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	3	AND THROUGH HIS NATURAL] GUARDIAN AND PARENT AS NEXT]
	4	FRIEND, DAWN MacADAMS,
994 - 2000 	5	Plaintiffs,
e e constante e	6	vs.] CIVIL ACTION NO.
- Annor	7	BLAKE CHRISTOPHER POLEYNARD,] [JURY DEMANDED]
	8	CLINIC, P.C.,
	9	Defendants.
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. ð,	11	DEDOCTATION OF MICHAEL C DADERCDY N D
	12	January 25, 1999
i al	23	500 Marquette Avenue, Northwest, Suite 280
: :	14	Albuquelque, New Mexico
	15	DIDCHANT TO THE GEODGIA CIVIL DRACTICE ACT this
	16	deposition was:
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	18	Attorney for the Plaintiffs
- 194	29	REPORTED BY: DANNA SCHUTTE EVERETT, RPR, NM CCR #139
	20	Professional Court Reporting Service
	21	Albuquerque, New Mexico 87102
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MR. BERGEN: This will be the deposition of 1 2 Michael Radetsky, M.D., taken in the MacAdams vs. Poleynard case. The deposition is taken pursuant to 3 notice and agreement of counsel, if any, for all purposes 4 allowed under the Georgia Civil Practice Act. 5 Reserve all objections except as to form and 6 responses, if that's agreeable. 7 MR. MULLER: Yes. 8 MR. BERGEN: Dr. Radetsky, I assume you want to 9 read and sign your deposition? 10 I would like to. THE WITNESS: 11 MR. BERGEN: It's agreeable to sign before a 12 notary here in this state or if you're traveling. 13 That's fine. THE WITNESS: 14 15 MR. BERGEN: No problem. THE WITNESS: Did you want me sworn in? 16 MR. BERGEN: We'll do that in just a minute. 17 We're going to stipulate to the qualifications 18 of the court reporter and the means and manner of taking 19 deposition. 20 Are there any other qualifications that need to 21 be stated? 22 MR. MULLER: That's fine. 23 24 25 SANTA FE OFFICE

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* * * * * * * * 1 MICHAEL S. RADETSRY, M.D., 2 after having been first duly sworn under oath, was 3 questioned and testified as follows: 4 EXAMINATION 5 BY MR. BERGEN: 6 Dr. Radetsky, as you know, I'm Fred Bergen. 0. Ι 7 represent Mrs. MacAdams on behalf of her son in this 8 lawsuit that's been filed in Savannah. 9 I sent you a notice to take your deposition, and 10 I understand you brought your file materials connected 11 12 with this case. Is that what you brought with you today? Yes, sir. 13 Α. Let me just do this. Let's go through the 14 Ο. Notice to Take Deposition, which I'll attach as Exhibit 1 15 to this deposition. 16 (Exhibit 1 marked.) 17 And do you have with you all letters you have Ο. 18 received from Mr. Muller or any member of his law firm or 19 any representative of the Illinois National Insurance 20 21 Company related to your review of materials relative to 22 this case? 23 Α. Yes. Where is that correspondence? 24 Ο. They're all in my working file. 25 Α. In fact, just

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to orient you, these are the records I was sent, 1 including depositions, and then my working file has 2 everything else in it. 3 Ο. It might be easier, instead of going through 4 each and every one, I'll just look at what you have, and 5 if 1 have any guestions --6 MR. MULLER: Actually, I think, Fred, I think to 7 speed things up, I think there's probably a -- Well, 8 actually, is this all one thing? 9 That's Johns Hopkins. 10 THE WITNESS: MR. MULLER: I thought everything was mixed in. 11 That's the Johns Hopkins records you've been Ο. 12 provided. Depositions? Deposition of Dawn MacAdams, 13 Harvey Kessinger, Gail Kessinger, Blake Poleynard, M.D., 14 Fernando Perez, M.D., Dr. Meislin, M.D., Charles 15 Schleien, M.D. Okay. So those are the depositions you 16 reviewed in this case? 17 Yes, sir. Α. 18 Ο. And then some document production, Schleien 19 hospital records; it looks like more Schleien hospital 20 records; Dr. Cohn's records of Ryan MacAdams. This is 21 from the Greater Baltimore Medical Center records, the 22 Memorial records. Anything else? 23 24 Α. And then in my working file, what I did was to take out certain records so that there would be more 25

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available to me. Here's all the St. Joseph's records here for the two emergency department visits, and then selected records from Memorial Hospital and from Johns Hopkins, which I pulled out just to have in my working file.

Q. Okay. And then there's also Montgomery General
Hospital records. Also, I see some literature search,
some articles. What else?

Α. Yes. Well, your request of me was to bring all 9 medical or scientific literature or sources that I used 10 11 in my analysis of Ryan. I must tell you, sir, in all frankness, since my area is infectious disease, issues 12 regarding severe infections, meningococcal infections, 13 and children with fevers are part of my daily scholarly 14 life, and, therefore, I brought to bear much more in 15 terms of general literature than is in this folder. 16

But I did bring what I thought were either the leading articles or articles which I felt shed light on the case. But I just want to make the statement that this is not an exclusive compiling of articles. These are the articles I felt were most relevant.

Q. I understand. I take it, then, because of your area of interest and expertise, you have like a working file on pediatric fevers and things of that nature, but case specific for this file, would these be the articles

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1 that you have referenced?

A. well, these are the article's I feel are most *3* relevant.

Q. Okay.

4

A. But I do use information contained in other
articles because they're part of my daily scholarly life.

7 MR. MULLER: Also, I got the impression from his 8 answer there are articles that may not be in a file out 9 there that he's seen that could come to bear on it, but, 10 obviously, he couldn't have all of them with him today 11 because he reads them.

12 Q. There is a vast amount of literature on the 13 subject matter. I understand that. Let's just look at 14 ones that you've got with you today.

A. Sure. I'll just give you the lot of them, andthen you can certainly go through.

Q. In one of the copied articles, you have some handwriting on the "Epidemic Meningococcemia and Purpura Fulminans with Induced Protein C Deficiency."

20 A. Yes.

Q. Is that relative to this case, or is that just some general working knowledge, you had your handwriting on top of that article?

A. When I first reviewed that case, I took notes on the article for myself, and they're, of course, on the

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original, so they copied when I made a Xerox copy of 1 them. 2 Why don't we do this, if you don't mind. I'd Ο. 3 like to get a copy of these. And why don't I just get 4 you to read the titles and lead authors, and then we can 5 move on. Okay? 6 Α. Okay. Sure. 7 I'll get a copy before we leave today. 8 Ο. "Osteonecrosis Following Meningococcemia and 9 Α. Disseminated Intravascular Coagulation in an Adult: 10 Case Report and Review"; lead author, Wayne N. Campbell. 11 "Epiphysiometaphyseal Changes in Children after Severe 12 Meningococcic Sepsis"; lead author, Francisco Fernandez. 13 "Late Sequelae of Infantile Meningococcemia in Growing 14 Bones of Children"; lead author, Heidi Patriquin. 15 "Skeletal Lesions Following Meningococcemia and 16 17 Disseminated Intravascular Coagulation"; lead author, Minhard Robinow, "Incidence of bacteremia in infants and 18 children with fever and petechiae"; lead author, Kenneth 19 20 Mandl. "Epidemic Meningococcemia and Purpura Fulminans with Induced Protein C Deficiency"; lead author, Darlene 21 Powars, "Experimental treatments of meningococcal 22 sepsis"; Michael Levin. 23 Then there is a series of slides which I 2425 compiled for a talk some time ago that recount

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information from a number of other articles, and I just
 gave you a copy of the handout.

Q. Okay.

3

15

A. "Febrile children with no focus of infection: a
survey of their management by primary care physicians";
lead author, Ronald Jones. "Practice Guideline for the
Management of Infants and Children 0 to 36 Months of Age
With Fever Without Source"; lead author, Larry Baraff.
And finally, "The febrile infant and the assumption of
risk," and I'm the author.

11 Q. You've got some correspondence in your next 12 section of your file?

A. Well, actually, to begin with, I had some notesthat I took that you wanted me to bring.

Q. All right. Thank you.

16 All right. The notes appear to be your notes
17 taken from the ER admissions -- the two ER admissions at
18 St. Joseph's.

19 A. That's correct.

Q. We'll just get a copy marked as an exhibit. If
you need to refer to them in the deposition, that's fine
A. The remainder are invoices, correspondence, and
a canceled check.

24 Q. When were you first contacted in this case? Was 25 it May of 1998?

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1	A. I honestly don't know, sir. It would be around
2	the time of the first correspondence.
3	Q. Okay. That's what I was trying to make out. It
4	may be May 14th. It says, "I appreciate your willingness
5	to look at the file for me." For Mr. Muller.
6	A. Yes.
7	Q. The only invoice that you have submitted so far
8	is one it looks like May 27th of 1998. Is that the
9	only invoice so far to date?
10	A. Yes, that's the only one.
11	Q. I take it there's been additional work since
12	then?
13	A. Yes.
14	Q. And your charge was \$350 per hour for case
15	review?
16	A. It is.
17	Q. Does it change in any degree for depositions?
18	A. Deposition time itself is \$400 an hour, and
19	trial testimony at \$450 an hour.
20	Q. What about travel?
21	A. Travel is \$350 an hour with a maximum of 12
22	hours per day.
23	Q. So it's not portal-to-portal, but it could be
24	for travel, I take it?
25	A. That's correct.

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So during travel and trial testimony, it's] Q. generally 12-hour days? 2 Yes. What I do is, actually, I bill Α. door-to-door with a maximum of 12 hours a day, and that's 4 in addition to the actual time I spend testifying. 5 And it usually works out that way. 6 Okay. Let's put this aside. I'll get copies of ο. 7 these letters, as well, later when we take a break or a after the deposition. 9 At the time of your first invoice, May 27th of 10 1998, were your opinions final in this case? 11 Α. Let me just look at that first invoice, if I 12could. 13 14 0. Sure. 15 Α. No. My opinions weren't final in the sense that 16 I had not read the depositions of any of the participating individuals, both the family and the doctors, so my expression of an opinion was based, 18 really, just on my record review, And then I think I 19 nentioned at the time that I would like to reread the 20 further depositions, and if there were a revision in my 21 opinions, then I would certainly be contacting them if 22 chat took place, based on a review of depositions, which 23 I had not at that time received. 24 25 Okay. You provided me a copy of your CV. I Q.

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1 understand the one I have in my hand is the most current 2 one. Correct?

A. That's correct.

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2.36

Q. Let me see if it compares to the one that I have, if I can find it. This one was apparently faxed -it was probably a little bit out of date. It was May of 1998 -- from you, I guess to Mr. Muller. Would the only additions be with regard to publications?

9 A. I think there's been an addition in
10 publications. I think I've gotten a couple more honors.
11 I haven't gotten the Nobel Prize or anything like that.
12 Q. Okay. I understand. Let me ask you this now.

What's your present address now, home address?

A. My home address?

Q. Yes.

16 A. 1217 Rockrose Road, Northeast, in Albuquerque,17 87122.

18 Q. And you're still at Lovelace Health Syseems, I 19 take it?

A. Yes. Actually, we say Lovelace.

21 Q. Lovelace, I'm sorry.

A. It's the way the founder pronounced his name.

23 Q. He's the one that gave the bread, so you do it

24 the way he wants to hear it.

A. That's right. That's the way.

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I Q. I see you went to law school at some point. It 2 looks like you graduated from Harvard in 1967. Is that 3 correct?

4 A. Yes. And I went to law school for one year, in 5 1967 to 1968.

Q. Why is that? What happened there?

Well, I went to law school because I thought 7 Α. that that's what I was going to be doing with my life. 8 At the end of that first year of law school -- And it was 9 a tumultuous year because of the fact that Martin Luther 10 King and Robert Kennedy were both shot during thac year, 11 It was the only year in the history of Harvard Law School 12 that final examinations were not given because of the 13 multiple murders of esteemed people, and it was a time of 14some disillusionment for some of us, so I took a leave 15 of absence and enlisted in the United States Peace Corps. 16 Q. And you went into the Peace Corps. And you're 17 18 going to have to help me with this word. I was a malariologist in the Peace Corps. 19 Α. 20 Ο. Was that for malaria or something?

A. Yes, there was a malaria eradication campaign run by the World Health Organization in Thailand, and I was a middle-level bureaucrat from this particular system.

25

6

Q. How old were you then?

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Twenty-one, twenty-two. Α. 1 Q. And then you went -- What is this? What is the 2 Alternative National Service (Conscientious Objection)? 3 Α. I was drafted after I left the Peace Corps, but 4 there was a provision in the Selective Service law that 5 was passed after the Second World War that if, in 6 relationship to a supreme being, an individual was 7 conscientiously opposed to war in any form, and if thac 8 belief was validated by their local draft board, they 9 would be granted alternative service in the national 10 interest outside of the military, and that was my 11 situation. 12 So, in other words, you objected to the Vietnam 13 0. War? 14 15 Α. No, I objected to my participation in war in any 16 form. 17 Ο. Regardless of whether it was Vietnam or any type of war? 18 That's correct. 19 Α. Were you drafted? 20 0. I was drafted, but rather than going into the 21 Α. military then, I worked at the Children's Hospital in 22 23 Boston. 24 Q. Did you avoid the draft? 25 Α. I didn't avoid the draft. The way the Selective

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Service law read was that if you were drafted but you 1 were a conscientious objector, you would perform national. 2 service in a nonmilitary capacity for the duration of 3 your obligation. 4 Would law school grant you a college deferment Ο. 5 for the Vietnam War? 6 7 Yes, education granted you a deferment as long Α. as you were continuously in an educational institution. 8 But when you were not, then, of course, you were fully 9 eligible to be drafted. 10 And so when you took your leave of absence from Q. 11 the Harvard Law School, was that on good standing? 12Yes, it was. And, in fact, I was invited to Α. 13 come back, but as one thing leads to another thing, I saw 14 the light and went into medicine instead. 15 But that leave of absence was a voluntary 16 Ο. 17 decision by you? Α. 18 It was. 0. If you did not go into the Peace Corps or the 19 conscientious objection program, then you would have been 20 required to serve a military commitment? 21 22 Well, I was drafted, and if I had not been Α. granted the status of conscientious objection, then I 23 24 would have served military service rather than alternative service. In either case, I was drafted. 25

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	17
r1	0. Wh× WiP you go to school in Canada in 1973?
\sim	A. Right. When I was applying to medical school, I
т	was accepted both in the United States and in Canada, and
4	CEnapa Seing the only country wPich has total reciprocity
ហ	with the UniteD States, so yow're not a foreign medical
Ś	${\bf t}$ ${\bf u}{\bf \pi}$ aduate, and , the ricans would go to Canadian schools, and
٢	Canadians woulΩ go to Am™rican schools My Wacision to
ω	go to CanaDa was that the tuition was \$800 a year
σ	Q Was it an <u></u> thing to Do wit > awoiDing ODD Draft at
0	all?
Ч	A No I had already been drafted and had served
12	my obligation Nut as a conscientious onjeceor in
m	alternati e e ze r eice So this was after Hy Draft
44	obligation.
ы П	Q. Did you haws to take the FLEX sxam if you go to
Тe	Canada?
17	A. No. You take the national examinations just the
8	way American students do. It's not considered to be a
6	∎¤parate Bysten from the poin of Jiew of li<∞nswre
0	ຊີ Th⊵n I take it it aທຸµະars th≞c aftະr meDical
12	school anû your rotating internahên, yowr focua Qecame in
[7]	tbe ares of p epiatrics and in Anfectious Dispases
m S	A. That's correct.
4	Q. What did you do as the malariologist in
ى د	Thailand? Obviously, you worked not you weren t a
* * * * * * * * * * * * * * * * *	SANTA FE OFFICE DEAN 119 East Marcy Suite 110 Marchen W. Suite
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doctor at that time. What were you doing, exactly? I'm
 trying to get a feel for that.

A. The program I was involved in was run by the World Health Organization, and we were given three months of training in Manila at the WHO center for malaria eradication there to acquire the organizational and laboratory skills whereby we could be put into an existing program and contribute to the work of eradication of malaria.

In my area of Thailand, which was up near the 10 Laotian border where Laos, Burma, and Thailand come 11 together, i supervised DDT spraying crews, supervised 1213 crews that would obtain blbod samples from villagers to see who had malaria, and then the distribution of malaria 14 medication to eradicate the infection. And this was the 15 three-pronged attack that the World Health Organization 16 had used successfully in other countries. 17

18 Q. And that was satisfying the conscientious19 objection issue?

A. No. When you were in the Peace Corps, you were given a deferment for the draft. The minute I left, the Peace Corps, I was drafted.

Q. And then you went into the alternative national service?

25

A. That's correct. I worked in a hospital in

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1 Boston.

Q. And that was a -- Was there a local draft board that had to approve that or --

Yes, the local draft board, which for me was in Α. 4 Denver, had to do two things. They had to grant you the 5 status of being a conscientious objector -- and I was the 6 first, and to my knowledge the only, one ever granted by 7 8 my draft board -- and secondly, once you were drafted, 9 you had to work in a civilian capacity in a way that contributed to the national service, really, and so they 10 11 had to approve your site where you found work that was acceptable under the guidelines that they published. 12

Q. Okay. Then in 1982/1987 you were the associate
director of infectious disease at Denver General?

A. Actually, the Denver Children's Hospital.

16 Q. Did you have any interaction with Dr. -- Is it 17 Barkin?

A. Well,, I knew Dr. Barkin during the time that I
was in Denver. He's still in Denver, but I moved away.
Q. Then you moved on to Tucson?

A. Well, I was engaged to be married, and my wife, who is a pediatrician, was going to do her internship in Tucson, so I certainly wanted to be near her, and made the move.

25

15

Q. Right. I understand. Did you know Dr. Harvey

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Meislin out there then? 1 Α. Yes, I did know him. 2 Have you kept up with him since? ο. 3 No, I haven't. Α. 4 Did you always get along with Dr. Meislin in 5 Ο. 6 working out in Tucson? I think, in all honesty, we probably only met a Α. 7 few times. He primarily worked at University Hospital. 8 I primarily worked at Tucson Medical Center, although I 9 did work some at the University Hospital. He was 10 primarily in the emergency department. I was primarily 11 12 in the pediatric intensive care unit. So there weren't a 13 bot of places in which our practice lives overlapped. How about by consults? 14 Q. Again, it was only infrequent, I must say, in 15 Α. all honesty. 16 Q. And then back to Denver in 1989? 17 Α. Well, yes. It was my hometown, and my wife was 18 19 able to transfer her training back to Denver, so we were able to move back home. 20 Then I see where you went to California for a 21 Ο. couple years. What was the reason for that? 22 Well, my wife, after she finished her pediatric 23 Α. training, went to Johns Hopkins to get her master's in 2425 public health. She then got a job in Sacramento at the

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A Never. Q Has any state ever rewoken or suspended your Santa Fe OFFICE 119 East Marcy Suite 110 Santa Fe. NM 87501 505) 890-1049 EAX (305) 820-6349 FAX (305) 820-640 FAX (305) 820-640 FAX (305) 820-640 FAX (305) 820 FAX (305) 820-640 FAX (305) 820 FAX (305) 820-640 FAX (305) 82	ひ ひ 4
сжіme of any tyke?	5 3
Q Have you pupt seen arrystaton or convicted of any	22
A Never.	7
in any way?	20
where your privileges were DenieD, suspended, or ReGkeD	ц 1
ο xave you ewer bad anx oroalems at any hospitats	4 1
A. That's correct.	17
landed, so to speak?	9 H
Q. You were searching, anD this is where you	10
Albuquerque.	44
tbr west where we were oth from, and fourd it in	13
joa Anp we starten looking for some other position in	Ц Ц
atate conscitution. She was last in/fir∋d out of har	
A They have a DalanceD DuDget amenument to their	10
Q. Right.	ማ
California. You max bave remenerre	ω
A Yes In 1993 there was a budget crisis in	7
Q. And then back here to Albuquerque?	Q
around, and I went to California	IJ
think I ve spant Host of Hy life Sollowing Hy wife	4
in Atlanta at the Contors for Dispash Control So I	M
two plum jo23 in the UnitRO StatR3 The otYer one Deing	N
Stace Department of PuOlic Health . Wich was one of the	
21	

privileges --1 2 Α. Never. -- or your licenses? 3 Ο. Α. Never. 4 What family do you have here in Albuquerque? 5 Ο. None. Well, we have two children, but there are Α. 6 just the four of us here. 7 How old are your children? Ο. 8 Α. Five and seven. 9 Q. First marriage? 10 First and only, I hope. 11 Α. Q. Got around to having children eventually? 12 Well, we're late bloomers. 13 Α. I've got you. No family in Chatham County? 14 Ο. Some fellow we knew had the same last name, but no family 15 members from your side or your wife's side in Chatham 16 County? 17 No. My wife has a cousin who lives in Savannah. 18 Α. Who is that? 19 Ο. I'm trying to think of her name. I'm sorry. 20 Α. 21 It's Susan something, but I can't remember her last name. I believe she is, as 22 Her husband is in real estate. well. But I just can't think of their last name. 23 I'm 24 sorry. If you would ask your wife and maybe tell 25 Ο.

