turned it down, because I can't
20 travel that far for personal reasons.

21 Q. I assume that you probably have to
22 travel for an entire day when you testify at
23 trial, is that fair?

1 A. I do everything I can to turn it into a
2 day trip. I had a trial this spring in Birmingham,
3 Alabama, and I went from here to Birmingham and
4 back in the same day. That's hard to do.

5 Q. It is hard to do. I've done that
6 myself many times. What do you charge to review a
7 case, Doctor?

8 A. 350 an hour.

9 Q. And to testify at deposition?

10 A. When we do them here, it's \$2,000 for a
11 half-day deposition. If I have to travel
12 somewhere, it might be a little more.

13 Q. Do you charge expenses as well?

14 A. I'm not charging you for the coffee, if
15 that's what you mean.

16 Q. I'm sorry, bad question. I'm asking
17 you if you're traveling --

18 A. Oh, yes.

19 Q. -- you're charging expenses, but do you
20 charge your travel time as well?

21 A. But just for coach airfare and a cheap
22 hotel, nothing fancy.

23 Q. But what I'm getting at is if you're

1 traveling for a deposition, you're traveling from
2 door-to-door, I assume, from the time you leave
3 your home to the time you return home?

4 A. No. I'm aware that some experts do
5 that. I've never felt comfortable with that
6 concept.

7 MR. BULLOCH: Off the record.

8 (Discussion off the record.)

9 Q. Doctor, what percentage of your income
10 do you believe is derived from your work as a
11 medicolegal expert?

12 A. 20 percent.

13 Q. And would you estimate that that's
14 consistent with the amount of time that you spend,
15 20 percent of your professional time is spent as a
16 medical expert?

17 A. No, time-wise it's probably more like
18 10 per 15 percent of my time.

19 Q. Do you remember being deposed last year
20 by a Chris Troy of this law firm in a case called
21 Gabrick versus Marymount Hospital?

22 A. I don't remember that. I know there
23 was a case called Gabrick. I don't remember who

1 the lawyers were. Who was the plaintiff lawyer,
2 that might help me?

3 Q. I have no idea, sir. I'm sorry. Has
4 the amount of time that you spend -- I don't think
5 it has because you told me you're still working on
6 50 cases a year, but has the amount of time
7 changed in the past year that you spend?

8 A. It might be down a little bit this
9 year, maybe 10 percent.

10 Q. All right. But 10 to 15 percent is
11 roughly the amount of time that you spend of your
12 professional time serving as a medicolegal
13 expert --

14 A. Correct.

15 Q. -- is your best estimate as we sit here
16 today?

17 A. Yes.

18 Q. Have you ever served as a medical
19 expert for the Becker & Mishkind law firm?

20 A. Yes.

21 Q. How many times, do you know?

22 A. I think we had one trial in Cleveland.

23 Q. Do you know the name of the case?

1 A. No. I remember a little bit about what
2 the case was about.

3 Q. That's all right.

4 A. But not the name. This was five to ten
5 years ago, and I believe it was in Cleveland, and
6 I've probably given a half dozen depositions over
7 the years.

8 Q. Do you know how many times you've been
9 asked to review cases for the Becker, Mishkind law
10 firm?

11 A. About double of that. It seems like I
12 find merit to about half their cases.

13 Q. So about 12 times you've reviewed cases
14 on behalf of Mr. Becker; attorneys in his office?

15 A. Pretty much it's only Mr. Becker. I
16 don't know the other people.

17 Q. Do you know Howard Mishkind?

18 A. I've talked to him once on the phone,
19 but I've never met him.

20 Q. Did he send you a case?

21 A. He sent me one case, I turned it down.
22 I looked at it, and they didn't use me. He didn't
23 like what I said, I guess. I didn't see any merit

1 to it.