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23 Mr. Muller, he can let me know, or if it comes to you 1 2 later today in the deposition. Okay? Α. Sure. 3 4 Ο. Have you ever practiced in Georgia? No. 5 Α. Q. I know you've spoken in several lectures. Have 6 you ever lectured in Georgia? 7 I don't believe I have. 8 Α. Have you ever been a party to a lawsuit before? 0. 9 Is that a way of asking if I have been sued? 10 Α. Q. Well, it doesn't matter. Either way. 11 Have you ever sued anybody, or has anybody ever sued you for any 12 13 reason? I'm going to boil it down to where it might --I've been sued for medical negligence twice in 14 Α. 15 my life, but I've sued no one. Have you had any other type of litigation? Like 16 Q. contract disputes, anything like that? 17 18 Α. No. Where were the two cases where you were sued 19 Ο. 20 regarding medical negligence? These were two cases which arose during the time 21 Α. that I was a pediatric resident in my training in Denver. 22 23 Both of them occurred in the early 1980s. And I was named along with a large number of other people in each 24 of them, and I was discharged in both of them early on in 25

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1 the proceedings of the lawsuit.

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Q. What was the nature of the complaint?

The complaint in one of them was concerning a Α. 3 seven-year-old girl who had kidney failure who was 4 admitted to the pediatric intensive care unit in which I 5 was working as a pediatric resident who, in the early б morning hours after her admission, had a cardiac arrest, 7 and the allegation was that it should have been 8 anticipated and it should have been conducted in a more 9 expeditious manner. 10

The second one was a child who had an unusual 11 condition, called hemolytic uremic syndrome, who was at 12 the Children's Hospital of Denver and who went into 13 kidney failure and needed to be transferred to the 14 University of Colorado where the dialysis unit was. 15 Ι dictated the transfer summary. Later on, he developed a 16 blood clot in his hand, and it was thought that the 17 catheter that he had in his radial artery contributed to 18 the blood clot, and he lost a portion of a finger, as I 19 recall. And I was named along with everyone else who had 20 21 ever cared for him.

Q. Those are the only two times a claim has everbeen made against you?

A. That's right.

Q. That includes any sort of notice of claim,

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25 regardless of whether it concluded in suit or not? 1 I've never been noticed. 2 Α. Q. Now, you've had experience in testifying in 3 medical/legal cases serving as an expert witness before, 4 5 have you not? 6 Α. Yes. 0. Let's talk about -- On how many occasions have 7 8 you? 9 Well, I started reviewing cases in 1982 when I Α. first went into the faculty of the Children's Hospital, 10 and in the 17 years I have probably reviewed 200 to 250 11 cases overall. 12 So a little bit more than -- What's that come 13 Ο. 14 out to a year? Averages 10 to 15 a year. 15 Α. 16 Ο. That's what I was going to say. Ten is a little 17 light, and 15 is about -- Between 10 and 15 a year? And 18 where have those cases -- To your knowledge, which states have they been in? 19 Well, they've arisen from many states, frankly. 20 Α. I think the bulk of them have been in the western states, 21 22 but I have been consulted by attorneys who represented 23 clients from many states, both in the east and in the 24 southeast. 25 Ο. Let's talk about which states you recall

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Ч	warticularlx
0	A. I'm going to compose a mental map of the United
Μ	States, and then I'll go through.
4	Q. Okay.
ហ	A Hawai.i Washington state California Idaho
0	NawaDa Arizona w Mwxico ColoraDo Nyoming Montana
5	I cant recall Texar Kanras South Dakota Wisconsin,
00	Illinois Missowri, Missirsi p yi, Tennezspe, Kentucky
σ	Florida, Georgia, North Carolina, New Jørsey
0 T	Connecticut.
11	Q. Okay.
12	A OP apy Ogio. I'H Sorry, sir.
1 Э	Q. Thank you.
4	A. And I may have missed something in there, but I
ы Ц	tried to do my best.
9 H	Q I appreciety it And in wach of tbosy states
17	have you testified bx Deposition as well?
00 1-1	A I Dontt think in wach of those states J ve
<u>б</u>	tartified By deposition necessarily but I can't recall
20	Bpecifically which
5	ч Q. How øbout trials? In which statøз havø you
52	testified in courtrooms?
23	A Xawai i California Nºw Mexico IDaho
24	<olorado, florida="" gworgia="" illinois="" kansas="" north<="" td=""></olorado,>
52	<arbivencess +="" and="" are="" deliewe,="" jersey="" new="" td="" the<=""></arbivencess>
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	only one	s.
	Q.	In Georgia, where did you testify in Georgia?
	Α.	In Atlanta.
	Q.	Okay. And do you know what the name of the case
	was?	
E	Α.	Yes, Adams versus Kaiser.
	Q.	Which side were you on?
٤	Α.	I was retained by the attorneys representing the
c,	defendan	t Kaiser-Perrnanente.
1 C	Q.	Mr. Pound in Austin appeared in that case?
11	Α.	Excuse me?
12	Q.	Was that Mr. Pound, Ted Pound?
13	Α.	I believe he was involved. That was the law
14	Eirm. B	ut the lead attorney was a female attorney, and I
15	just don	't remember her name. I'm sorry.
16	Q.	Was that a meningitis case?
17	Α.	No, it was a meningococcemia case in a child
18	:hat had	a cardiac arrest in a car going to the hospital.
19	Q.	And that was in favor of the plaintiff for like
20	a hundre	d
21	А.	A considerable verdict.
22	Q.	Yeah. Do <i>you</i> remember Mr. Malone?
23	Α.	Oh, yes.
24	Q.	Was that the only case you've testified in in
25	Georgia?	
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Antonio Carlos

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Α. I think so, sir. 1 2 Ο. And what were you testifying -- What issues did 3 you testify to in that case? The main issue was the issue of causation in 4 Α. 5 that case. The allegation on the part of the plaintiffs was that the cardiac arrest that occurred meaningfully 6 led to the limb damage that this child suffered. I tried 7 to point out that the disease itself could have and 8 9 probably did cause all of the limb injury and that the cardiac arrest itself did not increase the risk of limb 10 injury. 11 Is that the only time you've testified in 12 Ο. Georgia? 13 14 Α. I think it is, sir. How many cases are you currently reviewing? 15 0. I honestly don't know. 16 Α. 17 Would it be on the 10-to-15-a-year range? Ο. 18 I think so. Sometimes there is a case that I Α. 19 haven't heard about for a year or more, and I don't know whether it's settled or not settled -- sometimes people 2.0 don't have the courtesy to let me know -- so it's hard to 21 22 estimate how many cases are truly active. 23 Q. Okay. But has that been your general case 24 review load, 10 to 15 a year? 25 I think that's the way it's averaged. I think, Α.

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as with all things, there are probably times that it's 1 been more and times that it's been less. 2 Do you know how Mr. Muller got in touch with you 3 Ο. in this case, how he learned of you, or anything? 4 5 Α. i don't know. Do you advertise your services in any way? Б Ο. 7 Α. i do not. Have you generally had a reputation within your В Ο. 9 area of expertise as someone who does get involved in 10 medical/legal cases? 11 I don't quite know how to answer that. Α. Q. Well, i mean, like if you may know someone who 12 reviews cases that you see at meetings and you might say, 13 "Well, you know, that guy reviews cases"? 14 15 Α. Actually, I never know who is involved in case reviews unless I've encountered them in the context of a 16 lawsuit. It's not the sort of thing that's talked about Ε7 at professional meetings, at least not the ones that I go 18 19 to. 20 What's your annual income regarding Ο. 21 medical/legal cases a year? 22 Well, I can't give you a -- I cannot give you a Α. 23 dollar figure, but I would say that over the years it's 24 averaged, now, about 20 percent, perhaps in some years as 25 high as 25 percent of my total income.

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Q. well, I mean, what would it be on the range of 1 Dollar figures? 2 an average year? Α. I don't think I can give you that figure, sir. 3 Q. Well, do your tax returns report 1099's or a 4 different source of income per year? 5 Α. Well, I will receive 1099's in any given tax 6 7 year, which I send to my accountant. I understand. But what I'm try to understand ---8 0. 9 I'm just throwing numbers out. I mean, do you make \$25,000? \$50,000 a year? A hundred thousand? 10 MR. MULLER: I don't think this is relevant. 11 12 MR. BERGEN: Sure is. I don't think it is. 13 MR. MULLER: MR. BERGEN: 14 Well, we can take it up with the I'm just trying to get a range of what your 16 Ο. 17 annual income has been in medical/legal cases. Well, I think the only way I can really answer Α. 18 19 it is the percentage answer that I gave you, sir. I 20 can't give you any numbers. 21 Ο. Who's your accountant? 22 Α. I don't think I'm going to provide you with that information, sir. 23 24 So you refuse to tell me that information? 0. 25 I'm not going to provide you with that Α.

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1 information.

Do you know how many hours you spend a year in 2 Ο. medical/legal cases, review and otherwise? 3 Α. Well, not exactly. This is all done on my own 4 time pretty much, either in the early morning or weekends 5 or whatever, or on a day off, like today. 6 Are you able to estimate? That's all. 7 Ο. No, I can't give you a real estimate. 8 Α. As to your professional life, how do you break 9 0. it down into academics, hands-on, clinical, things of 10 11 that nature? The Lovelace Clinic that I'm associated with was 12 Α. founded in the 1920's by a fellow from the Mayo Clinic. 13 He wanted to make it the Mayo Clinic of the Southwest. 14 And it has always been an independent entity without the 15 strong presence of medical students or house staff, so we 16 do not have any intermediary individuals between us and 17 the patients. Consequently, of my 120 percent time, I 18 will spend about two-thirds of my time directly seeing 19 patients, both -- actually doing all three things: 20 21 ambulatory pediatrics -- I have my own clinic, my own patients who grow up in my office; I do their primary --22 hospital pediatrics, including a busy nursery, a busy 23 24 hospital ward; and I'm the only one trained to do intensive care medicine. 25

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Q. Is it a PIC unit, or is it considered a nursery? A. It's a level 2 nursery. It does not -- Let me put it this way. We have not elected to have long-term ventilated patients there.

5

Q. Okay.

A. And then I do consultations in infectious
diseases. I also teach at the university, where I will
a attend on their pediatric ICU and step-down unit and
g teach in infectious disease, and then I have some
scholarly activities that I'm involved in. But since I'm
also the department chair, there's a lot of
administrative work that I must do, of course.

13 'Q. So probably two-thirds patient care, one-third 14 academic and administrative?

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A. I think that's a good breakdown.

16 Q. In the cases that you have testified in and 17 appeared as an expert witness, have you testified in any 18 cases where you've expressed an opinion that earlier 19 intervention would have made a difference in the outcome 20 of the patient with meningococcal infection?

A. I don't think I've been involved in a case in
which I felt the facts could lead me to that conclusion
regarding a meningococcal infection.

Q. Of the cases where you have testified in, howdoes it break down plaintiff versus defendant, patient

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versus doctor or medical center, whatever? 1 Α. Sure. 2 3 MR. MULLER: Are you talking about he's 4 testified --Uh-huh. MR. BERGEN: 5 -- in either deposition or at MR. MULLER: 6 7 trial? Yeah, deposition and trial. MR. EERGEN: a Understanding, since I don't advertise, people 9 Α. 10 just call me, and the proportion in which they call has very little to do with myself, it's just who ends up 11 12 contacting me, I'd say in terms of review of cases, probably 80 to 85 percent have been sent to me by 13 attorneys representing defendants of one sort of another; 14 15 to 20 percent injured parties. At deposition, I would 15 say the disproportion increases. I'd say probably 90 to 16 17 95 percent of depositions are involved with my expressing opinions supporting defendants in a lawsuit. And I've 18 only testified one out of 20 to 25 times at trial for an 19 20 injured party. 21 What type of case was that time when you Q. 22 testified for an injured party? Bacterial meningitis. 23 Α. 24 Q. Where was that case? Right here in Albuquerque. 25 Α.

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1	Q. Do you remember the style of the case?
2	A. I do. It was called Little versus Parrino. I
3	think that's P-A-R-R-I-N-0.
4	Q. And do you know what court it was filed in in
5	Albuquerque?
6	A. I honestly don't know, sir. It was some years
7	ago.
8	Q. Of the 20 to 25 times you've actually testified
9	in court, only one occasion is where you testified for
10	the injured patient?
11	A. That's correct.
12	Q. And that was the Little case?
13	A. That's correct.
14	Q. And has most of your testimony been in cases
15	regarding causation or causal relationship or whether
16	earlier intervention would have changed the outcome? Has
17	that kind of been your area of expertise in testifying?
18	A. No, I wouldn't say so. As a practicing
19	physician, I'm called upon to express opinions on the
20	standard of care as well.
2 1	Q. Okay. Have you ever expressed any concern about
22	the medical/legal system where a plaintiff can sue a
23	defendant medical provider for compensation?
24	A. By "expressed," what do you mean?
25	Q. Well, in lectures or writings or letters to

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1 editors, things of that nature.

2	A. Well, you understand, I've gone to law school
3	for at least a year, and I honor the system by which an
4	injured party can seek compensation. I have expressed in
5	writing in letters or primarily letters, I think
6	that I feel that there is an untoward amount of
7	litigation directed towards doctors which, at least in my
8	experience, has been ill-founded in truth or science.
9	And I think in one letter I even called it rampant
10	litigation. But I think it is true that there are many
11	lawsuits that don't have merit that are filed. But I
12	certainly honor the lawsuits that do have merit.
13	Q. I think the rampant litigation statement dealt
14	with timing of meningococcemia treatment. It was a
15	response to a doctor in Texas, an article done by some
16	physicians in Texas, a lady by the name I can't
17	remember her first name.
18	A. Yes, I think you're right, sir. It was a letter
19	written following the publication of a review article
20	regarding the experience of meningococcal disease at
21	Parkland General Hospital in Dallas.
2 2	Q. Right, timing of therapy for meningococcal
23	infection. It was a letter that you wrote after a review
24	of an article by Kirsh
25	A. Yes.

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-- in which you referred to rampant litigation. Ο. 1 Α. Yes. 2 Any other articles or letters that you've Q. 3 expressed this opinion that you can refer to? 4 Α. Not, perhaps, in the same tone of voice, no. 5 You've provided me a number of articles dealing Ο. 6 7 with the subject matter relative to Ryan MacAdams on your opinions in this case. Is there any other sort of 8 standard textbook or treatise that you consider to be 9 authoritative considering this type of infection that 10 11 Ryan had? Would you define "authoritative" MR. MULLER: 12 here? 13 That he would consider to be 14 MR. BERGEN: How about that? 15 gospel. MR. MULLER: I mean, because your doctor had a 16 specific feeling about it, and I didn't know whether you 17 were defining it the same way your doctor states if it is 18 an authoritative text, and that means that every single 19 thing in that text is gospel. 20 MR. BERGEN: That's right. I mean, basically, 21 it's a term of art, and it's a term of legal consequence. 22 MR. MULLER: You're talking about everything, is 23 there a text where everything in that text is gospel? 24 MR. BERGEN: 25 Right.

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I don't think so. Dr. Meislin put it nice when Α. 1 he said textbooks are informative, but not authoritative. 2 Q. Do you agree with that proposition? 3 I would. He pointed out that many chapters are Α. 4 written with some delay before they get published so 5 they're somewhat out of date, perhaps, and also you're 6 7 limited in your space and in the depth to which you can probe in a textbook article, so you're always shying away 8 from controversies, and you're always, in a sense, just 9 reinforcing certain standard observations that may not be 1011 entirely direct. 12 Q. It's a good reference source, but there can be -- it can be out of date, things like that? 13 I think it's a good place to start if you know Α. 14 nothing about a subject, but it's not the place to end. 15 16 Ο. How about as to any literature that you consider to be authoritative -- and I'm talking about 17 literature -- in the management of a febrile infant? 18 And I notice you had Dr. Baraff's article, the guidelines 19 that was in Pediatrics, I think, in 1984 --20 21 Α. It was 1993. I'm sorry' 1993. Excuse me. -- which had been Q. 22an accumulation of guidelines that had been in the works 23 for years, as I understand. **Is** that your understanding? 2425 Α. Well, Dr. Baraff and his colleagues tried to

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create some order out of what they perceived to be 7 disorder, and come out with a viewpoint that could be a 2 source of guidance for practicing physicians, but as 3 you'll notice in the last paragraph of their article, 4 they very specifically state that theirs are not binding 5 recommendations; they're only a starting point, and that 6 7 every practicing physician should approach the individual patient with individual analysis. 8

9 I've written a couple times on the febrile
10 infant. I think that the things that I said in my own
11 publications are my best attempt to approach it. It's an
12 area in which there are now over 800 or 900 separate
13 publications, and, therefore, it's one for which no
14 single publication can stand alone.

Q. Okay. And you anticipated my next question. I guess you saw me look at your journals. I think you did have one of your journal articles which dealt with the clinical evaluation of the febrile infant and certain recommendations that one might want to go through to evaluate the febrile infant.

A, There have been two articles I've written aboutthat subject.

Q. I see one on primary care. What was the other one?

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A. I actually provided you a copy of it. Yes. It

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1.	was published in the Current in 1996.
2	Q. Is that the assumption of risk?
3	A. That is the one.
4	Q. Have you provided Mr. Muller with any written
5	reports regarding your opinions in this case?
б	A. No.
7	Q. Did you consult with any physician in
8	formulating your opinions in this case?
9	A. No.
10	Q. Did you discuss it with any other physician or
11	colleague?
12	A. No.
13	Q. To date, how much time do you think you have
14	spent in reviewing the materials and formulating your
15	opinions in this case? To date?
16	A. The only invoice I sent was for a total of
17	seven-and-a-quarter hours. I think that within the
18	additional materials sent to me, the depositions, the
19	preparation for this deposition, the travel time, there's
20	probably an additional 15 to 20 hours involved.
21	Q. Okay. So 23 hours, 24 hours would be
22	comfortable?
23	A. I think that order of magnitude, without being
24	held to an exact number.
25	Q. Sure. I understand. I'm just trying to get a

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1 reference of how much time you've spent.

A. But I must say, in all fairness, there were a considerable number of depositions, and the follow-up hospital records here were quite complex because of the nature of the child's injury.

Q. Did you make any notes in the deposition7 transcripts or anything?

In the mother's deposition, I did some 8 Α. underlining -- excuse me -- highlighting, and I don't 9 think I did it in anyone else's deposition than hers. 10 And then in some of the medical notes here I did some 11 This would be an example on the page on 12 highlighting. 13 which I highlighted just some references to the rash, which, of course, is a matter of contention in the 14 lawsuit. 15

Okay. As to practice guidelines of a febrile Ο. 16 infant, a fever for an infant, I notice Dr. Baraff and 17contributors talk about a low range of 100.4 rectal as 18 being a fever, and I notice in one of your articles you 19 give 100 degrees by mouth and 101 by rectum. I mean, I'm 20 asking you, where do you consider the cutoff point for 21 fever of an infant, definition of a fever of an infant of 22 Is the scale zero to 36 months? **Is** that the 23 Ryan's age? range you Like to use, or is it a shorter time period? 24 25 I think the zero to 36 months is too large a Α.

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span because a freshly minted newborn you approach differently than a child who's four months of age, that this child almost was, or a child who's three years of age.

But I should tell you, in all fairness, that the definition of fever has no consistency about it, and it has no agreed point. It is truly one of those arbitrary thresholds. I will tell you currently what the two most common breakpoints actually are.

Q. Okay.

10

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A. One is 100.6 degrees Fahrenheit -- that's 38 degrees centigrade -- and the other is 101 degrees Fahrenheit, or 38.3 degrees centigrade. But in truth, I think the only reason that they're chosen is because they are round numbers. One is 101, a round number Fahrenheit number; the other is 38, a round centigrade number.

Q. From what we have in the chart regarding Ryan's first emergency room admission, could you and I agree that, from what's in the chart, at all times Ryan had -would classify as having a fever?

A. He would.

Q. And by that definition, then, at all times, from what we see in the chart, Ryan was a febrile infant in the emergency room at St. Joseph's Hospital on the afternoon of May 7th, 1992?

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Α. He was. 1 And based upon your review of the chart, can we 2 Ο. agree that the fever that Ryan had was without known 3 source during that first emergency room visit? 4 Α. Yes. 5 With infants of Ryan's age that are febrile, Ο. 6 isn't there an -- there is a risk of bacteremia? 7 Α. There is. 8 And am I correct in this statement, that Q. 9 meningitis is almost always preceded by a bacteremia? 10 That's correct. Α. 11 And isn't it agreed or thought medically in your 12 Ο. area of expertise that one of the reasons why you want to 13 determine if the patient may have bacteremia in an infant 14of Ryan's age is so that you can treat it so it won't go 15 16 into possible meningitis? Am I making sense there? Yes, although I think there's a broader Α. 17 perspective than just that. Bacteremia most of the time 18 is self-resolving, but there will be children who will 19 have a progressive illness -- that is, septicemia -- or 20 in whom the bacteremia will lead to a focal infection. 21 And the idea there is to either try to terminate the 22 bacteremia before it progresses to a more severe 23 generalized infection or before it actually causes focal 24 infection, 25

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Q. Okay.

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A. Understanding that most cases of bacteremia will *3* resolve on their own.

4 Q. When you talk about a focal infection, you're 5 talking about a specific organism causing the disease?

A. No, what I meant by a focal infection, meaning a
focal part of the body or a specific part of the body.
8 For example, meningitis, a bone infection, a series of
9 places in which the infecting organism may lodge and then
10 cause disease in a particular organ.

Q. Well, Ryan had an occult bacteremia, did he not? A. Well, a blood culture was not performed, as you know, and, therefore, no one can know for sure whether on his first visit he did, in fact, have a bacteremia.

Q. Knowing what we know now, looking back retrospectively, would you agree that more likely than not he did have a bacteremia when he first went to the hospital?

19 A. I think he had a bacteremia at some point during 20 that three-hour stay in the emergency department, 21 because, as you point out, seven hours after his 22 discharge he's back with septicemia. And I think that in 23 retrospectively weighing the chances of a bacteremia 24 existing at the first ER visit, it would come out greater 25 than 50-percent likelihood.

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Q. Would you agree, then, in infants older than two
 or three months of age the higher the temperature, the
 higher the risk of bacteremia occurs?

4

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A. Yes.

Q. In the converse, of infants less than two to
three months of age, the height of the fever has no
relationship to the relationship of bacteremia?

8 A. I think I have to revise that now and probably9 say less than one month of age.

Q. Okay.

Α. The most recent information -- I think 11 Dr. Baraff talks about this as well. A child who is one 12month of age or less is really quite different than a 13 child who is older than one month. And probably the 14 three-month differentiating point, which has been used 15 since the early 1970's when the first articles came out 16 about the febrile infant, is probably the wrong 17 differentiating point. It probably should be one month 18 of age. 19

And I might say in passing, sir, that although I personally feel that the higher the fever the greater the risk of bacteremia, there are some very capable people who have published quite the opposite of that.

Q. But as to what you think, the height of the temperature has **a** correlation to the increased risk of

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bacteremia if the child is three months -- greater than 1 2 three months? Α. Yes, I believe so. But probably in light of 3 these conflicting publications, it's probably less 4 5 reliable than I thought it was at one point. Q. Well, isn't it true that the medical literature 6 talks about that with each increase in degree of 7 temperature in an infant, the increased degree of risk of 8 bacteremia exists in three-to-eight-month age infants? 9 10 Α. I don't think I understood that, sir. I'm 11 sorry. Okay. If the temperature -- high temperature in 12Q. a child who is three to eight months -- using that window 13 there --14 15 Α. Three to eight months? Three to eight months. -- that that puts them at Ο. 16 17 higher risk of bacteremia? I don't know the literature that deals with an Α. 18 19 age range of three to eight months. I'm just unfamiliar 20 with that. But greater than three months you're familiar 21 Ο. with? 22 Well, I'm familiar with the topic in general at 23 Α. all ages, and it is my personal conclusion that at all 24 25 ages the height of the fever probably increases the risk

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of bacteremia once you're above a month of age, but with less reliability than I thought at a former time due to the publications of articles which do contradict what I've just told you.

Let me ask you this. In 1992, the general Ο. 5 thought was greater than one month or greater than three б 7 months with temperature increased the risk of bacteremia? I think in 1992 most people thought that as a Α. а general proposition the higher the fever, the greater the 9 risk. And understanding that even then it was known that 10 there were certain breakpoints. For example, a 11 12temperature of 39 degrees -- greater than 39 degrees seemed to have a distinctly different risk than 13 temperatures less than 39 degrees. That was based on an 14 15 old study, but a uniquely useful study, published in 1975 from Boston Children's Hospital -- excuse me -- from 16 Boston City Hospital. Of 600 consecutive patients seen 17 with fever, none of them had a bacteremia if their 18 temperature was less than 39 degrees. 19

And I think people felt somewhere around 104 to 105 there was another breakpoint. Above that break level, the risk went up from the three- to four-percent level to a higher level of maybe five, six, or seven percent. But, of course, no one knew exactly where the breakpoints were at 1992 or even today.

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Is cyanosis one of the clinical pictures of 1 Ο. toxicity in febrile infants? 2 Well, it's one of the items that people look for Α. 3 in assessing the general state of an infant, but you'd be 4 looking for central cyanosis, not peripheral cyanosis. 5 Would you agree that the standard of care in Ο. 6 1992 in all children from zero to 36 months with fever 7 without known source and with some toxic appearance 8 requires hospitalization with antibiotic treatment and 9 sepsis workup? 10 MR. MULLER: Objection to the form of the 11 question. Fails to define "toxic." 12 No, I wouldn't agree with that as a straight Α. 13 statement. 14 What do you consider to be toxic 15 Ο. Okav. appearance of children with bacteremia? 16 That's one of those words that's very hard to 17 Α. define, "toxic," but the one -- the definition that I've 18 always used, which I think is the best, comes from a 19 pediatrician named Sidney Gellis, who was the chairman of 20 21 the department at Tufts for many years, and is now in his When he was asked to define toxic, he said, 2.2 eighties. "It's the child who looks and acts damn sick." It 23 conveys, I think, what the issue is; that is, it's a 24 child who is distinctly unwell, more unwell than just the 25

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height of the fever can account for, more unwell than the 1 2 time of day can account for, more unwell even than a possible focus of infection can account for. And for 3 people who do this day in and day out and see any numbers 4 5 of children every day, that conveys a message. Do you agree with me that with the type of Ο. 6 7 disease that Ryan was known to have, that symptomatology can change acutely? It can be a rapid-changing 8 condition? 9 A lot depends on the use of the word "rapid." Α. 10 It is a dynamic illness, and children can become ill 11 12 progressively, but different individuals will become ill at different rates, 13 Ο. Okay. Well, you've read the deposition of 14 Mrs. MacAdams in this case, have you not? 15 I have. 16 Α. And you read where she testified that a rash was 17 Ο. present after Dr. Perez examined Ryan, but when 18 Dr. Poleynard came back in, it came on. Would that be 19 consistent with this type of disease that this child had, 20to have a rash that suddenly appeared? 2 1 MR. MULLER: Is that the only thing that you're 22 23 asking? MR. BERGEN: Yeah. 24 25 MR. MULLER: Is the sudden appearance of a

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2	MR. BERGEN: Right.
3	MR. MULLER: consistent?
4	A. Well, I guess as a general truism, any child
5	that has a rash has to have a moment when that rash
6	appears. In that case it's instantaneous. It's no
7	different than any other illness, such as measles or drug
8	reaction or German measles. There's a point where you
9	don't have a rash; there's a point where you do have a
10	rash. I don't think it's any more abrupt here than it
11	would be in any of those other illnesses.
12	Q. When Ryan left the emergency room, more likely
13	than not you'd agree that the meningococcemia had not
14	been diagnosed? He had an undiagnosed case of
15	meningococcemia, more likely than not, when he left the
16	emergency room the first time?
17	A. Let me try to answer. When he left the
18	emergency room at 8:15 that night, I believe he had
19	meningococcemia, but I do not believe he had
20	meningococcal bacteremia, which is a word that says
21	you're not only bacteremic, but you are also severely
22	clinically ill.
23	Q. Okay. He still would have had the presence of
24	the organism of Neisseria meningitidis?
25	A. Meningitidis.

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Thank you. That organism would have been Ο. 1 present in his blood? 2 I believe more likely than not that he had it in 3 Α. his bloodstream. 4 Do you have an opinion as to whether or not, Ο. 5 6 based upon what you have reviewed, Ryan should have been admitted to the hospital during the first emergency room 7 visit? 8 Α. I have an opinion. 9 What is your opinion? 10 Ο. I don't believe there was an indication for 11 Α. obligatory hospitalization, no. 12 Do you have an opinion as to whether or not 13 Ο. antibiotics should have been used during the first ER 14admission? 15 Α. I have an opinion. 16 And what's your opinion? Ο. 17 It's my opinion that the use of antibiotics was 18 Α. 19 an acceptable but not an obligatory management option. 20 Q. Okay. Now, if you suspect a child to have -make sure I'm correct in this -- meningococcal 21 22 bacteremia, what do you consider the standard of care to be to treat that condition if the patient presents -- I'm 23 24 talking about 1992 because that's what this case deals 25 with.

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MR. MULLER: Object to the form of the question. 1 It's a hypothetical that fails to supply sufficient facts 2 for the doctor to form an opinion. 3 You can go ahead and answer. With an infant of Ο. 4 5 Ryan's age. Α. I can't answer it because I don't know Sure. 6 what you mean by the word "suspect." It means different 7 things to different people. а Ο. How about differential? In your differential 9 diagnosis? 10 The reason I have trouble with that is when Α. 11 12 you're dealing with small children, you also suspect bad things, If you find no evidence for them being present, 13 14 then you don't act upon them being present and, therefore, no treatment would be necessary for a 15 suspicion. Only if someone had clinical disease would 16 17 you begin therapy. 0. If clinical disease -- what do you mean by that? 18 "Clinical disease"? 19 20 Α. Well, a child who had clinical septicemia. You know, you can't look at the child and say when a child --21 excuse me -- what organism may be causing the septicemia 22 Q. 23 I understand that. But if you saw a child who had clinical 24 Α. 25 septicemia or clinical meningitis, then there are

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standard ways of treating those individuals. But
2 suspicion is not a word which has a common definition,
3 I'm afraid.

Q. Okay. Let's take the example where you believe this child does have meningococcal bacteremia. What do you do?

7 A. Well, if you believe the child has clinical
8 septicemia, because you don't know what might be causing
9 it until you get the results of your testing back -10 Q. Right.

A. -- then the proper approach to that is first and foremost to get the child to a place where they can deal with serious illness in small children, You then establish an intravenous line. You assess for the presence or absence of respiratory failure and shock, and if present, you deal with those first.

You obtain all the specimens that you will need to diagnose the disease, which is your clinical diagnosis. That would include under most circumstances a blood culture, probably a urine culture, and if safe to do so, a spinal tap, although it may not be safe to do so at the particular time.

Q. Okay.

A. And then you would initiate antimicrobialtherapy. And there are a number of choices there.

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Okay. Did you try to make a determination in 2 your mind as to whether or not a rash was present with 3 Dr. Poleynard?

A. I tried to make a determination based upon the available information which I possessed.

Q. Did you reach a conclusion, or are you not sureone way or the other?

I think that the available information to me 8 Α. suggests that no rash was present despite the fact that 9 the mother, in fact, does remember a rash being there. 10 Ι couldn't find confirmation of that reliably in the 11 contemporaneous notes written at the time that the child 12 became terribly ill and was admitted to the hospital and 13 14 so on.

Q. If a rash was present with Dr. Poleynard based on the patient's findings in the chart, would that change what Dr. Poleynard should have done in this case?

18 MR. MULLER: Object to the form of the question.
19 It fails to outline what you mean by "rash."

Q. You can go ahead and answer.

A. Well, if a rash is present in any child, the child requires a reevaluation, and that's the only thing that would have been required. And then based on the reevaluation, then there might be different management decisions made.

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Q. If it was determined to be a petechial rash, a
 nonblanching petechial rash, what would the standard of
 care have required in this case?

The standard of care would have required a Α. 4 reevaluation and a reexamination. One of the articles 5 that I supplied to you is an article from Boston 6 Children's Hospital looking at fever and petechial rashes 7 and how rare it actually is that that is a harbinger of a a serious illness. Consequently, a petechial rash is not 9 an uncommon finding in children with fevers due to many 10 different causes, but it does demand a reevaluation' 11

Okay. And reevaluation, what does that include? 12 Q. Well, it includes another systematic 13 Α. examination. It may include repeat blood testing. The 14 15 child did have some blood testing done earlier on, but 16 that was at 7:15. It's now an hour or so later. That 17 would be an option. Blood culture could be obtained, further period of observation could be obtained, the 18 child could have been given presumptive antimicrobials 19 and then sent home. 20

The child could have been admitted to the hospital for observation only, The child could have been admitted to the hospital for observation and antimicrobials. There are a number of different choices that one can make, and, of course, the goal is to try to

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1 match the level of risk that you perceive in the 2 individual patient. This type of disease that Ryan had also can Q. 3 demonstrate macular rashes, can it not? 4 It can. 5 Α. How do you describe a macular rash? 6 Ο. The definition of a macular rash is a flat 7 Α. blanching red rash. 8 And you would expect a physician practicing in 9 Ο. an emergency room in 1992 in the United States of America 10 to be able to differentiate between petechial and macular 11 rashes, would you not? 12 13 Α. Yes. An emergency room physician practicing in the 14 Q. United States in 1992 should recognize -- standard of 15 care would require that physician to be able to recognize 16 that macular rashes are associated with meningococcal 17 bacteremias? 18 That's an interesting question. Α. The main 19 article that showed that was published in an English 20 21 journal in and around that time. It may even have been after 1992. And I would not necessarily expect people to 22 23 associate macular rashes with meningococcal bacteremia in 24 1992, no. 25 Ο. But ysu would?

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A. That's my area of expertise.

Q. I understand. But there is a correlation between the two -- or can be -- a macular rash with meningococcal bacteremia?

A. Let me put it this way. The macular rash has
been reported as being a rash seen at some stage in some patients in meningococcal bacteremia. But in the
universe of rashes, it is an extremely rare component,
and, therefore, a macular rash in an otherwise
well-looking child would not raise the spectre of
meningococcal disease in a patient.

12 Q. Can the macular rash also turn into the 13 purpura-type rash --

14 A. Yes, it can.

15 Q. -- at later stages?

A. It can.

16

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17 Q. In the macular rash, a forerunner is the purpura 18 rash?

19 A. Not necessarily.

20 Q. Frequently, more often than not, a forerunner is21 the purpura rash?

A. Again, the macular rash is seen at times, but
not commonly, in meningococcal disease. And in this
particular case study that was done, many of those
children did evolve their illness into either petechiae,

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purpura, or both. But, of course, that is an evolution 1 of illness that is seen in children who have 2 meningococcal disease without a macular rash. 3 So it's unclear whether the rash is a precursor 4 or whether it is just another manifestation of the 5 disease. But as I say, it's an uncommon manifestation of 6 the meningococcal disease, and I would not want to raise 7 the spectre of meningococcal disease in a child who did 8 not look significantly ill. 9 You and I can agree that a child such as Ryan 10 Ο. with -- a febrile infant such as Ryan of unknown source 11 12who develops a rash in the emergency room -- that that 13 would at least -- standard of care -- require reevaluation of the patient? 14 15 MR. MULLER: You're talking about a child getting a petechiae rash? 16 17 MR. BERGEN: Just a red rash. 18 A diaper rash? MR. MULLER: 19 MR. BERGEN: I'm just asking about any rash on 20 his body. He's already told me rashes can develop 21 anywhere on the body. Does **a** rash raise a red flag in conjunction with 22 Q. 23 treating a febrile infant of Ryan's age, three-and-a-half 24 months? 25 Α. As a general answer to a general question, SANTA FE OFFICE MAIN OFFICE

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rashes can be clues to illness, and the rash certainly 1 2 should be looked at if it evolves in any setting, whether it's in an emergency room or in a clinic or whether the 3 child comes in with a rash. It's another clue to illness 4 that should be evaluated, but it doesn't necessarily 5 demand a full reexamination of every body part. 6 7 Q. Do you agree with me that Dr. Poleynard should have gotten other temperatures on Ryan? 8 Α. No. 9 So the two temperatures that you see in 10 Ο. Okay. 11 this case are adequate in your opinion? Oh, sure. We would only get one temperature. 12 Α. Even after treatment? 13 Ο. 14 Α. Sure. So you would use just the temperature at 15 Ο. admission, at triage, in evaluating this patient? 16 That's correct. We seldom get a second 17 Α. temperature on a child. 18 Ο. Did you see where Dr. Poleynard testified that 19 he gave another temperature but it wasn't recorded? 20 Yes, I saw that. 21 Α. 22 If you take a temperature, don't you record it? Q. Well, I don't personally take temperatures of 23 Α. 24 anybody. 25 Q. Well, if you order it.

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Again, I don't usually record it. It's usually 1 Α. a nursing function, if it's recorded at all. A lot 2 depends, sir, on what the object is for taking the second 3 temperature. If it's to convince myself that I've waited 4 long enough for the child to get the fever down so the 5 parents wouldn't worry, that would be one thing; if I'm 6 using it as a diagnostic maneuver, that might be 7 something else. 8

9 As I said, we so seldom order or take second 10 temperatures that it's very hard to answer your question. 11 Q. Let me ask you this. As to Ryan, from your 12 review of the first emergency room visit, I understand 13 you've expressed opinions that you think Dr. Poleynard 14 adhered to the standard of care during the first 15 admission.