2 Q. David Kulwicki, have you ever reviewed
3 any cases for a David Kulwicki in Cleveland?

4 A. I don't remember.

5 Q. How about a John Burnett?

6 A. I know I've talked to him at some time.
7 I've never met him.

8 Q. Larry Peskin?

9 A. Doesn't mean anything to me, I've never
10 heard that name.

11 Q. What about Pam Pantages?

12 A. I don't know her.

13 Q. Doctor, have you ever been a defendant
14 in a lawsuit?

15 A. Technically twice; realistically, only
16 once.

17 Q. Do you remember what the allegations
18 were in those lawsuits?

19 A. Well, that's why I said technically
20 twice. In one case there was no allegation. They
21 filed papers without a complaint, without
22 allegations, just to beat the statute of
23 limitations, you can do that in Pittsburgh.

1 Q. Okay.

2 A. And then after about six months, they
3 withdrew. There was never discovery, never a
4 settlement, never a complaint. There were no
5 allegations.

6 Q. Okay.

7 A. The other case did file a complaint, I
8 did give a deposition, and I was released from the
9 case just before it went to trial against two
10 obstetricians.

11 Q. Okay. I assume, then, you agree with
12 me that just because a doctor is sued does not
13 mean that the doctor was negligent, is that a fair
14 statement?

15 A. That's fair.

16 Q. And similarly, just because there's a
17 bad outcome, a child has a bad outcome, does not
18 necessarily mean the doctor was negligent either,
19 does it?

20 A. That's correct.

21 Q. And I assume then you would agree that
22 just because a child has cerebral palsy doesn't
23 mean necessarily that the doctor did anything

1 wrong either, correct?

2 A. Correct.

3 Q. In fact, most cases of cerebral palsy
4 are not caused by physician negligence, true?

5 A. True.

6 Q. You have this Clinics in Perinatology
7 coming out, the Perinatal Causes for Cerebral
8 Palsy in which you're the guest editor, correct?

9 A. Correct.

10 Q. In fact, you've had several
11 publications with Clinics in Perinatology, don't
12 you?

13 A. This is my second, and I've been in
14 contact with them about completing my trilogy in
15 2007.

16 Q. All right, I have obtained an advance
17 copy.

18 A. You have.

19 Q. Yes, I have.

20 A. How did you do this?

21 Q. I had to pay for it.

22 A. Did you really? I haven't even seen
23 it, isn't that terrible?

1 MR. BECKER: I've had some publishers
2 refuse to do that.

3 THE WITNESS: I'll be.

4 MR. BECKER: But if you work for a
5 defense firm, they make exceptions.

6 MR. MOSCARINO: Off the record.

7 (Discussion off the record.)

8 Q. You asked me how I found it?

9 A. No, is it any good? Could you find
10 anything interesting in it?

11 Q. A lot of interesting in it. It's very
12 good. It's a very interesting publication. I'm
13 sure it will sell well.

14 A. Okay.

15 Q. Anyways, you were asked to be the guest
16 editor of this publication, correct?

17 A. Yes.

18 Q. And in the preface you have analyzed
19 the contribution of each process that causes
20 cerebral palsy, correct?

21 A. Yes.

22 Q. And --

23 MR. BULLOCH: You know what, let's make

1 that an exhibit.

2 (Hermansen Exhibit No. 5 was marked for
3 identification.)

4 Q. Now, since we've marked this as an
5 exhibit, I'm not going to bother reading through
6 all of these, but where does Matthew Wagoner fall
7 in these?

8 A. The first, complications of
9 prematurity. That's the most common cause.

10 Q. Well, prematurity, though, itself can
11 cause cerebral palsy, correct? If a baby is born
12 early enough, the brain is not fully developed,
13 and the child can have cerebral palsy, correct?

14 A. It's probably more proper to say
15 complications of prematurity. Something has to go
16 bad for that brain to develop cerebral palsy.

17 Q. Well, there's a lot of children born
18 premature, low birthweight, and there's nothing
19 indicated in the medical record or in the mother's
20 prenatal record, there's genetic testing, there's
21 metabolic testing, and there's really no findings
22 of anything abnormal, correct?

23 A. There are some patients who are worked

1 up totally and you don't find a cause, that's the
2 final group on here, idiopathic, 5 to 10 percent.