A. Yes, I do.

16

19

22

Q. If you look at Ryan's material in the chart, he was over three months of age at the time, correct?

A, Correct.

20 Q. He had a fever, by your own definition, at all 21 times?

A. Definitely.

Q. And if you take the additional fact of a rash
being present during the examination by Dr. Poleynard, if
you take those triad of information, would you agree with

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question. 3 MR. BERGEN: Why don't we just reserve all 4 objections to all hypotheticals? I know you're going to 5 object to all hypotheticals I give him. 6 MR. MULLER: Okay. I'm reserving all objections 7 to hypotheticals, including form of the question, which 8 would include, but not be limited to, that it gives facts 9 that aren't proven, that it doesn't give sufficient 10 facts, et cetera. 11 MR. BERGEN: Right, Anything as to not assuming 12 all facts in evidence will be reserved. Let's just 13 leave --14 MR. MULLER: I may from time to time pipe up 15 anyway, such as this one where I have a problem with you 16 talking about a rash but not talking about what type of 17 18 rash. 19 Q. The rash that was described by Mrs. MacAdams in her definition. 20 Sir --21 Α. 22 Do you want me to rephrase it? Ο. 23 No. Any child with a fever has a possibility of Α. 24 having a bacteremia, sight unseen, but the level of risk of bacteremia changes as you get more information

me that Ryan had in his differential a bacteremia?

I'm going to have to object to that

MR. MULLER:

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regarding the child. Physical information, observation,
 laboratory testing.

3 So the answer to your question is generically 4 yes to any young child with a fever without an obvious 5 source, but the level of risk of that thought of 6 bacteremia or probability of bacteremia changes as you 7 gather more information during the period of evaluation.

8 And, of course, that's what your business is, is 9 to try to use physical findings, laboratory studies, and 10 the natural history to correctly assess what the risk 11 actually is.

12 Q. With a bacteremia, do you agree that one's white 13 cells can be consumed?

A. There are some children who have septicemia -meaning clinical illness, so they look and act damn
sick -- who have a bacteremia who have low total white
counts, yes.

18 Q. Well, isn't it with meningococcal bacteremia 19 that if it's fulminant you can have one's white cells 20 consumed?

A. Well, in any serious infection, white blood
cells are consumed, but the body makes up enough new ones
so that you have a steady state in the bloodstream,
There are situations of fulminant disease in which the
destruction of the white cells exceeds the capacity to

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replace them and the number measured in the bloodstream 1 2 will, in fact, go down, as it did here. Ο. In that instance, you get the shift to the right 3 in the band count, do you not? 4 5 Α. Well, no. Q. I'm sorry. Shift to the left. I'm sorry. 6 Increased neutrophils? 7 Excuse me. Well, under those circumstances you would have 8 Α. very few neutrophils of any sort because they are all 9 10 being destroyed. Okay. Let me ask you this. Do you get a shift 11 Q. to the left with this type of bacteremial infection that 12 Ryan had? 13 Well, you can have the presence of immature 14 Α. 15 forms with all kinds of infections, trivial and otherwise. Band counts in this age group are not useful. 16 You saw where he did have a shift to the left? 17 0. 18 Α. Well, he did have 29 percent band forms out of a 19 total of 9,600 total white cells, yes. But as I say, the 20 differential in this age group is not useful. 21 Well, I understand what you're telling me, but 0. 22 let me just ask you this question. If you have an 23 infection or bacteremial infection such as Ryan had, do 24 you get an increase in any neutrophils because they're 25 fighting off the infection -- or attempting to fight off

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1 the infection?

A. Well, with any infection, you can get an *increase in neutrophils.* It's not isolated to
meningococcal infections.

Q Does Ryan have a shift in his right?
A. No. If you look at his differential count, the
common interpretation of that would be a shift to the *8* left.

9 Q. As to viral infections, don't you more likely 10 than not expect the shift to be to the right rather than 11 the left?

A. Again, in this age group, the differential count
does not help you with regards to viral or bacterial
infections.

Q. Well, if you do have a viral infection, do you normally see a shift to the right as opposed to the left? A. No, not necessarily. That's why it's not a useful tool. And, in fact, if you look at the Baraff article, they specifically tell you not to use the differential counts because it is not a useful tool,

(The deposition recessed at 4:00 p.m. and resumed at 4:05 p.m. as follows:)

Q. In looking at Mrs. MacAdams' deposition, the portions you've highlighted, there's a series of questions in here regarding the rash that Mrs. MacAdams

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1 testified about. What was the significance of that to 2 you? Why did you highlight it?

3 Α. Well, as you've already pinpointed through your own questioning, I was concerned about the fact of the 4 rash and whether it were present, and if so, what type of 5 6 rash it was, and so on, and she was, of course, one of the main people who is involved in the memories of those 7 things, so I highlighted information regarding the rash 8 from her deposition. With Dr. Poleynard, it was easy 9 because he doesn't recall seeing the rash. 10

11 And then I highlighted references to the rash in 12 the medical records in order to formulate in my own mind 13 whether I thought a rash were present, and if so, what 14 kind it was.

Q. Okay. And if the rash was present as described by Mrs. MacAdams, how would you categorize that type of rash?

A. Well, she says that she has a memory of not only the rash being there on the chest, but of it being a small pinpoint rash. And when Dr. Poleynard looked at the rash and pressed on the rash, the color did not press out of the rash; therefore, it was a nonblanching rash. So if you only look at her memory alone, it would be classified as a petechial rash on the chest.

25

Q. Let me see if there's anything else while on

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this particular subject.

vou also had some squiggly lines by the question regarding had his muscle tone changed, and according to 4 Mrs. MacAdams, he was more listless and more subdued. 5 Did that have any significance to you and your opinions 6 in this case?

Well, the information was relevant to me because Α. 5 it seemed to me that Dr. Poleynard was obliged to be sure а that the child he was sending home was not a sick child. 9 He was not the admitting emergency room physician, but he 10 was the discharging emergency room physician. 11 He did say that he went in and examined the child, reexamined the 12 child, and the other reasons that he came in to see the 13 child 14

What I wanted to do was to gather information as 15 to what the child was likely to have looked like at the 16 17 time that the decision was made to finally send him home, and hers is, of course,, one of the important memories. 18 If you assume her recollection to be correct, 19 Ο. would you agree with me that Dr. Poleynard violated the 20 standard of care generally employed in the medical 21 22 profession in discharging Ryan without further workup in reevaluation and holding him there? 23

A. No, not necessarily. The words that she was
using -- and of course, so much depends on words here --

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1 were a mixture of observations that s ested to me that 2 the child was not as active as normally he is, and was 3 quieter than normal, But based on the words that she 4 spoke, at least in her deposition, and only looking at 5 those alone, I did not get the impression that this was a toxic or damn sick child.

Just to finish my sentence, under those 7 circumstances, it would have been acceptable for 8 Dr. Poleynard to send home a child who was not looking 9 toxic, given the fact of the other reassuring aspects of 10 the child's evaluation in the emergency department, 11 including the long period of observation, the reassuring 12 total white count, and the fact that the parents seemed 13 capable of reporting back any changes. 14

Q. Well, that white blood count -- you called it reassuring white blood count -- could that white blood count be one that was falling --

18 A. Well --

19 Q. -- based on knowing what his disease process was
20 later on?

A. Well, again, 1 have to look from the standard of care perspective. I have to look at it from that perspective. I can't hold Dr. Poleynard responsible for not seeing the future, If I knew 1 was going to have a heart attack next week, I would check into the hospital

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All you have is what you have at the time. Consequently, he sees a child who has a total count of 9,600. Now, that's a reassuring white count. It's the kind of white count that decreases the perceived risk of a serious illness. Not down to zero, but down to a lower level than when you first started off.

Plus, you have a child who's been there for three hours and has three hours' worth of intermittent observations. And if, in his opinion, the child was not becoming sicker while in the hospital, those are all reassuring elements that lead to an acceptable decision to send the child home.

14 Q. If you evaluate a child, do you make 15 documentation of that evaluation?

A. I don't understand the question, because thereare many points of evaluation of a child.

Q. Well, if Dr. Poleynard testified that he zvaluated this child and did an examination of the child in his deposition -- Do you remember reading that?
A. Yes.

Q. And if he did that, do you agree with me that the standard of care requires documentation of those indings about the examining physician?

A. No, I've never thought that the level of

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documentation and the standard of care were the same 1 2 thing. I believe they're two entirely different things. Q. Well, do you document if you would have examined 3 4 this child? Α. I think it depends on the clinical setting, but 5 if the child is really unchanged, there would be no 6 reason to document based on -- For the child's sake, 7 there would be no reason to document a child who has 8 exactly the same constellation of physical findings at a 9 10 later stage that they did at an earlier stage. Ο. Well, that's your determination that there was 11 no change in condition. 12 MR. MULLER: Which is what Poleynard has 13 testified to. 14 Ο. Well --15 My understanding of reading his deposition, 16 Α. 17 sir -- and all I have are the depositions --Right. Ο. 18 19 A' -- is that he claims that if there would have been a meaningful change, he would have documented it and 20 presumably even acted differently. And I must say, it is 21 the convention in medicine that things that are not 22 written down are taken to be normal or unchanged or 23 whatever the case may be. So the lack of any written 24 25 summary of the reexamination on the emergency room sheet

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1 is not unusual, in my experience.

2	ρ . You would agree with me that the onset of Ryan's
З	symptoms were sudden, according to the family, before
4	they arrived at the emergency room, on the triage note?
5	MR. MULLER: You mean the parents have said on
6	the triage note that they were sudden?
7	MR. BERGEN: No.
8	Q. According to the notes, it gives a complete
9	complaint and history of sudden onset of expiratory
10	grunting, tense, nausea, elevated temperature, began
11	approximately one hour ago, 4:30 p.m. You're aware of
12	that?
13	A. Yes.
14	Q. That would be a sudden onset by history, would
15	it not, of temperature? Expiratory grunting?
16	A. You used the word "sudden," If someone has got
17	a fever, it's got to begin sometime.
18	Q. It's not chronic?
19	A. At that moment is onset. To call that sudden is
20	really no different than any febrile child. Fever in a
21	child must begin at a certain point.
22	O. You've seen different studies that talk about
23	~ chronic fevers over two weeks of age, sudden onset, or
24	acute fevers?
25	A. Oh, I would call this an acute febrile illness.

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but the characterization of this as being a sudden 1 illness -- fever must begin at a certain point. 2 Ο. Okay. Do you practice emergency room medicine? 3 Let me answer that in two ways. The answer is Α. 4 yes in the sense that I see patients in the emergency 5 room, but only pediatric. I don't do adult emergency 6 room work. And I'm called in to consult on patients in 7 But -- if I could just finish -the emergency room. 8 this particular kind of illness that this child has is 9 bread-and-butter pediatrics that occurs in offices, 10 clinics, and emergency rooms, and it's not 11 quintessentially an emergency room issue the way an 12 automobile accident would be. 13 Ι 14 Ο. Well, I mean, do you see patients -- I say don't want to take this literally. Do you work emergency 15 room shifts? 16 No, I do not, but our emergency room is situated 17 Α. in a way that they can just call me when a patient 18 arrives if it's a patient that, for example, I've brought 19 in to be seen or that they want me to see. 20 Is that by consult? 21 Ο. Α. No. There are children that I will see 22 primarily in the emergency room, but also I will be asked 23 24 to come in to see children that have been already evaluated by an emergency room doctor. We do it all ways 25

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Q. What I'm trying to get at, when I think about an emergency room physician, I'm thinking of the front-line emergency room physician who takes all kinds. You don't work that kind --

A. No. I don't see adults. I see only pediatric patients.

Q. When you see a pediatric patient, if you see a pediatric patient in the emergency room, it's at the request of the emergency room physician?

11 A. Sometimes it is, but sometimes I will see the12 child as the primary physician.

13 Q. You say, "Meet me in the emergency room," and 14 you see the patient there?

A. Yeah, either I will, or the nurse who takes the telephone call will say, "Call to the emergency room and ask for the pediatrician," and I'm the poor old pediatrician who sees that child.

Q. Did the fact that Ryan had been irritable 15 minutes prior to arrival, by crying, have any significance to you in formulating your opinions as to whether or not further workup should have been done on Ryan during the first emergency room admission? A. All historical elements have importance, but

25 that fact in and of itself I don't believe would alter my

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opinion that the correct workup was, in fact, given to the child 2 Okay. If Mrs. MacAdams asked Dr. Poleynard to Ο. 3 look at the rash that she described on Ryan in her 4 deposition, would you have expected Dr. Poleynard to make 5 a note entry about that rash or not? 6 Α. Not necessarily. 7 Did the standard of care require for the 8 Ο. differential to come back and be reviewed before 9 discharge on Ryan? 10 You mean the differential white count? Α. 11 Q. Yes. 12 No, because as I've said, in the age group, the 13 Α. 14 differential count is not of importance in making a medical decision as to what to do with the child, as is 15 clearly stated in the Baraff article. 16 17 Then why would you order it? Q. I don't. 18 Α. In this case it was ordered. 19 Ο. It was. 20 Α. Do you see any reason why it should have been 21 Q. ordered in this case? 22 Well, I don't know what their normal routine is, 23 Α. sir, in this emergency department. There are many 24 25 emergency departments that I know of, including ours. In

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some of them, it goes together like peanut butt and jelly. You order a white count; you get a differential. Consequently, you don't have the option of ordering only one or the other.

In our clinic, however, it's different. We can 5 order either a general hemogram that only has the total 6 7 white count or a differential. But in the emergency department, it's done, 1 think, as a matter of course. а Т don't know whether this was a conscious decision that was 9 made by Dr. Perez to order the differential or whether 10 it's just what we do in our emergency department. 11

Q. Okay. And as to the temperature of this child, the 104.3, was enough to be done in this case to meet the standard of care Just one temperature on admission and that's it?

A. That's correct, because, you see, fever's only useful as a stop sign to tell you to look at the child, that this child has a febrile illness and needs an evaluation. Further measurement of the temperature is not useful from a diagnostic point of view.

And, in fact, I must say in all honesty, we have in pediatrics a condition that has been called fever phobia, where parents become quite apprehensive when they have fever alone, as if fever was going to harm the child. And we feel in our department that fixation on a

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temperature level for no good reason just reinforces that
 in the family.

3 Q. As to what's in the chart, I take it that you
4 considered everything in the chart on the first emergency
5 room admission to be accurate and correct?

6 MR. MULLER: I'm sorry, what was that question 7 again?

Q. You considered everything in the chart as being9 accurate, did you not?

10 A. I didn't see any contradictions that would have
11 made me suspect that it was inaccurate.

Q. Okay. Now, you've also been provided opinions 12 13 regarding Ryan's prognosis. Let's see. We'll mark this 14 as an exhibit. The interrogatory responses you've given us about your opinions is that you expect to testify to 15 the issue of standard of care and caution based upon 16 various facts which have been presented in the medical 17 records and depositions, and based upon test results, 18 that Dr. Poleynard acted within the standard of care in 19 his diagnosis, treatment, and discharge of Ryan MacAdams. 20

I think we've covered that aspect of your opinions, have we not, in its entirety, or is there anything else that you need to add that supports, in your mind, that Dr. Poleynard acted in the standard of care in discharging Ryan?

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A. Well, just by way of summary, I believe that Dr. Poleynard did in fact re-examine the child, did not find a child who had a meaningful deterioration from when he was seen by Dr. Perez, did not find a child who was toxic who needed a more comprehensive reevaluation or alternate management plan.

I do not believe that a rash was present, but if 7 it were present, I believe that Dr. Poleynard did, in a fact, look at it and made a judgment in his mind that 9 this did not alter the acceptability of sending him home. 10 And I believe that he provided good follow-up information 11 and instructions for the family. So under those 12 situations, I believe he met the standard of.care. 13 14 Ο. Okay. Even if a petechial rash was present on Ryan, he met the standard of care, is what you're telling 15 16 us, too?

A. If the child had a petechial rash but was a nontoxic child -- which I'm not admitting was the case. This is a hypothetical situation -- I believe under the circumstances that it was an acceptable treatment option to send the child home with close follow-up if the child were, in fact, nontoxic, as I state that I believe the child was.

Q. And when you say "nontoxic," you just said that you think the child doesn't look sick?

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76 Exactly, does not have a global appearance of a Α. 1 2 child who is seriously ill. 2 Q. Okay. Remember, Dr. Perez was guite eloquent in what 4 Α. he wrote down in making a description of the child. 5 Consolable, alert, tracking, appropriate response, 6 et cetera. If the child did, in fact, have those things 7 at the time that he was being discharged or did not have 8 a meaningful alteration from that state, that is a 9 10 nontoxic child, Ο. But if the child did have a toxic appearance 11 with petechiae rash, then the child should not have been 1213 discharged? 14 Α. I agree. 15 MR. MULLER: As defined by Dr. Radetsky on what a toxic-appearing child is. 16 17 MR. BERGEN: Right. Right, using my own concept of toxic. 18 Α. Which I still don't have a good feel for except 19 Ο. 20 it's just the child looks sick, is what you're telling 21 ne? 22 Α. Well, it's not just that, sir. It's one of 23 those terms which is inexact to people who don't see 24 children as a care provider day in and day out. But for someone who's in the business, when you say "toxic" or 25

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"looks and acts damn sick" or "child looks seriously ill, " you know what you're talking about.

There has been extensive investigation in trying to come up with other words for these things and also to 1 come up with a profile. One of the most famous of them was done in the early 1980s at Yale by Dr. Paul McCarthy, and it's called the Yale Observation Scale, and it looks at the child's color, sociability, state of hydration, ٤ consolability, breathing pattern, and the like in order C to define toxic, as well. 10

But I must tell you, despite the fact that it's 11 been used as a research tool, the Yale Observation Scale 12 has never come into general use, and what has remained is 13 this concept that in looking at a child who looks 14 inappropriately ill for a -- as a constellation of 15 features, of their alertness, their interactiveness, 16 their sociability, their color, their breathing, their 17 activity, their body tone, and the like, and you must 18 understand it's very age-specific, then that child is 19 nontoxic. 20

21 If a child, however, you know, is withdrawn, flaccid, mottled, breathing rapidly and heavily, has cool 22 23 extremities and so on, then you consider that child probably to be a toxic child. It means something to 24 people who actually do the work. 25

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1 Q. What about the fact that there is a history of 2 this child grunting? Doesn't that cause some kind of 3 concern for respiratory compromise?

A. The initial reporting of that historical fact
does, but it was not confirmed by the subsequent
examinations of the child in the emergency room
department. And I believe the mother has testified that
it was not present in the emergency department.

9 Q. But the history would be of some importance to 10 you?

A. It would be of importance, but unless it were validated by a finding at the time of examination, it probably could be discounted since serious things that cause grunting do not go away.

In reading about toxic appearances, I've come Ο. 15 16 across the word "listlessness." Is that one of the things that is equated to one of the clinical symptoms of 17 18 toxic? "Listless child," I think it's been referred to. 19 Yes, "listless" is another one of those words Α. for which is there is no good definition. "Lethargic" is 20 That's why you really have to get a better 21 another one. 22 description of what the child is or is not doing in order 23 to make an independent assessment, and that's why in reading the mother's deposition it's a little bit 24 difficult to know what she thought of her child, because 25

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she uses so many different words to describe the child including "somewhat subdued," which is -- again, it's only a word, but it conveys a different feeling than a child who is, perhaps, listless or a child who's lethargic.