3 Q. So the premature brain where all the
4 neurons are not fully formed itself is not a cause
5 of cerebral palsy is your opinion, correct?

6 A. Correct. That brain may be vulnerable
7 and susceptible to insults more than a term brain,
8 but it's the insult to that premie brain.

9 Q. In another article you talk about
10 gasping. Is that equivalent to grunting?

11 A. No, totally different.

12 Q. What is gasping?

13 A. Medically we use that term the same as
14 laypeople do, it's taking a gasp for a breath.
15 Usually it happens late in asphyxia. I -- I've
16 never seen it written, but I've always believed if
17 I see somebody gasping, a baby, their pH is below
18 7. I think it happens with severe asphyxia and
19 severe acidosis. They take a gasp to breathe,
20 attempt to breathe.

21 Q. And you didn't see any evidence of
22 gasping in Matthew Wagoner's medical record,
23 correct?

1 A. Correct.

2 Q. Do you have to take that, Doctor?

3 A. No, I was going to tell you gasping is
4 to catch one's breath with an open mouth as in
5 exhaustion or astonishment. So to catch one's
6 breath with an open mouth.

7 Q. This is a late symptom, correct?

8 A. With asphyxia, yes.

9 Q. Now, you say prematurity and
10 intrauterine growth rate restriction. You're
11 talking about small babies for gestational age?

12 A. Yes. That's what the latter of those
13 two terms means. There's one chapter in the book
14 about that concept, and here we're dealing with
15 chronic hypoxia causing poor growth.

16 Q. And what percentage is caused by
17 prematurity and which percentage is caused by
18 intrauterine growth rate restriction, if you
19 know -- the percentage of CP is caused by
20 prematurity?

21 A. Of those two, it's probably 80 percent
22 prematurity, 20 percent growth restriction. It's
23 predominantly prematurity, but I lumped them

1 together.

2 Q. So 30 to 40 percent is caused by what
3 you're calling complications of prematurity,
4 right?

5 A. Yes.

6 Q. Now, if I look in the rest of this
7 text, I'm going to find a chapter, apparently, on
8 prematurity?

9 A. Yes, a doctor from the University of
10 Chicago.

11 MR. BECKER: While he's looking, I'm
12 just going to run into the bathroom here.

13 MR. BULLOCH: I could use a little
14 break, too. Why don't we take a five-minute
15 break.

16 (Discussion off the record.)

17 MR. BULLOCH: Let me ask you this one
18 question before I take a break, because I could
19 use one, and I'm sure the court reporter could use
20 one, too.

21 Q. Doctor, I'm handing you the table of
22 contents from the upcoming release of your
23 textbook, Perinatal Causes of Cerebral Palsy, and

1 you told me a moment ago that there was a chapter
2 in there on prematurity. Could you point out to
3 me what chapter you're referring to that deals
4 with complications of prematurity?

5 A. It's this bottom one on the first page.

6 Q. So the chapter you're referring to that
7 would deal with complications of prematurity is
8 captioned the Panorama of Cerebral Palsy After
9 Very and Extremely Preterm Birth: Evidence and
10 Challenges, correct?

11 A. Yes.

12 Q. And that's by a doctor you said was out
13 of Chicago by the name of Doctor Michael Msall?

14 A. Yes.

15 Q. Now, the chapter says very and
16 extremely preterm birth. Was Matthew very preterm
17 or extremely preterm?

18 A. No.

19 Q. In fact, Doctor Msall described very
20 preterm as less than 32 weeks, correct?

21 A. That's reasonable.

22 Q. And extremely preterm as less than 28
23 weeks, correct?

1 A. That's reasonable.

2 Q. And Matthew was dated by Dubowitz and
3 other methods as about 35 to 36 weeks at birth,
4 correct.

5 A. Most references in the chart say 35.
6 I've seen 34, I've seen 36, but I'm going to say
7 35, give or take a week.

8 Q. And I'll compromise with you, Doctor,
9 35 is appropriate.

10 So I asked you the question -- well,
11 maybe I didn't ask you the question. Do you
12 believe that the information in this chapter is
13 related to Matthew Wagoner?