6 Words mean what the people say that they mean, 7 to tell you the truth. And in looking at the chart at 8 the time, it is my judgment that to the examiners at the 9 time, the child did not look and act seriously ill, and I 10 cannot find a contradiction to that in the mother's 11 deposition.

12 Q. So even when she says that the child was13 lethargic, you rule that out?

MR, MULLER: Mom never said that,

15 Α. I didn't see that in her deposition. I saw "somewhat subdued"; I saw the word "listless." I'm 16 trying to think of some of the other words that I saw in 17 there. But I never saw "lethargic." So, again, I'll 18 have to confront that word. But that's another one of 19 those words that means different things to different 20 21 people. I have nurses who come in and say they feel really lethargic. 22

Q. What does lethargic mean for an infant?
A. I try not to use the word because it doesn't
have a common definition. That's why I will write down

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words that mean something to me A child that is tracking, a child who's consolable, a child who' alert, these things mean something to me.

Q. Do you know if he actually wrote those down or
those were complete sentences made by the Carswell
dictation system that was being used at the emergency
room?

I knew that they had a dictation system. Α. I was a 9 unfamiliar with the fact that it generates its own words, And if so, I'll need more information as to what you key 10 But I presume that -- Dr. Perez seems to support in. 11 that in his deposition, that these words accurately 12 described the child. He was accurately describing the 13 child he was seeing. 14

Q. What does "subdued" mean to you, if a child is subdued?

I think it means to me that the child is just a 17 Α. little bit less interactive than usual, less interested 18 in social overtures than usual, a little bit more quiet 19 than usual, a little less demonstrative than usual. 20 Mind you, we're getting on to 7:00 and 8:00 at night, and a 21 child with a fever being a little less than **all** of those 22 things does not seem to me to be out of the question for 23 a four-month old child. 24

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Q. Is there anything else that you base your

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opinion that this child -- that it was acceptable to discharge this child that you haven't --

A. I think we've talked about all those things.
And again, I think that the follow-up information
presented in written form during the time of the
reevaluation in the emergency department and the
hospitalizations supports my conclusion that, more likely
E than not, a rash was not present.

Q. What do you base that on?

10 Α. Well, there are a number of references in the subsequent chart. There, for example, is the reference 11 in the chart that Dr. Poleynard fills out at the second 12 admission that states the child developed a petechial 13 rash. So that would have occurred after the child had 14 Then there are a number of references by been sent home. 15 physicians and nurses in the Memorial Medical Center 16 17 notes that, with only one exception, all refer to the 18 rash developing after the time of the emergency department visit. 19

Q. And what exception are you referring to?
A. The only exception is a note that is written on
:his sheet called Nursing Progress Note.

Q. Right.

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A. And it's 5:30 a.m. to 6:20 a.m. -- and I can't read the handwriting, so I don't know who wrote it -- and

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it refers to the following statement by the mother. "Mom 1 states that baby was treated for viral illness and 2 released. Also, mom noted (quote) petechial rash to 3 abdomen which did not impress M.D.'s at St, Joseph's." 4 And there's no other quotation marks, so I don't know 5 whether that ends. That's the only deviation from what I 6 7 see as a consistent recording of the rash occurring after the emergency department visit in multiple places in the Я hand of various other individuals. 9

10 11 Q. So that's the only exception that you noted?A. That's correct.

Q. But, I mean, obviously, for some reason, you went through the medical records carefully and were noting the rash. You were concerned about the rash because you know that's a red flag in this type of treatment of these patients, right?

A. I needed to know the facts before expressing an
opinion, and I tried my best to get at them.

19 Q. If a rash was present, would that change your 20 opinions?

A. I think that any new physical finding that was not claimed to have been seen by the examining doctors would have to go into my opinions, so, of course, I was concerned about that, as well as any new physical finding that might have been there.

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But you knew the rash was an important finding Ο. 1 in this type of disease process? 2 Α. In retrospect, yes. 3 Q. Well, you would know that, if you were treating 4 a patient, a new rash has an important consequence in 5 this type of infection? 6 All rashes are important in that child with a 7 Α. fever because they may be a clue to the cause of the 8 fever. 9 Q . Now, the next point on the interrogatory answers 10 states, "On the matter of causation, Dr. Radetsky is 11 expected to testify that there was no causal connection 1213 between Ryan MacAdams not being diagnosed with meningococcal infection or treated with antibiotics 14 during his first ER (admission), and his development of 15 orthopedic and other problems. Treatment with 16 antibiotics during the first (admission) would not have 17 prevented such problems. Also, it is likely that the 18 orthopedic problems were unrelated to the meningococcal 19 infection." 20 I take it -- Let's do this, Let's take that in 21 22 subsections, if we can. 23 Sure. Α. 24 Ο. If Ryan had had a blood culture done during the

25 first admission, more likely than not it would have shown

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84 meningococcal infection -- Neisseria -- You have to --Α. Just call it the meningococcus. It's a lot easier. I believe more likely than not that that is ٤ true. And with that in hand, standard of care requires ο. f immediate antibiotic therapy? MR. MULLER: You mean when the blood culture Ε finally comes back? ŝ Well, if you suspected it beforehand. But you 10 0. could start antibiotics even before you got the result 11 back, right? 12MR, MULLER: Well, now you're skipping around. 13 you're talking --14 Peter, just calm down. I'm asking MR. BERGEN: 15 a good question. 16 You testified that if -- or you've purported to 17 Ο. testify that if he was treated with antibiotics at the 18 first ER admission, it wouldn't have made a difference 19 with any of the development of orthopedic or other 20 21 problems. That is correct. Α. 22 That's where I'm trying to get to. Tell me 23 Q. what's the basis of that opinion. 24 25 Α. Sure. If his bone injuries and his injury to

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bone growth were caused by the meningococcus, as I think it could have been caused -- and there are four articles in my packet that someday you'll get that have to do with this -- it is an extremely rare complication of meningococcemia.

And the reason I say it's extremely rare is that б in the best of those articles published from Baltimore by 7 the University of Maryland Group, they can only come up а with somewhere between 12 and 21 cases of this ever being 9 reported in the literature. They refer to 12, then they 10 mention an article in which 9 more cases could have been 11 due to meningococcemia, but they didn't do all the 12 cultures to find out one way or the other. 13

But at any rate, that's an extremely small 14 number of cases in a disease which is not that uncommon. 15 Thousands of cases a year occur of meningococcal 16 infection in the United States alone, much less the 17 That's an extremely rare complication, world. 18 Consequently, we're dealing with a serious illness that's 19 not uncommon, but not terribly common, and an extremely 20 21 rare complication of that illness. Consequently, there is absolutely no information regarding duration of 22 23 illness, timing of antibiotic therapy in meningococcal infection and the risk of full-blown disease. There's 24 25 just no information on that at all and no one possesses

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1 that information because it's so rare.

Consequently, you have to look at the relationship between the timing of therapy in the context of illness and outcome in other areas of meningococcal infection to try to come up with biological principles you can apply to this case.

Are you with me so far?

7 1 Q. Somewhat. Do you see anything in all the 9 medical records you've reviewed that would explain any 10 other cause, other than the meningococcal disease, to 11 Ryan's infarcts?

12Α. I'm not an orthopedic surgeon, and I don't pretend to be an orthopedic surgeon. I believe the 13 meningococcal infection could have caused those bony 14infarcts, but there are many other cases that I'm 15 unaware. And, of course, the treating doctors who are 16 dealing with his bones, they don't care what the original 17 cause might have been. They're just trying to deal with 18 the cause that he has these growth arrests on these 19 various long bones. 20

Q. Do you think the people treating his condition now are less qualified to express opinions as to the origin of the bone infarcts?

Less qualified than what?

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Q. Than you,

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I don't believe they're less qualified than me 2 no. Are they more qualified than you? 3 Ο. They might be. I don't really know any of the 4 Α. people who are treating him, so I can't really answer. 5 Q. But, I mean, you aren't ruling out that the bone 6 infarcts was caused by the disease that he had? 7 I'm not ruling it out, sir. I just think it's Α. 8 an awfully rare complication and, therefore, there may be 9 some other condition that causes this. I just don't 10 11 know. Do you have any other explanation as to what Ο. 12 caused these bone infarcts? 13 14 Α. I honestly don't, because I'm not an orthopedic surgeon, and it's not an area in which I express any 15 16 expertise. 0. And the articles -- and I haven't reviewed the 17 ones that you have here today -- do they associate the 18 bone infarcts with the **DIC**? 19 They do. 20 Α. Could one possible explanation be, as to why you 21 Q. have not seen as many case reports of bone infarcts, is 22 23 because, A, generally this disease progresses to 24 amputation from necrosis and gangrene; and, B, death? 25 Α. well, I think you raise an important point, In

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purpura fulminans, which this child had, the combination of shock, DIC, and purpura in meningococcal disease, there is a considerable death rate and there is a substantial risk of loss of some portion of the body, whether it's a digit or a limb or something of this sort.

6 But the answer is no, I don't believe that that 7 fact alone would deprive the world of medical literature 8 of a number of case reports of this delayed diagnosis of 9 bony infarcts or growth plate arrests in meningococcal 10 infection if it were a common occurrence, no. The 11 answer's no.

So the fact that the best of all articles can 12 only report a handful of cases implies that .it must be a 13 rare complication, because there are articles -- and I 14 supplied one of them to you -- about purpura fulminans, 15 just in -- just from one measly institution that's 16 recording or reporting scores of cases of purpura 17 fulminans in meningococcal infection. So clearly it must 18 be much more uncommon. So this is a rare condition. 19 And 20 I've never seen one in my lifetime, and I've seen hundreds of cases of meningococcal infection. 21

Q. Has antibiotic therapy been effective in thecases that you've treated?

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A. What do you mean, "effective"?

Q. I mean successful, retarded any long-term

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1 sequelae.

A. Well, in the cases that I've seen, I've seen a
3 spectrum of outcome.

Q. I'm talking about the cases you've been actuallyhands-on, not what you've consulted.

A. I understand. In the cases I've seen, I've seen
7 the gamut of outcomes, including death, limb
8 dismemberment, burn-like injuries in the skin because of
9 large amounts of skin loss, meningitis, organ
10 infarctions, or total survival without any problems
11 whatsoever. So I've really seen the gamut.

12 Q. Do you believe that timing of antibiotic therapy13 is important to retard meningococcal CB?

Α. I think that the timing of antibiotics does not 14 influence many of the more serious outcomes of 15 meningococcal infection; that the timing of 16 antibiotics -- Clearly, you're worse off without 17 antibiotics than you are with antibiotics. But in terms 18 of the sensitivity of outcome to the timing of 19 antibiotics, I believe that there is a general grace 20 21 period within which the timing of the antibiotics leads 22 to an equivalent outcome.

Q. Okay. And you've criticized studies that have promulgated -- you know, in your writings -- timing of antibiotic therapy having a better outcome. The earlier

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1 the treatment, the better the outcome.

A. What I have criticized is unsubstantiated
3 claims.

4

Q. Right, scientific proof.

In other words, it is an old saying that Α. Yes. 5 the earlier you treat, the better off you are. And in 6 general that is true, but what it ignores is the fact 7 that there are certain conditions in which antibiotic 8 therapy is helpless to prevent death or severe outcome. 9 And in general, there is a wide grace period within which 10timing leads to an equivalent outcome. 11

So what I complained about is the fact that people say this phrase or make this statement when the truth is more complex than that, and they say it in an unsubstantiated way.

Ο. Isn't it true that there's no scientific proof 16 due to the fact that you don't hold off antibiotic 17 therapies with patients like Ryan if you know that that's 18 what they have? You can't do a study on them because 19 then you're doing it to the detriment of the patient? 20That's correct, you can't do a prospective Α. 21 randomized trial in which one of the choices is not 22 giving antibiotics. 23

Q. Right, like giving one antibiotics and givinganother a placebo?

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A. That's correct. So what you have to do is get
 at information some other way.

Q. What? What other ways have you read or learned -- That's probably a bad word to use. Are you familiar with the increased LPS's in individuals with meningococcal disease or with high levels associated with worst prognosis?

8 A. Yes, it is true that the worst prognosis is 9 associated with increased levels of endotoxin, or LPS as 10 you've expressed it, in the bloodstream. That is a bad 11 prognostic sign in those studies that actually looked at 12 it. And there haven't been that many studies, but there 13 have been some.

Q. Have there been studies that look at early administration of antibiotics and that prove the level of LPS falls, and prove that antibiotic therapy does make them fall in meningococcal disease?

Actually, there have been two sets of studies. 18 Α. One set of studies showed that there was a spike in the 19 level of LPS or endotoxin in the serum right after they 20 gave the antibiotic. In other words, there was more 21 present. And then there are other studies that show that 22 after you get the antibiotics, you will, in fact, then 23 see at a certain point a decay in the amount of LPS that 24 you measure in the serum. 25

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Q. You've read studies where the conclusion was reached that because you give antibiotic therapy and the LPS falls, then, therefore, antibiotic therapy does help meningococcal disease?

5 A. I have seen that claim in the study that --6 Again, there have only been a few studies of these sorts, 7 anyway, that saw the decline in LPS levels. The trouble 8 with that is that the bad outcome in meningococcal 9 disease has more to do with the host than it does with 10 the level of the LPS.

Q. And that's kind of where you've written or you've promulgated that this is the host response, not the timing of the antibiotic therapy?

Α. That's correct. The timing of antibiotic 14 therapy doesn't alter the host response, so that if you 15 have a host that responds in a particular malignant way, 16 injuring its own body in the process of fighting the 17 infection, the antibiotics don't seem to help that kind 18 of person as much as they would help, for example, a 19 20 person who can kind of respond in a more modulated 21 manner, and under those circumstances long delays in withholding antibiotics could lead to a deleterious 22 23 outcome.

Q. You would agree with me that since the advent orinception of antibiotic therapy in treating

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meningococcemia, that the mortality rate has decreased
 from greater than 80 percent to less than 20 percent?
 A. That is correct.

Q. And talking about the host response concept that you espoused, that if you have less than 20 white blood cells in the CSF fluid, that is a prognosticator that the host -- there's a failure of the host neutrophils to mount an appropriate response?

The usual viewpoint on these matters is that if Α. 9 an individual with meningococcal infection has 10 meningitis -- meaning greater than 10 or 20 white 11 cells -- it's thought to be a good prognostic sign 12 because it means that their body has been able to control 13 the disease long enough for them to develop the 14meningitis. That has actually been looked at now a 15 second time around, and the most recent studies do not 16 17 show that the presence of meningitis puts the individual in a better prognosis category, so there now seems to be 1.8 19 some disagreement amongst investigators regarding this 20 issue.

Q. Okay. But as to the white blood cells in the CSF fluid, are you familiar with any literature or studies that give an account --

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A. Well --

Q. -- that are significant? Regardless of the

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2	A. If you have a small number of white blood cells
3	in the CSF, people oftentimes interpret that as not
4	having meningitis. So I would actually have to look at
5	the study to know how they made the diagnosis of
6	meningitis but with no white cells in the spinal fluid.
7	Q. But you have to have explain to me In
8	order to have white blood cells in the CSF, you have
9	meningitis?
10	A. That's correct.
11	Q. And the finding of meningitis in the CSF or not
12	now is thought not to be a good indicator or a bad
13	indicator of outcome?
14	A. That's correct. The most recent studies to look
15	at it do not find that the presence or absence of
16	meningitis can give you a prediction of how well or
17	poorly the person will do.
18	Q. Okay. Did you make a determination as to
19	whether Ryan had meningitis in his blood count
20	A. Yes.
2 1	Q or CSF?
22	A. Yes, I did make that determination.
23	Q. What was your determination?
24	A. He did have meningitis.
25	Q. And did you look at the white blood cells?

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A. Yes. He had a spinal fluid that showed 48 white
 blood cells.

And wouldn't that be an indication that the host Q. 3 neutrophils -- there was not a failure of the host 4 neutrophils to fight off or appropriately respond if 5 earlier intervention had been instituted in this case? 6 Well, I'm sorry, but the problem with this child Α. 7 was not a neutrophil problem. The problem with this 8 child was the blood vessel disease. And under those 9 circumstances, the level of neutrophils is a very poor 10 11 quide.

Q. I'm going back. If antibiotic therapy had beenintroduced initially, not later. The first admission.

14 A. Then I'm confused as to the role of neutrophils15 in your inquiry,

16 Q. Okay. So the fact that he had 48 white blood 17 cells in the CSF is not -- does it weigh in your opinion 18 one way or the other?

19 A. That's correct.

20 Q. And the fact that he had meningitis was 21 originally thought would be a good prognosticator of 22 earlier antibiotic therapy, better outcome?

A. Well, no, it's not a prognosticator that early antibiotic therapy should or shouldn't have been given in that sense. Just taken as a prognosticator, it has been

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1 thought in the past that the presence of meningitis was a 2 good prognosticator. More recent studies looking at it 3 could not confirm that.

Okay. Do you differ with the earlier studies, Ο. 4 or where do you line up on that school of thought? 5 Α. School of thought. It's been my experience that 6 meningococcal disease is not good no matter what you have 7 in the way of spinal fluid findings. a I've seen individuals develop malignant brain swelling with 9 meningococcal disease and die because of the meningitis, 10 11 so I'm not convinced that the presence or the absence of meningitis is of any clinical use at all. 12

Q. When you mentioned earlier about endotoxins, isthat the real culprit in this disease?

15 A. Which, sir?

Q. The real culprit to causing injury and death isthe release of endotoxins in the cell wall?

Α. Well, the real culprit in this disease is the 18 way that the child reacted to the presence of the 19 20 meningococcus. You see, this was a serotype B It's actually a less virulent form than 21 meningococcus. 22 other forms of meningococcal infections, particularly the serotype C, which is seen now commonly in the United 23 24 States, which has worse outcomes in general,

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The real problem here was this child had zero

blood vessels reactivity, a severe vasculitis. 1 Ιt certainly was started by the presence of the 2 meningococcus -- no question about that -- but took on a 3 life of its own that continued despite the fact that you 4 killed off the meningococcus. And therein lies the 5 difficulty of ascribing tremendous advantages to the 6 timing of antibiotics in a disease in which it is the 7 host blood vessel injury and excessive clotting of blood 8 vessels which led to the DIC, purpura fulminans, and 9 possibly the bone injuries in this child, which is not 10 11 subject to modification with antibiotics.

12 Q. Let's back up, if you don't mind, to help me 13 understand that. You're saying that he had the type B, 14 which is usually not as strong as other types of 15 meningococcemia?

Α. That's correct. The serogroup B meningococcus 16 is a less virile type organism than other serotypes. 17 18 Q. Now, what you're telling me, then, 1 take it is because he had -- he was not able to fight off DIC --19 20 Α. No, what I'm saying is that DIC and the purpura fulminans that this child has is not the result of a 21 poison excreted by the germ or a destruction of blood 22 23 vessels because of the infection. It is the result of 24 his reaction to the infection. And he reacted to the infection with inflammation in the small blood vessels as 25

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well as clotting within these damaged vessels, and it is 1 this kind of reaction that he had that led to all of the 2 subsequent problems that he encountered both with regards 3 4 to the initial management at Memorial Hospital -- which I think they did a fine job of -- his kidney failure, the 5 need for blood products, and attempts to slow down the б DIC, and then ultimately the skin disease, and perhaps 7 even the bone disease if, in fact, his bone disease was 8 due to the meningococcus. 9

Q. When do you think the window of opportunityclosed on this child to stop that type of reaction?

I honestly believe, sir, that once the infection Α. 12 gets established in a patient who's going to react that 13 way, that antibiotics don't alter the process. 14 In other words, to restate it, from the time that one could have 15 reasonably made the diagnosis and reasonably instituted 16 therapy, the die is cast if someone is going to react 17 this way. 18

Q. When do you think, then, infection wasestablished with Ryan?

A. I just have no idea, and no one really can know. That's an impossible task to know when the first germ arrives or, you know, when it first gets into the bloodstream. These are imponderables that no one can know.

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You agree that meningococcus is a very rapidly 2 dividing organism?

A. It divides as rapidly as any other organism,
4 sir.

Q. Does it divide every thirty minutes?
A. No one knows actually the division rate in the
human body. In a laboratory beaker of growth broth with
no defenses, it probably divides every 30 to 40 minutes
depending on the temperature and the nutrients
Meningococcus needs particular nutrients.

But, you see, the body is different. The body has white blood cells, it has a liver and a spleen, it has lymph nodes, it has bone marrow, and it has serum factors which inhibit the growth of bacteria, so that the doubling time in the body is different than the doubling time in a beaker in broth in a laboratory.

Q. Have you ever read about where they took -where the endotoxins double every 30 minutes, and if you
start one meningococcus at ground zero, in 30 minutes
it's one, at an hour it's two, at an hour and a half it's
four, and then on out?

A. Sure, but this is all done in laboratorybeakers, not in the human body.

Q. Well, you don't test human bodies in this type of disease because it would **be**, basically, watching them

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1 die before your eyes, wouldn't it, potentially?

Α. Yeah, you can't get that information from a real 2 human being. Actually, what is known in human beings is 3 that the numbers of organisms usually plateau at a 4 certain level and they do not keep rising until your body 5 becomes clogged up with the organism, so that there is a 6 steady state that is reached, but it's different in 7 different individuals. And this has been done in animal 8 experiments. To put it another way, you can't use 9 doubling times to backtrack to say, "Ah-hah, this is when 10 the disease begins." 11

Q.

12

Right.

A. The reason you can't is twofold. One is, you don't really know the doubling time in a particular human body, so you don't know what that number is. And the second is, the disease is not Just the organism, it's the response to the organism, and different people respond with different vigor.

19 So someone who responds more quickly, more 20 dramatically, and I might say more malignantly, will 21 actually come down with symptomatic clinical disease 22 earlier in the course of their illness than someone who 23 responds in a very subdued fashion, since the illness is 24 not the organism; the illness is the response.

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You see, we will treat children who have

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1	meningococcal bacteremia that's been untreated for weeks
2	and they just don't respond very violently; consequently,
3	they don't get very sick. They have what's called
4	chronic meningococcemia, which is, if they sampled their
5	blood over one or two or even three weeks, they would
6	remain blood-culture positive, but they wouldn't be
7	terribly ill, They would get rashes that would come and
8	go and fevers that would come and go, but they would not
9	be materially ill. They're not the kind of children that
10	react violently, the way that this fulminant kind does.
11	Q. Isn`t it the endotoxins that stimulate the
12	response from the body's immune system?
13	A. It's endotoxins and other parts of the cell wall
14	of the coccus that is the stimulus for these reactions.
15	Q. And with vigorous response from the immune
16	syst,em, such things could be DIC?
17	A. That's one of the ways the body can respond.
18	And, of course, it's not a very intelligent way to
19	respond, since you're harming yourself in response to
20	this organism.
21	And one of the articles that you'll have someday
22	is by Mike Levin, who is a researcher, the professor of
23	pediatrics at St. Mary's Hospital in London, whose area
24	of research is meningococcal disease, and it's ${f a}$ recent

25 study of the state of the art of dealing with these

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issues. And you can see the perplexity he feels in, one, 1 2 knowing what actually causes the reaction; two, why certain people react more strongly than others; and, 3 three, what to do about it when you've identified it, 4 because there's just a limited number of means at your 5 disposal to try to deal with it. They did everything 6 they could for this child, but you're limited in what you 7 can do. 8

9 Q. And from what you've told me, you would agree 10 that the body's response is directly related to the 11 avenue of endotoxins to which the body's exposed?

A. No.

12

13

15

Q. You would not?

14 A. No.

Q. Okay.

A. The body's response is in response to the
endotoxin, but the timing of the response and the vigor
of the response is peculiar to the individual.

Q. And in this case what you're telling me is that once the infection was established in Ryan, whenever that was, the die was cast?

22 A. I believe so.

Q. Okay. But you can't tell me when the infection was established with Ryan within a reasonable degree of medical certainty?

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A. Well, I believe he had the bacteremia at the

3	Q. Okay. Was the infection established then?
4	A. Well, I believe it was, because I can't find any
5	other explanation for his fever. Consequently, here's a
6	child who has a fever. It's got to be due to something.
7	We know that just a handful of hours later he has
8	terrible purpura fulminans that anybody could diagnosis.
9	So it's my opinion that when he was being seen
10	on the first emergency room visit, unknown to the
11	treating doctors, who don't have a crystal ball, he had
12	meningococcal bacteremia that was symptomatic, but he
13	hadn't had that violent reaction yet.
14	But even if antibiotics had been indicated at
15	that time I do not believe that they were indicated.
16	I do not see an indication for antibiotics, although some
17	people might have given antibiotics because different
18	people have different management styles. Even with that,
19	I don't believe that the process that had begun could
20	have been switched off by the antibiotics.
21	Q. At any time during the first admission?
22	A" That's correct.
23	Q. And do you agree with me that antibiotic therapy
24	can retard the occurrence of DIC in this type in a
25	patient with meningococcemia?

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Α. It's very, very hard to answer that question 1 2 because if you're looking at this very rapid onset disease, I don't believe antibiotics can alter the 3 incidence of DIC. But if you're looking at untreated 4 meningococcal disease before there are antibiotics -- And 5 nowadays with treated meningococcal disease, clearly 6 7 there are some people who over the course of many days would have developed DIC who aren't going to develop it 8 because they're getting their antibiotics. But that's a 9 different form than an explosive form, this purpura 10 fulminans form. This I don't believe is subject to 11 antibiotics altering the incidence of DIC, particularly 12 if you're dealing here with just a matter of hours. 13 14 Ο. When you say a matter of hours, what do you mean? 15 Well, I mean, he's discharged home at 8:15 that Α. 16 night, and he comes back in at 3:30 that morning with an 17 18 illness that's clear to everybody that he's terribly ill. You're only talking here about seven hours. 19 That's why I wanted to find out what's your Q. 20 understanding when you talk about "a couple." 21 I said a handful of hours. 22 Α. Seven hours, 15 minutes? Ο. 23 But it's not a bunch of days. He's not going 24 Α. two, three, four days without therapy, ill and 25

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languishing and deteriorating in front of your eyes.
 This is a child who only has a fever at this point.

On the second visit, he just looks sicker than all get-out. In retrospect, you can't explain his fever at all due to anything else but the meningococcal infection.

7 It does take time to develop the DIC. I mean, 8 it doesn't happen instantaneously. But even killing off 9 the bacteria, you know, you've got a spike in the 10 endotoxins when you kill the bacteria. That's going to 11 foment the process that's well in place by seven hours 12 later.

I just don't think there's any support to the notion that in this kind of illness that the timing of antibiotics makes a significant difference in outcome. And I think there's good literature support for that contention, and you'll have some graphs to even look at that.

19 Q. In that one case where you did testify on behalf 20 of the plaintiff, would antibiotic therapy have made a 21 difference in that particular case?

A. That was a real peculiar case because this was a child who was admitted to the hospital with meningitis caused by H flu, which used to cause most of the meningitis in most cases of the old days, who was put on

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the wrong antibiotic, and they keep him on the wrong 1 antibiotic, even though he's not getting any better, for 2 ten days. And then he's not better, so they do a spinal 3 tap, and lo and behold, the antibiotic wasn't working. 4 Then he switched the antibiotic, and he has a stroke. 5 That's clearly a different thing. Ten days go by on the 6 wrong antibiotic before you're wise to the fact that it 7 wasn't a wise choice. I think it's different than this 8 situation. 9

10 Q. Have you testified in cases where there was a
11 12-hour delay?

A. Well, as I've already said, I've never testified
in a meningococcal case in which I felt that the timing
of the antibiotics could have altered the outcome.

Q. And of all those cases, what time periods werewe talking about? From a range of what to what?

A. On which cases?

18 Q. In the cases that you've testified that it would 19 not have made a difference when antibiotic therapy --20 What I'm trying to find out is what was the delay in 21 treatment ranges in those cases?

A. Again, without recalling exactly the cases, my impression **is** they're all hours, of the same sort as this -- I mean, a handful of hours or a half day or something like that -- in which I felt that the infection

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1 was clearly already established, in retrospect. In 2 retrospect. And then by then the process which would 3 have led to the limb injury or the bad outcome had 4 already been started and the momentum was in place such 5 that antibiotics couldn't have altered it.

Q. In each of those cases, you're basing that on
7 the individual's response to the disease process?

Well, I'm basing it on two things. One is what 8 Α. I know about the biology of the disease, which is that 9 10 the vigor of the reaction is an individual issue, because, clearly, meningococcal infections give you a 11 whole spectrum of illnesses, including some case reports 12 of children who had meningococcal bacteremia who cured 13 themselves, came back in, and they were fine, and you 14 culture them, and it's gone, So it's really a whole 15 spectrum, with a very bad portion of the spectrum on one 16 side, the septic shock, purpura fulminans side of it. 17 So the biology of the disease is one part. 18

But the second part is a series of reports and case series that show that with the disease in general the really bad outcomes seem to be insensitive to the dosing of antibiotics. This means death or severe damage. And, in fact, it's quite clear that from the very beginning, in retrospect, they were kind of set up for these bad outcomes because their decease was very

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explosive. They were usually sick a very short period of time if petechiae or purpura were present. They were less than 12 hours. They were diagnosed usually very early on because they looked so ill so fast, but they just have that bad outcome.

And I've supplied you with some of that stuff in the materials that I had in my folder here.

a Q. Are you critical of Mrs. MacAdams of how she9 followed her child that night?

10 A. Oh, gosh, no. You know, she's doing the best11 she can as a parent, of course.

Q. When was the last time you treated a patient with this type of disease that Ryan had? What's your experience per year?

A. You mean this spectrum --

16 Q. Yes.

-- or just meningococcal infection in general? 17 Α. Meningococcal infection in general of an infant. 18 Ο. Well, just a couple of months ago we had child 19 Α. come into the emergency department who looked well. He 20was three or four months old. He had a blood culture 2 1 done, he was sent home. The next day the blood culture 22 was positive for meningococcus. You already know that 23 24 all the alarm bells went off all over the place.

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The child was rushed back in. He still had a

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1 fever. He didn't look particularly sick. We-gave him 2 antibiotics. He was fine. He was obviously a child who 3 didn't react from strictly the meningococcus, because 4 there was 12 or 14 hours between when he was seen and 5 when the blood culture was positive.

6 Q. In that particular case, why did you get a blood7 consult in the patient?

A. I didn't. It was done by the emergency room
folks. And I don't know what all went into the decision
to get the blood culture. But here, of course, even if a
blood culture had been taken, he got ill so fast that the
blood culture would not have been read until a number of
hours after he was seen the second time around.

Q. And in those cases where you suspect and do blood cultures, don't you treat with antibiotics after blood culture immediately?

Most people make those two separate 17 Α. No. decisions. In other words, the blood culture acts as 18 kind of a fail-safe. Well, I still want to be sure that 19 this child doesn't get out of my hands, so I'll do a 20 21 blood culture. I don't think he's very sick, so I won't 22 treat him, but I know I have a blood culture present as kind of a -- what's the right word? -- as kind of a 23 24 safety valve for my concerns because I worry about small concerns, I'm just going through an internal monologue 25

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MAIN OFFICE 500 Marquette NW, Suite 280 Albuquerque, NM 87102 (505) 843-9494 FAX (505) 843-9492 1.800.669.9492 1 in someone who might order a blood culture.

I do it as a safety net for the child. But I 2 really don't think they're ill, so I'm not going to start 3 Two separate decisions. If I do think you're therapy. 4 5 ill enough to start therapy, I will get a blood culture first. No question about it. But there are often times 6 you get a blood culture even if you don't plan on 7 treating it. 8 I must tell you, with small children who come in 9 with fevers, we often get a blood culture, because if you 10 get a good stick and get a CBC, you might as well get a 11 culture at the same time. Kids being so tough to get a 12 13 blood culture on. Isn't it true you promulgated the theory of 14 Ο. 15 using infants -- of using three different criteria: clinical judgment, total white blood count, and -- you'll 16 17 have to help me -- erythrocyte sedimentation --The SED, or SED rate as we say for short. 18 Α. 0. Yeah. 19 20 Α. I don't think I promulgated that. What I've done, in 1984 I reviewed all the information present up 21 to that time which concerned investigations in the 22 clinical judgment, total white count, and SED rates, and 23 24 I tried to show clinical judgment is superior to white 25 count and SED rates.

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MAIN OFFICE 500 Marquette NW, Suite 280 Albuquerque, NM 87102 (505) 843-9494 FAX (505) 843-9492 And in 1996, I did analysis another way, looking at risks, and I tried to show how you make a judgment sequentially. First you listen to the history, then you get the temperature, then you do your examination, and at each stage you're saying, "All right, now, what do I think?"

7 And then if you're still iffy, you might order a 8 blood count. If you're clear it's not a -- unclear it's 9 a sick child, you may not order a blood count. You're 10 going in stages with a maneuver that you take in order to 11 clarify what you should do next in sequence.

12 Q. Well, in the management of the febrile infant,13 do you get a SED rate?

14 A. No, 1 do not. It's takes an hour to get back.15 It's usually not very useful for us.

16 Q. You've written that that doesn't predict the 17 presence of bacteremia, but it rules it out.

A. Yes, a normal sedimentation rate decreases the
probability of a bacteremia to a very low level, but not
zero, obviously, and it can be reassuring if you're kind
of on the fence.

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Q. Did they obtain a SED rate in this case?

A. They did not.

Q. Are you critical of them not getting a SED rateduring the first admission?

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Α. I don't think so. As I said, the white count 1 was normal, the total white count was normal, and that, 2 3 from their point of view, was reassuring. And they had three hours to observe the child, and I think from 4 Dr. Poleynard's point of view the child did not look 5 seriously ill after that period of time and, therefore, I 6 don't criticize him for not getting a sedimentation rate, 7 because I don't feel it would have added anything to the 8 9 management. Would you agree with me that Ryan did sustain Q. 10 serious illness from meningococcal disease in this case? 11 12 Α. Yes. And how would you summarize his injuries as a Ο. 13 result of the meningococcal disease? What injuries did 14 he sustain? 15 Α. You mean long-term injuries or during? 16 Let's talk about short-term. Q. 17 During the acute course, he had a prolonged Α. 18 hospitalization with shock, depression of cardiac output, 19 20 DIC, significant purpura fulminans, kidney failure, need for artificial ventilation, and meningitis. 21 22 If you just look at his presentation, his prognosis for death was about 30 percent. So from my 23

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point of view, this was extremely good medical management

in the acute phase, because he didn't die. I mean -- And

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1 he had a pretty awful likelihood of dying. But he had 2 3 long period of time, as you well know.

4

O. Long-term?

A. Well, long-term is as best I can understand it, and I don't have each and every one of his follow-up records, but I do have some. I think that the thing that will affect him the most will be his limb discrepancies and his -- for want of a better word, the malformation of his lower limbs that has resulted from his many surgeries and his growth arrest.

12 And as I said, I think meningococcal infection 13 can cause those bony injuries. I'm not an orthopedic 14 surgeon, and I don't know whether there are a number of 15 other things that can cause it. The treatment I don't 16 think would be any different no matter what the actual 17 cause is.

Blessedly, he did not have any brain damage that I can see. His hearing seems to be fine. I note that he had an eye examination done, but I don't believe that they ascribe any eye injury to his meningococcal infection.

And then clearly, any child who has any disfiguring injury is going to have an emotional side to that injury, and anyone who takes care of children

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realizes that and knows that there's going to have to be 1 a good collaboration between the family and the treating 2 health team to try to get him over those emotional 3 hurdles. 4 Do you have any opinion as to how tall he'll Ο. 5 ever grow? 6 I don't have an opinion. I just don't know. 7 Α. Q. Let's take a short break. I think I may be 8 finished. I just want to check a couple of things, if 9 you don't mind. 10 (The deposition recessed at 5:20 p.m. and 11 12 resumed at 5:25 p.m. as follows:) (Exhibit 4 marked.) 13 If an infant develops DIC with this type of Ο. 14 disease that Ryan had, is it often your opinion that 15 because it goes to DIC, that earlier antibiotic 16 intervention would not have changed the course? 17 No, not necessarily. Looking at it in Α. 18 retrospect, because that's the only way I think you can 19 20 make judgments about matters of causation, he had explosive illness, short period of time of illness, 21 22 presents with purpura fulminans, develops septic shock, kidney failure, DIC. I mean, that kind of package. 23 24 Because of his age, it's known that children of this age group typically have worse blood vessel disease and blood 25

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MAIN OFFICE 500 Marquette NW. Suite 280 Albuquerque, NM 87102 (505) 843-9494 FAX (505) 843-9492 vessel clotting and damage to tissue or organs just because of his age. And the reason is that there are a couple of substances in the blood that prevent excessive clotting that are at low levels in four-month old children. Protein C and protein S. Consequently, children of this age in general do worse than adults would with regards to this.

But it's the explosive nature of this disease, a having it so quickly over a short period of time, that 9 10 makes me feel that by the time he developed a symptomatic illness -- which in retrospect I believe the fever and 11 the illness that brought him into the emergency 12 department was a symptomatic illness -- the portion of 13 his system that reacts to these infections was already 14 geared up and starting to go. It just hadn't guite 15 expressed itself yet. 16

So, even if you killed the germ with an 17 antibiotic, you wouldn't have stopped that. One, it was 18 already in place and beginning to go; and, secondly, the 19 endotoxin, which is one of the stimulus for it, would 20 still be around, because, in fact, the levels get higher 21 right after your treatment with antibiotics. 22 And he's 23 just in seven hours later with full-blown disease. I don't even see physiologically how the antibiotic could 24 have made any difference. 25

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MAIN OFFICE 500 Marquette NW, Suite 280 Albuquerque, NM 87102 (5051 843-9494 FAX (505) 843-0492 Q. So if I'm to understand the basis of your opinion -- and that's what I'm trying to do here today -is that based on the fact of his age and the short time period that made, in your opinion, it less likely to have a change of his course with antibiotic therapy being instituted during the first admission? Is that --

A. Not exactly. But if I could just restate it.
a First of all, I don't believe antibiotic therapy is
9 necessarily indicated on the first admission. I think
10 we've talk about that.

11

Q. We've talked about that.

Α. But there are two general bases for my opinions 12regarding causation. One is what's known about the 13 biology of how this disease expresses itself. I believe 14 he was at the far end of the spectrum on the fulminant 15 side as opposed to some other place on the spectrum, and 16 17 the biology of that kind of fulminant course is one in which the reactivity of the body happens abruptly and 18 seemingly without any ability of medical management to 19 20 alter.

21 Q. Okay. Can I stop you right there?

22

A. Okay.

Q. Was he at the fulminant stage when he first hitthe emergency room on May 7th, 1992?

25

A. He did not have symptomatic purpura fulminans or

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MAIN OFFICE 500 Marquette NW, Suite 280 Albuquerque, NM 87102 (505) 843-9494 FAX (505) 843-9492 1 septic shock or kidney failure, but he's the same child, 2 and in that sense, you have -- have to look at it, at any 3 rate, as one piece which has a clinical expression that 4 is truncated into a very short time interval. And even 5 though he wasn't expressing it at the time of his 6 emergency room visit, all of the elements were in place 7 which could not be reversed by antibiotics.