14 A. We'd have to look specifically. I'm
15 sure part of that chapter would be and part of it's
16 not.

17 Q. All right.

18 MR. BULLOCH: Let's take a break.

19 (Recess taken.)

20 BY MR. BULLOCH:

21 Q. Doctor, before we broke we were talking
22 about the preface that you had generated in the
23 Clinics in Perinatology, and we've also marked an

1 exhibit marked Hermansen No. 2, Exhibit No. 2,
2 that I believe you represented to me that you
3 drafted after you read Doctor Alder's testimony,
4 correct? It would be this document --

5 A. Yes.

6 Q. -- I'm showing you? And in that
7 document you listed various causes of cerebral
8 palsy, correct?

9 A. Yes.

10 Q. Now, the document that you did in the
11 Clinics in Perinatology, you said that you
12 analyzed data from the authors that were in the
13 issue -- in that particular issue of Clinics in
14 Perinatology, right?

15 A. Yes.

16 Q. You developed this list --

17 A. Yes.

18 Q. -- on Exhibit 5 after analyzing all the
19 articles that were contained in that volume,
20 correct?

21 A. Yes.

22 Q. Where did this information come from
23 that's been marked as Exhibit 2? And I'll show it

1 to you, again, sir, so you can --

2 A. I had a draft of the preface on my
3 computer, and I went back to that draft to come up
4 with this list.

5 Q. Okay. Now --

6 A. I did that yesterday.

7 Q. Some of the numbers are different?

8 A. That's because it was a draft, and the
9 final, it got a little bit changed, but it's not
10 much different.

11 Q. You don't have a copy of the final
12 draft?

13 A. No, I didn't have it in my computer. I
14 looked.

15 Q. Who would have made the changes between
16 the draft you have in your computer and the final
17 draft that is being published?

18 A. Oh, I did. I did. Those numbers
19 aren't precise. In fact, that's what's better
20 about the final draft is it gives some leeway into
21 every category, just to point out the lack of
22 precision. I don't think you're going to find many
23 significant differences.

1 Q. I think you mentioned earlier that a
2 neonatal hypoxic event that's sufficient to cause
3 PVL brain damage is rare, is that correct?

4 A. Yes.

5 Q. And, in fact, you didn't list that as
6 one of the causes of PVL brain damage or CP in
7 either your article or this document that we
8 marked as Exhibit 2, correct?

9 A. Well, PVL would be a complication of
10 prematurity. Premature CP cases are to a large
11 extent PVL.

12 Q. Okay.

13 MR. BECKER: Are you going to mark
14 those articles?

15 MR. BULLOCH: No, I'm going to give
16 those back to the Doctor.

17 MR. BECKER: Be sure to give them back
18 to him.

19 Q. The injury that -- in your report that
20 you believe Matthew Wagoner sustained is PVL brain
21 damage, correct?

22 A. At this point it appears not. It
23 appears he does have some white matter damage, but

1 some people with more expertise than I are
2 concluding it's not PVL, as I understand it. At
3 least --

4 Q. Where are you obtaining that
5 information from?

6 A. From expert reports, and I even think
7 Doctor Alder in his deposition. I don't think he's
8 calling it PVL, and I know the neuroradiologist
9 says it's not PVL, that there's white matter damage
10 in the brain, but it's not PVL.

11 Q. Well, one of the things you said causes
12 PVL is hypoxic ischemic injury, correct?

13 A. I said ischemia to the brain.

14 Q. And does white matter injury that you
15 believe Matthew is now suffering from, what is the
16 cause of that white matter damage?

17 A. The pneumothorases on the evening of
18 the 25th.

19 Q. Okay.

20 A. Associated with severe hypoxia over a
21 prolonged period of time.

22 Q. Do most babies that suffer from
23 asphyxia or hypoxic ischemic events end up with

1 cerebral palsy?