8 Q. Even if he -- I guess -- I take it, it's your 9 opinion, even if those symptoms were expressing 10 themselves during the first admission, even a 11 seven-hour-and-15-minute delay of antibiotic therapy 12 wouldn't have made a difference --

13

14

A. No, not --

Q. -- or treating it early?

No, not in terms of this, because of where he 15 Α. was on the spectrum of disease, this explosive reaction. 16 17 But the second basis for my opinion is a long series of articles that look at antibiotics in general in 18 19 meningococcal infection, and outcome in general, particularly bad outcome, like death. And as I read the 20 literature -- As I read the literature, I do not find any 21 2.2 correlation between outcome and the timing of antibiotics 23 or outcome and the duration of illness before antibiotics except to say that in the fulminant disease, the shorter 24 25 you're ill, that seems to be a bad prognostic factor for

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MAIN OFFICE 500 Marquette NW, Suite 280 Albuquerque, NM 87102 (505) 843-9494 FAX (505) 843-9492 1-800.660-0492 1 bad outcome.

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2	Q. There is also a school of thought that would
3	differ with what you just told me?
4	MR. MULLER: What part of that? Because he just
5	told you about three or four things.
6	\mathbb{Q}_{\cdot} Well, that earlier antibiotic therapy could make
7	a difference in a case similar to Ryan MacAdams.
8	A. I have had some not correspondence, but a
9	letter to the editor and responses from two sets ${f of}$
10	individuals, the group in Dallas who published the
11	article in a pediatric infectious disease article, and
12	the group in Barcelona, Spain, that there was a large
13	article in the <u>Journal</u> of the Medical Association.
14	Q. Right.
15	A. And the group in Dallas disagreed with my
-	
16	conclusions regarding death death and the timing of
16 17	conclusions regarding death death and the timing of antibiotics based on some studies that they referred to
16 17 18	conclusions regarding death death and the timing of antibiotics based on some studies that they referred to that took place in England.
16 17 18 19	conclusions regarding death death and the timing of antibiotics based on some studies that they referred to that took place in England. I believe they're interpreting these studies
16 17 18 19 20	<pre>conclusions regarding death death and the timing of antibiotics based on some studies that they referred to that took place in England. I believe they're interpreting these studies incorrectly. And there has now been an additional study</pre>
16 17 18 19 20 21	<pre>conclusions regarding death death and the timing of antibiotics based on some studies that they referred to that took place in England. I believe they're interpreting these studies incorrectly. And there has now been an additional study from the same English group which shows no difference in</pre>
16 17 18 19 20 21 22	<pre>conclusions regarding death death and the timing of antibiotics based on some studies that they referred to that took place in England. I believe they're interpreting these studies incorrectly. And there has now been an additional study from the same English group which shows no difference in death rate based on the timing of antibiotics that's</pre>
16 17 18 19 20 21 22 22 23	<pre>conclusions regarding death death and the timing of antibiotics based on some studies that they referred to that took place in England. I believe they're interpreting these studies incorrectly. And there has now been an additional study from the same English group which shows no difference in death rate based on the timing of antibiotics that's appeared since their letter appeared.</pre>
16 17 18 19 20 21 22 23 24	<pre>conclusions regarding death death and the timing of antibiotics based on some studies that they referred to that took place in England. I believe they're interpreting these studies incorrectly. And there has now been an additional study from the same English group which shows no difference in death rate based on the timing of antibiotics that's appeared since their letter appeared. So although they disagreed with me at the time</pre>

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letter to the editor, I think that my view is the correct
 one.

Q. I understand. Could we agree there's a debate?
A. Well, as I said, there's been a subsequent study
from England by the same group that did the studies that
they use as their response to my letter that I believe
undercuts the validity of their interpretation of the
earlier studies.

9 Q. But they hadn't changed their opinion?
10 A. Well, the subsequent study from England came out
11 after that interchange, so 1 don't know what they're
12 thinking these days.

With regards to the group in Barcelona, I 13 14 actually asked for clarification of their data which had to do with the use of oral antibiotics in an over 15 500-person series of meningococcal disease and outcome, 16 and their response to my letter clarified some of the 17 points of information that I didn't know about, and I 18 would think it's safe to say that they acknowledged my 19 point of view, but they expressed the hopeful belief that 20 antibiotics in some cases could alter the outcome. 21 Ιt was not an acrimonious debate, by any means. 22

Q. Okay. You said it wasn't an acrimonious debate?Friendly interchange?

A. Sure. Medicine is one of those things where

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MAIN OFFICI 500 Marquette NW, Suite 28 Albuquerque, NM 8710 (505) 843-949 FAX (505) 843-949 people do try to interchange and express viewpoints in
 order to clarify things. That's the beauty of the
 colleagueship that you have in medicine.

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£ .

Q. The one in Barcelona is the <u>Barquet</u>?

A. That's correct.

Q. But there is a debate among your colleagues as
to the prognosis with early antibiotic therapy with these
types of patients? I mean, that's been going on?

A. Well, you know --

10 Q. I know you have your opinion. I'm not debating 11 that with you. But there's a difference of opinions as 12 to whether antibiotic therapy makes a difference in this 13 type of treatment of infants, like Ryan's age, that has 14 existed through the years?

The only debate, so-called, that I've had have 15 Α. been these two letters to the editor that I wrote 16 following the publication of articles. I believe that I 17 have overwhelming support for my point of view, and I 18 think that anybody who has a different viewpoint must 19 20 mount the argument based on what's known and what has been shown in studies, and I think that that's incumbent 21 upon somebody who does have a different viewpoint. 22 Ι 23 think I make my case based on solid information, and I 24 think my interpretation's correct.

25

Q. As to the money that you're paid in reviewing

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cases like this, is that money that goes to you 1 personally, or do you pay it to the school or any 2 education or research or anything like that? 3 No, it comes to me personally, because it's done Α. 4 on my own time. 5 You don't have -- Some schools have contracts Q. б 7 where you have to submit the money to the school, and things like that, but I didn't know what the case was in 8 your situation. 9 Inasmuch as it's done on my own time, it's Α. No. 10 not anything that would concern the Lovelace Clinic. 11 MR. BERGEN: Okay. I believe that's all I have. 12 Thank you for your time. All right. 13 (A discussion was held off the record.) 14 You mentioned you knew of Dr. Meislin when you Q. 15 were at Arizona. 16 That's correct. Α. 17 Have you had any interaction with him since you 18 Ο. were at Arizona? 19 No. You asked me that, and the answer's no. 20 Α. Do you know that he lectures nationally on the 21 Ο. field of emergency medicine? Do you know anything about 22 what he does? 23 24 Α. Well, I know that he's an emergency medicine physician who has an academic career and lectures widely, 25

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1	sure.
2	Q. Okay. Do you know whether or not he's respected
3	in this field of emergency medicine?
4	A. I honestly don't know. That's not my field.
5	MR. BERGEN: Okay. That's all.
6	(The deposition concluded at 5:35 p.m.)
7	(Exhibits 2 and 3 marked.)
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PROFESSIONAL COURT

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	IN THE STATE COURT OF CHATHAM COUNTY		
	STATE OF GEORGIA		
2	RYAN MacADAMS, A MINOR, BY1AND THROUGH HIS NATURAL1GUARDIAN AND PARENT AS NEXT1FRIEND, DAWN MacADAMS,1		
Ľ	Plaintiffs, 1		
E	vs. 1 CIVIL CTION NO,		
٤	BLAKE CHRISTOPHER POLEYNARD,] [JURY DEMANDED] M.D., AND SAVANNAH SOUTHSIDE 1 CLINIC, P.C., 1		
с	Defendants.		
10			
11	CERTIFICATE OF COMPLETION OF DEPOSITION		
12	Ι ΠΑΝΝΑ SCHUTTE ΕVERETT CCR #139 ΠΟ ΗΕΡΕΒΥ		
13	CERTIFY that on January 25, 1999, the deposition of		
14	request of, and sealed original thereof retained by:		
15	MR. FREDERICK S. BERGEN Attorney for the Plaintiffs		
16	Marist Place, Lafayette Square 123 Charlton Street, East		
17	Savannah, Georgia 31401-4603		
18	I FURTHER CERTIFY that copies of this Certificate		
19	record and parties not represented by counsel:		
20	MR, PETER D. MULLER Attorney for the Defendants		
21	BOUHAN, WILLIAMS & LEVY Post Office Box 2139		
22	Savannah, Georgia 31498-1001		
23	•		
24			
25			
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	(505) 989-4949 FAX (505) 820-6349 FAX (505) 843-9494 FAX (505) 820-6349 FAX (505) 843-9492		

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I FURTHER CERTIFY that examination of this 1 transcript and signature of the witness were required by the witness and all parties present. On 2 ____ a letter was mailed or delivered to DR, MICHAEL S. RADETSKY regarding obtaining signature 3 of the witness. 4 I FURTHER CERTIFY that the recoverable cost of the original and one copy of the deposition, including 5 exhibits, to MR. FREDERICK S. BERGEN is \$____ 6 I FURTHER CERTIFY that I did administer the oath to the witness herein prior to the taking of this 7 deposition; that I did thereafter report in stenographic shorthand the questions and answers set forth herein, and 8 the foregoing is a true and correct transcript of the proceeding had upon the taking of this deposition to the 9 best of my ability. 10 I FURTHER CERTIFY that I am neither employed by nor related to nor contracted with (unless excepted by the 11 rules) any of the parties or attorneys in this case, and that I have no interest whatsoever in the final 12 disposition of this case in any court. 13 ´ -14 (huc DANNA SCHUTTE EVERETT, 15 CCR, RPR Certified Court Reporter #139 License Expires: 12/31/99 16 17 18 19 2.0 21 22 23 24 JOB NO. 5500-6 (DSE) 25 Proofread by: LR MAIN OFFICE SANTA FE OFFICE 500 Marquette NW, Suite 280 119 East Marcy, Suite 110 Albuquerque, NM 87102 Santa Fe, NM 87501 (505) 843-9494 (505) 989-4949

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1	MacAdams vs. Poleynard, et al.
2	DEPONENT SIGNATURE/CORRECTION PAGE
3	If there are any typographical errors to your
4	deposition, indicate them below.
5	PAGE LINE
6	Change to
7	Change to
8	Change to
9	Change to
10	Change to
11	Any other changes to your deposition are to be listed
12	below with a statement as to the reason for such
13	change.
14	PAGE LINE CORRECTION REASON FOR CHANGE
15	
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18	T NIGULARI (DADREGDY N.D. do horoby cortific that
19	I have read the foregoing pages of my testimony as
20	transcribed, and that the same is a true and correct transcript of the testimony given by me in this
2 1	deposition except for the changes made.
22	MICHAEL S. RADETSKY, M.D.
23	
24	JOB NO. 5500-6 (DSE)
25	Date Taken: January 25, 1999 Proofread by: LR
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IN THE STATE COURT OF CHATHAM COUNTY STATE OF GEORGIA

RYAN MACADAMS, A MINOR, BY	J
AND THROUGH HIS NATURAL	1
GUARDIAN AND PARENT AS NEXT	l
FRIEND, DAWN MACADAMS,	1
]
Plaintiffs,]
	3
vs.	1 CIVIL ACTION NO.
	1 I-97-3091-G
BLAKE CHRISTOPHER POLEYNARD,	[<i>JURY</i> DEMANDED]
M.D., AND SAVANNAH SOUTHSIDE	1
CLINIC, F.C.,	
	1
Defendants.	1

Defendants.

NOTICE TO TAKE DEPOSITION

TO: Michael S. Radetsky, M.D. Department of Pediatrics Lovelace Medical Center 5400 Gibson Boulevard, SE Albuquerque, New Mexico 87108

Please take notice that counsel for the Plaintiff in the above-styled case will take the Deposition of Michael S. Radetsky, M.D. at Bean & Associates Court Reporters located at 500 Marguette Northwest, Suite 280, Albuquerque, New Mexico, on the 25th day of January, 1999 beginning at 2:30 p.m. The Deposition will be taken before a Certified Court Reporter and Notary Public. The Deposition will be taken pursuant to Georgia Civil Practice Act (Code §§9-11-16 and 9-11-30) and for the purpose of discovery and preservation of testimony for trial. The oral examination will continue from day to day until its completion. You ${\tt may}$ attend and examine, as allowed by law.



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Michael S. Radetsky, M.D. is further ordered to bring the following:

- 1. Letters received from Peter D. Muller, or any member of the law firm of Bouhan, Williams & Levy, L.L.P., or any representative of Illinois National Insurance Company relating to your review of the materials relating to the case of Ryan MacAdams, a Minor, By and Through His Natural Guardian and Parent, Dawn MacAdams vs. Blake Christopher Poleynard, M.D. and Savannah Southside Clinic, P.C.
- 2. Medical Records which you have reviewed to formulate your opinions in the case of Ryan MacAdams, a Minor, By and Through His Natural Guardian and Parent, Dawn MacAdams vs. Blake Christopher Poleynard, M.D. and Savannah Southside Clinic, P.C.
- 3. Materials which you have reviewed to formulate your opinions in the case of Ryan MacAdams, a Minor, By and Through His Natural Guardian and Parent, Dawn MacAdams vs. Blake Christopher Poleynard, M.D. and Savannah Southside Clinic, P.C.
- 4. Medical literature which you have reviewed in formulating your opinions in the case of Ryan MacAdams, a Minor, By and Through His Natural Guardian and Parent, Dawn MacAdams vs. Blake Christopher Poleynard, M.D. and

Savannah Southside Clinic, P.C.

- 5. Reports, memorandums, written letters, or any other writing which expresses your opinions in connection with the case of Ryan MacAdams, a Minor, By and Through His Natural Guardian and Parent, Dawn MacAdams vs. Blake Christopher Poleynard, M.D. and Savannah Southside Clinic, P.C.
- 6. Records reflecting time spent in reviewing the materials in formulating your opinions in the case of Ryan MacAdams, a Minor, By and Through His Natural Guardian and Parent, Dawn MacAdams vs. Blake Christopher Poleynard, M.D. and Savannah Southside Clinic, P.C.
- 7. Invoices submitted for payment for your expert review and copies of payments received from Peter D. Muller, or any member of the law firm of Bouhan, Williams & Levy, L.L.P., or any representative of Illinois National Insurance Company relating to your review of the materials relating to the case of Ryan MacAdams, a Minor, By and Through His Natural Guardian and Parent, Dawn MacAdams vs. Blake Christopher Poleynard, M.D. and Savannah Southside Clinic, P.C.
- 8. Depositions which you have reviewed in the case of Ryan MacAdams, a Minor, By and Through His Natural Guardian and Parent, Dawn MacAdams vs. Blake Christopher

Poleynard, M.D. and Savannah Southside Clinic, P.C. This 20 May of January, 1999..

JOSEPH B. BERGEN Ga. Bar No. 054200 and FREDERICK S. BERGEN Ga. Bar No. Ø54160 By: m Frederick S. Bergen Ga. Bay No. 054160

123 East Charlton Street Savannah, Georgia **31401** (912) **233-6600**

1 16

CERTIFICATE OF SERVICE

This is to certify that I have this day served counsel for all parties with a true copy of the foregoing pleading by placing a copy of same in the United States Mail with adequate postage thereon to insure prompt delivery to:

> Peter D. Muller, Esquire BOUHAN, WILLIAMS & LEVY, L.L.P. Attorneys at Law Post Office Box 2139 Savannah, Georgia 31402-2139

This 20 day of January, 1999.

Bergen eder s.

123 East Charlton Street Savannah, Georgia 31401 (912) 233-6600

- Meningurocciemia 15 210 1/2/92 Seen ER St. Juseph Husp. Tal 1755 2N: Alert T = 1043Fing wet diapers skin "writ" 17:30 Tylenul sopp-MD: conscluble, alert, tincking, J 18:30 appropriate response to Noxious stimulus; notor tenre (n) Slein. @ turgon, worm, day, without egamosis, without petechiap No rash is 18:07 T-1013 R) present. A. Vie Syndreme. Dr Perez Pulse Ox- (L) big toe-98% Traires 19:00, DR LAB . 491 POLEYMAND (9(2))11.2 9600 33 (445, 293, 252 955 comps UA- Sp 9 R. 1.007. D.p O Sp02(RA) 98% Home instructions: Push Eluids Blenol Fly family MD, F. Not well a 2 or 39 call or return 2 Any problem 201.15 DIC 1930-20:45

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IN THE STATE COURT OF CHATHAM COUNTY

STATE OF GEORGIA

RYAN MACADAMS, A Minor, By)
and Through His Natural) _
Guardian and Parent as Next)
Friend, DAWN MACADAMS,)
)
Plaintiff,)
)
VS .) CIVIL ACTION NO. 19710910
)
BLAKE CHRISTOPHER POLEYNARD,)
M.D., FERNANDO JAVIER PEREZ,)
M.D., and SAVANNAH SOUTHSIDE)
CLINIC, P.C.,)
))
Defendants.)

DEFENDANT SAVANNAH SOUTHSIDE CLINIC, P.C.'S SUPPLEMENTAL RESPONSES TO PLAINTIFFS' FIRST INTERROGATORIES

COMES NOW Defendant SAVANNAH SOUTHSIDE CLINIC, P.C., and supplements

its responses to Plaintiffs' First Interrogatories as follows:

 (a) Michael S. Radetsky, MD CM Department of Pediatrics Loveiace Medical Center 5400 Gibson Boulevard SE Albuquerque, NM 87108 (505) 262-3542

Dr. Radetsky is expected to testify on the issue of standard of care and causation, based upon various facts which have been presented in medical records and depositions.

EXHIBIT 3	
BEASSOCIATES.	
10741	

Based upon the test results and findings of Dr Perez and Dr. Poleynard, Dr Poleynard acted within the standard of care in his diagnosis, treatment, and discharge of Ryan MacAdams.

On the matter of causation, Dr. Radetsky is expected to testify that there was no causal connection between Ryan MacAdams not being diagnosed with meningococcal infection or treated with antibiotics during his first ER visit, and his development of orthopedic and other problems. Treatment with antibiotics during the first visit would not have prevented such problems. Also, it is likely that the orthopedic problems were unrelated to the meningococcal infection.

DATED this <u>4</u> day of January, 1999.

BOUHAN, WILLIAMS & LEVY LLP

By:

PETER D. MULLER Georgia State Bar No. 528233 ATTORNEYS FOR DEFENDANT SAVANNAH SOUTHSIDE CLINIC, P.C.

Post Office Box 2139 Savannah, Georgia 31402-2139 (912) 236-2491

IN THE STATE COURT OF CHATHAM COUNTY

STATE OF GEORGIA

RYAN MACADAMS, A Minor, By)	
and Through His Natural Guardian and Parent as Next)	
Friend, D A W MACADAMS,)	
)	
Plaintiff,)	
)	
VS.)	CIVIL ACTION NO. 1971091G
DI LUE CUDICEODUED DOI EUNLADD)	
BLAKE CHRISTOPHER POLEYNARD,)	
M.D., FERNANDO JAVIER PEREZ, M.D., and SAVANNAH SOUTHSIDE)	
CLINIC, P.C.,)	
)	
Defendants.)	