2 A. No.

3 Q. In fact, most babies that have hypoxic
4 ischemic injuries are totally normal, is that
5 correct?

6 A. Well, I -- I don't like it when you put
7 the word injuries in there. Injuries implies that
8 damage occurs. But most babies following birth
9 asphyxia with hypoxia and ischemia turn out normal.

10 Q. Okay.

11 A. But once they are injured, it's a
12 little awkward saying they're normal. Injury to me
13 implies damages.

14 Q. Good point. Do most children that
15 experience hypoxic events -- and I'm talking about
16 babies that experience hypoxic events in
17 hospitals -- do they end up with white matter
18 damage consistent with what Matthew has?

19 A. No.

20 Q. Do most babies that experience
21 pneumothoraxes experience any type of white matter
22 damage consistent with what Matthew has sustained?

23 A. Not most, but many do. It clearly is a

1 risk factor. But most will recover, but many
2 suffer damages.

3 Q. Well, do you have any idea of what
4 percentage of children with pneumothoraxes end up
5 with -- I'm sorry -- let me go back. Yeah, with
6 pneumothorax experience white matter damage?

7 A. I'd have to look up in the literature.
8 I think I would probably find numbers like 15 or
9 20 percent, but I am not confident with that
10 estimate. I might be off quite a bit there.

11 Q. In your CV, you were a participating
12 investigator in Vermont Oxford Network in 1997,
13 correct?

14 A. Yes.

15 Q. Have you been involved with that
16 organization since that time?

17 A. Not really.

18 Q. What is the Vermont Oxford Network?

19 A. It's a large coalition of hospitals
20 sharing outcome data.

21 Q. Respected entity?

22 A. Yes.

23 Q. Well-controlled studies?

1 A. Some, yes.

2 Q. Do you know a Doctor Soll?

3 A. Yes.

4 Q. S-O-L-L?

5 A. From Vermont.

6 Q. There is a database, Vermont Oxford
7 database, correct?

8 A. Yes.

9 Q. Are you familiar with the Vermont
10 Oxford database in which there were 3,505 infants
11 between 1,400 and 1,500 grams, these are babies
12 with RDS, and significant proportion -- or a
13 proportion, I shouldn't say significant, I retract
14 that -- but a certain number of those children
15 never received surfactant?

16 A. That's probably true.

17 Q. Do you believe that physicians that
18 failed to administer surfactants to these tiny
19 babies are negligent?

20 A. You would have to look at each
21 individual case to decide that. I would hope that.

22 MR. BECKER: Let me just stop there and
23 say that we'll produce Doctor Hermansen for a

1 continuation at a mutually agreed date, and I
2 apologize for leaving early.

3 MR. BULLOCH: That's all right. We
4 will finish this up later. Have a safe trip back.

5 (The deposition was adjourned at 11:37 a.m.)
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C E R T I F I C A T E O F W I T N E S S

I, MARCUS C. HERMANSEN, M.D., do hereby certify that I have read the foregoing transcript of my testimony, and further certify that it is a true and accurate record of my testimony (with the exception of the corrections listed below):

Page	Line	Correction
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1. The first part of the document is a header section containing the title "THE EFFECTS OF THE 2008 FINANCIAL CRISIS ON THE UK ECONOMY" and the author's name "JAMES H. M. SMITH".

[illegible]

[illegible]

MARCUS C. HERMANSEN, M.D.

Sworn and subscribed to before me this _____ day
of _____, 2006.

Notary Public

My Commission expires: _____

1 CERTIFICATE

2 I, Pamela J. Carle, Registered
3 Professional Reporter, do hereby certify that the
4 foregoing is a true and accurate transcript of my
5 stenographic notes of the deposition of MARCUS C.
6 HERMANSEN, M.D., who was first duly sworn, taken
7 at the place and on the date hereinbefore set
8 forth.

9 I further certify that I am neither
10 attorney nor counsel for, nor related to or
11 employed by any of the parties to the action in
12 which this deposition was taken, and further that
13 I am not a relative or employee of any attorney or
14 counsel employed in this case nor am I financially
15 interested in this action.

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Pamela J. Carle, CCR, RPR