CERTIFICATE OF SERVICE

I, PETER D. MULLER, **do** hereby certify that I have this day served a true and correct copy of the above and foregoing DEFENDANT SAVANNAH SOUTHSIDE CLINIC P.C.'S SUPPLEMENTAL RESPONSES TO PLAINTIFFS' FIRST INTERROGATORIES by causing same to be placed in the United States mail, postage prepaid, to:

Frederick S. Bergen, Esquire 123 E. Charlton Street Savannah, GA 31401

This ______ day of January, 1999.

BOUHAN, WILLIAMS & LEVY LLP

By: PETER D. MULLER

Post Office Box 2139 Savannah, Georgia 31402-2139 (912) 236-2491

IN THE STATE COURT OF CHATHAM COUNTY

STATE OF GEORGIA



DEFENDANT SAVANNAH SOUTHSIDE CLINIC, P.C.'S <u>SUPPLEMENTAL</u> <u>RESPONSES TO PLAINTIFFS' FIRST INTERROGATORIES</u>

COMES NOW Defendant SAVANNAH SOUTHSIDE CLINIC, P.C., and supplements

its responses to Plaintiffs' First Interrogatories as follows:

 (a) Michael S. Radetsky, MD CM Department of Pediatrics Loveiace Medical Center
 5400 Gibson Boulevard SE Albuquerque, NM 87108 (505) 262-3542

Dr. Radetsky is expected to testify on the issue of standard of care **and** causation, based upon various facts which have been presented in medical records and depositions.

Based upon the test results **and** findings of Dr Perez and Dr. Poleynard, Dr Poleynard acted **within** the standard **of** w e **in** his diagnosis, treatment, and discharge of Ryan MacAdams.

On the matter of causation, Dr. Radetsky is expected to testify that there **was** no causal connection between Ryan MacAdams not being diagnosed with meningococcal infection or treated with antibiotics during his first ER visit, and **his development of** orthopedic and other problem. Treatment with antibiotics **during the first visit** would not have prevented such problems. Also, it is likely hat the orthopedic problems were unrelated to the meningococcal infection.

DATED this _____ day of January, 1999.

BOUHAN, WILLIAMS & LEVY LLP

By:

PETER D. MULLER Georgia State Bar No. 528233 ATTORNEYS FOR DEFENDANT SAVANNAH SOUTHSIDE CLINIC, P.C.

Post Office Box 2139 Savannah, Georgia 31402-2139 (912) 236-2491

- 2 -

IN THE STATE COURT OF CHATHAM COUNTY

STATE OF GEORGIA

RYAN MACADAMS, A Minor, By)
and Through His Natural Guardian and Parent as Next	> -)
Friend, D A W MACADAMS,)
Plaintiff,))
VS.) CIVIL ACTION NO. 1971091G
BLAKE CHRISTOPHER POLEYNARD, M.D. FERNANDO JAVIER PEREZ, M.D. and SAVANNAH SOUTHSIDE CLINIC, P.C.,))
Defendants.	,)

CERTIFICATE OF SERVICE

I, PETER D. MULLER, do hereby certify that I have this day served a true and correct copy of the above and foregoing DEFENDANT SAVANNAH SOUTHSIDE CLINIC P.C.'S SUPPLEMENTAL RESPONSES TO PLAINTIFFS' FIRST INTERROGATORIES by causing same to be placed in the United States mail, postage prepaid, to:

Frederick S. Bergen, Esquire 123 E. Charlton Street Savannah, **GA** 31401

This _____ day of January, 1999.

BOUHAN, WILLIAMS & LEVY LLP

By: PETER D. MULLER

Post Office Box 2139 Savannah, Georgia 31402-2139 (912) 236-2491

CURRICULUM VITAE

NAME:

MICHAEL S RADETSKY MD CM

DATE AND PLACE OF BIRTH:

November 19, 1945 Denver, Colorado USA

PROFESSIONAL ADDRESS:

Department of Pediatrics Lovelace Medical Center 5400 Gibson Boulevard SE Albuquerque, NM 87 108 Telephone (505) 262-3542

CURRENT POSITION

Chairman, Department of Pediatrics Director of Newborn Medicine Consultant, Pediatric Infectious Disease Consultant, Pediatric Critical Care Lovelace Health System Albuquerque, New Mexico

Attending Physician Department of Pediatrics 'University of New Mexico Health Science Center

Clinical Professor of Pediatrics University of New Mexico School of Medicine Albuquerque, New Mexico

Fellow, Center for Public Policy and Contemporary Issues University of Denver Denver, Colorado

JOURNAL EDITOR

Editor

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Section on Pediatric and Neonatal Infections Current Opinion in Infectious Diseases 1993-1996

JOURNAL REVIEWER

Pediatric Infectious Disease Journal American Journal of Diseases of Children Journal of the American Medical Association Pediatrics Journal of Pediatrics Pediatric Emergency Care Yearbook of Pediatrics

UNDERGRADUATE EDUCATION:

Harvard College - A.B. cum laude, 1967

POSTGRADUATE ACTIVITIES

Harvard Law School - 1967-1968 (Courses passed. Leave of Absence)

United States Peace Corps, Malariologist Thailand - 1968-1969

Alternative National Service (Conscientious Objection) - Boston Floating Hospital for Children/Tufts New England Medical Center -1970-71

PREMEDICAL EDUCATION

University of Colorado Boulder, CO - 1972-1973

MEDICAL EDUCATION:

McGill University - Montreal, Quebec Canada - M.D.C.M. (Honors), 1973-1977

ROTATING INTERNSHIP:

San Francisco General Hospital, 1977-1978

PEDIATRIC INTERNSHIP:

University of Colorado School of Medicine Denver, Colorado, 1978-1979

RESIDENCY IN PEDIATRICS:

University of Colorado School of Medicine Denver, Colorado, 1979-1981

FELLOWSHIP TRAINING IN PEDIATRIC INFECTIOUS DISEASES

University of Colorado School of Medicine/The Children's Hospital Denver, Colorado, 1980-1982

<u>PRIOR POSITION:</u> 1991-1993

Director, Pediatric Critical Care Services Consultant, Pediatric Infectious Disease Kaiser Permanente Sacramento, CA

Attending Pediatric Intensivist University of California Medical Center Sacramento California

Pediatric Intensivist Sutter Memorial Hospital Sacramento California

Consultant, Pediatric Infectious Diseases University of California Medical Center Sacramento California

Clinical Professor of Pediatrics University of California School of Medicine Davis, California

Assistant Director of Pediatrics Director of Pediatric Hospital Services Director, Pediatric Critical Care Consultant, Pediatric Infectious Disease Denver General Hospital

Attending Physician Ambulatory Pediatric Clinic Denver General Hospital

PRIOR POSITION: 1989-1991

Attending Physician Intensive Care Unit The Childrens Hospital of Denver

Visiting Professor in Pediatrics Fitzsimmons Army Medical Center Denver, Colorado

Associate Professor of Pediatrics University of Colorado School of Medicine Denver, Colorado

Lecturer in Ethics Graduate School of Public Affairs University of Colorado, Denver Denver, Colorado

Lecturer in Medicine and Social Policy University of Denver Denver, Colorado

<u>PRIOR POSITION:</u> 1987-1989

Director of Pediatric Critical Care Services Tucson Medical Center Tucson, AZ

Chief of Pediatric Critical Care University Medical Center Tucson, **AZ**

Consultant in Pediatric Infectious Disease Tucson Medical Center, and University Medical Center Tucson, AZ

Clinical Associate Professor of Pediatrics University of Arizona School of Medicine Tucson, AZ

PRIOR POSITION:

1982-1987

Associate Director Infectious Disease Service The Children's Hospital Denver, Colorado

Attending Physician Intensive Care Unit The Children's Hospital of Denver

Assistant Professor of Pediatrics University of Colorado School of Medicine Denver, Colorado

Visiting Lecturer in Medical Ethics and Social Policy Department of Public Affairs University of Denver Denver, Colorado

Lecturer in Ethics Graduate School of Public Affairs University of Colorado, Denver Denver, Colorado

PRIVATE PEDIATRIC PRACTICE

Childrens Medical Center 1575 Vine Street Denver CO (Parttime coverage 1981-1987)

PROFESSIONAL LICENSURE:

State of New Mexico

State of Colorado

State of California (inactive)

State of Arizona (lapsed)

Medical Council of Canada

BOARD CERTIFICATION:

American Board of Pediatrics - 1983

Pediatric Critical Care American Board of Pediatrics - 1987 Recertification - 1995

Pediatric Infectious Disease American Board of Pediatrics - 1995

NATIONAL FACULTY:

Pediatric Advanced Life Support American Heart Association

HONORS AND AWARDS:

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Gold Medal Winner, Thames Cup Henley Royal Regatta, England - 1966

Haines' Memorial Award for Excellence in Athletics - Harvard College - 1967

Alexander Stewart Prize for Medical School Excellence, McGill Medical School, 1977.

Surgery Prize, McGill University, 1977,

Kaiser-Permanente Clinical Teaching Award Finalist, University of Colorado School of Medicine, 1983.

Kaiser-Permanente Clinical Teaching Award Finalist, University of Colorado School of Medicine, 1984.

Kaiser-Pemanente Clinical Teaching Award Finalist, University of Colorado School of Medicine, 1985.

The Gary Way Award for Outstanding Teaching Department of Pediatrics The Children's Hospital University of Colorado School of Medicine, 1985.

Kaiser-Pemanente Teaching Award Finalist in the Basic Sciences University of Colorado School of Medicine,
1986.

Commencement Speaker University of Colorado School of Medicine Hooding and Oath Ceremony, 1986.

Pediatric Housestaff Special Award Department of Pediatrics University of Colorado School of Medicine, 1987

Dean's List for Excellence in Teaching in the Clinical Sciences Award University of Arizona College of Medicine, Tucson, AZ, 1988

Finalist, Clinical Teaching Award University of Arizona College of Medicine, Tucson, AZ 1989

Commencement Speaker University of Colorado School of Medicine Hooding and Oath Ceremony, 1989

Attending Physician of the Year Family Practice Residency Program University of Arizona College of Medicine, 1989

Pediatric Residents' Teaching Award University of Arizona College of Medicine, 1989

The Gary Way Award for Outstanding Teaching Department of Pediatrics The Children's Hospital University of Colorado School of Medicine, 1990.

Honorary Faculty Membership Alpha Omega Alpha Honor Medical Society University of Colorado School of Medicine 1991

Joseph W. St. Geme Jr. MD Award for Exceptional Ability in Promoting the Overall Mission of the School of Medicine University of Colorado School of Medicine 1991

Excellence in Teaching Award Medical Student Council University of Colorado School of Medicine 1992

Clinical Teacher of the Year Department of Pediatrics University of California School of Medicine -Davis, 1992

Clinical Teacher of the Year Department of Pediatrics University of California School of Medicine -Davis, 1993

Inaugural Recipient William and Daniel Gelfand Lectureship The Childrens Hospital/University of Colorado 1994

Alpha Omega Alpha Annual Lectureship University of Arizona School of Medicine 1995

Selected The Best Doctors in America: Central Region 1996-1997

Sydney Gellis Lectureship Bostan Floating Hospital for Children Tufts University 1998

Selected Best Doctors in America 1999

VISITING PROFESSORSHIPS:

Hospital for Sick Children, Great Ormond Street, London, England 1982

Hospital for Sick Children, Great Ormond Street, London, England 1985

Duke University School of Medicine, Durham, NC 1987

Hospital for Sick Children, Great Ormond Street, London, England 1989

University of Florida School of Medicine, Jacksonville, Florida 1990

Akron Children's Hospital, Akron, Ohio 1990

Cincinnati Children's Hospital, Cincinnati, Ohio 1991

University of Minnesota School of Medicine 1991

Aglaia Kyriakou Children's Hospital Athens, Greece (invited only) 1991

Tkpler A m y Medical Center Honolulu, Hawaii 1991

Childrens Hdspital Omaha, Nebraska 1993

PROFESSIONAL ORGANIZATIONS

Infectious Disease Society of America

Pediatric Infectious Disease Society

Alpha Omega Alpha Honorary Fraternity

American Society for Law, Medicine and Ethics

PUBLICATIONS:

<u>Journals</u>

1) **Radetsky MS.** Recapturing the spirit in medicine. N Engl J Med 1978; 298:1142.

2) Radetsky MS, Istre GR, Johansen TL, Parmelee SW, Eauer BA, Wiesenthal AM, Glode MP. Multiply resistant pneumococcus causing meningitis: its epidemiology within a day-care centre, Lancet 1981; 2:771-773.

3) **Radetsky MS.** A diagnostic approach to Epstein-Barr virus infections. Pediatr Infect Dis 1982; 1:425-429.

4) **Radetsky MS**. Personal view: the hero in medicine. Brit Med J 1983; 287:493.

5) **Radetsky M**, Todd JK. Criteria for the evaluation of new diagnostic tests. Pediatr Infect Dis 1984;3:461-466.

6) **Radetsky MS**. The clinical evaluation of the febrile infant. Primary Care 1984; 11:395-405.

7) **Radetsky M**, Wheeler RC, Roe MH, Todd JK. Comparative evaluation of kits for rapid diagnosis of group A streptococcal disease. Pediatr Infect Dis 1985; 4:274-281.

8) **Radetsky M.** The rise of the academic clinician. Am J Dis Child 1985; 139:861.

9) **Radetsky M.** Sudden intimacies. JAMA 1985;254:1361; reprinted in: Dan BB, Young RK (eds). A Piece of My Mind, 1988; New York: AA Knopf.

10) **Radetsky M.** Laboratory evaluation of acute diarrhea. Pediatr Infect Dis 1986; 5:230-238.

11) **Radetsky M**, Wheeler RC, Roe MH, Todd JK. Microtiter broth dilution method for yeast susceptibility testing with validation by clinical outcome. J Clin Microbiol 1986;24:600-606.

12) **Radetsky M**, Soloman JA, Todd JK. Identification of streptococcal pharyngitis in the office laboratory: reassessment of new technology. Pediatr Inf Dis 1987; 6: 556-563.

13) **Radetsky M.** Duration of treatment in bacterial **meningitis**: a historical **inquiry.** Pediatr Infect Dis J 1990; 9: 2-9.

14) **Radetsky M.** Duration of Symptoms and Outcome in Bacterial Meningitis: An Analysis of Causation and the Implications of a Delay in Diagnosis. Pediatr Infect Dis J 1992;11: 694-8.

15) **Radetsky M.** Pediatric and neonatal infections: Editorial overview, Curr Opinion Infect Dis 1993;6:545-6.

16) **Radetsky M**. Infectious disease emergencies. Curr Opinion Pediatr 1994; 6: 310-6.

17) **Radetsky M.** The timing of antimicrobial therapy and outcome in serious bacterial infections. Curr Opinion Infect Dis 1994;7:341-4.

18) **Radetsky M.** The laboratory evaluation of newborn sepsis. Curr Opinion Infect Dis 1995;8:191-9.

19) **Radetsky M.** Use of antimicrobials for the prevention of recurrent urinary infection in children. Dialogues in Pediatric Urology 1995;18:7-8.

20) **Radetsky M.** The febrile infant and the assumption of risk. Curr Opinion Infect Dis 1996;9:171-5.

21) Radetsky M. The discovery of penicillin. Pediatr Infect Dis J 1996;15:811-8.

22) **Radetsky M.** The newborn at risk for serious infections. In: Britton JR (ed). Issues related to early discharged newborn infants and their mothers. Clinics in Perinatology, 1998;25:327-34.

23) **Radetsky M**. The emerging spectrum of tickborne infections. Curr Opinion Infect Dis 1998;11:313-8.

Letters to the Editor

1) **Radetsky M**. Timing of therapy for meningococcal infection. Pediatr Infect Dis J 1997;16:540-1.

2) Radetsky M. Prognostic factors in meningococcal disease. JAMA 1997;278:1658.

3) **Radetsky M**. The social missions of academic health centers. New Engl J Med 1998;338: 1232.

4) Radetsky M. On the front lines of medicine. Time, November 2, 1998;152 (18):22.

Book Chapters

1) **Radetsky M**. A clinical approach to the diagnosis of streptococcal pharyngitis. In: Barkin RM (ed.) The Emergently III Child, Rockville MD, Aspen Publishers, 1987.

2) **Radetsky M.** The Nature and History of Medical Ethics. In: Nussbaum E. (ed.) Pediatric Intensive Care, 2nd ed., 1989, Mt. Sisco NY, Futura Publishing. Reviewed in: Arch Dis Child 1990; 65:816.

3) Radetsky M. The Doctrine of Informed Consent. In: Nussbaum E. (ed.) Pediatric Intensive Care, 2nd ed., 1989, Mt. Sisco NY, Futura Publishing. Reviewed in: Arch Dis Child 1990;65:816.

4) **Radetsky M**. Decisions to Limit, Diminish, or Withdraw Therapy. In: Nussbaum E. (ed.) Pediatric Intensive Care, 2nd ed., 1989, Mt. Sisco NY, Futura Publishing. Reviewed in: Arch Dis Child 1990;65:816.

5) **Radetsky M.** The Definition and Determination of Death. In: Nussbaum E. (ed.) Pediatric Intensive Care, 2nd ed., 1989, Mt. Sisco NY, Futura Publishing. Reviewed in: Arch Dis Child 1990;65:816.

6) **Radetsky M.** Enterobacteriaceae. In: Patrick C (ed.) Infections in Immunocompromised Infants and Children, 1992, New York, Churchill Livingston.

7) **Radetsky M.** The Use of Antimicrobials and a Synopsis of Infectious Disease in the Pediatric Intensive Care Unit. In: Fuhrman BP, Zimmerman JJ. (eds.) Pediatric Critical Care, 1992, St. Louis, Mosby Year Book

8) **Radetsky M.** Exanthematous Viral Infections. In: McAnarney ER, Kreipe RE, Orr DP, Comerci GD (eds.) Textbook of Adolescent Medicine, 1992, Philadelphia, W.B. Saunders.

9) **Radetsky M.** Streptococcal Infections. In: Burg FD, Ingelfinger JR, Wald ER (eds.) Current Pediatric Therapy 14, 1993, Philadelphia, W.B. Saunders.

10) **Radetsky M.** Streptococcal Infections. In: Burg FD, Ingelfinger JR, Wald ER (eds.) Current Pediatric Therapy 15, 1995; Philadelphia, WB Saunders

11) **Radetsky M.** Staphylococcal Infections. In: Burg FD, Ingelfinger JR, Wald ER (eds.) Current Pediatric Therapy 15, 1995; Philadelphia, WB Saunders..

12) **Radetsky M, Overturf GD.** Epstein-Barr Infections in Adolescents and Young Adults. In: Overturf GD, Jacobs RF (eds.). Adolescent Medicine: Viral Infections of the Adolescent, 1995, Philadelphia, Hanley and Belfus.

13) **Radetsky M.** Antibiotic; use in pediatric critical care. In: Fuhrman BP, Zimmerman JJ. (eds.) Pediatric Critical Care, 2nd ed., 1998, St. Louis, Mosby Year Book.

14) **Radetsky M.** Shigella Infections. In: Stockman JA, Lohn JA (eds.) Ambulatory Pediatrics, Philadelphia, WB Saunders (in press).

D) Abstracts

1) **Radetsky MS**, Glode MP, Istre GR, Lauer BA, Wiesenthal AM. Emergence of multiply resistant pneumococcus. Presented at the 21st Interscience Conference on Antimicrobial Agents and Chemotherapy. Chicago, November 1981.

2) Todd JK, Parmelee SW, **Radetsky M**. Antimicrobial combination interactions with Streptococcus pneumoniae and Haemophilus influenzae. Presented at the 22nd Interscience Conference on Antimicrobial Agents and Chemotherapy. Miami, October 1982. Abstract 222.

3) Wheeler RC, **Radetsky MS**, Roe MH, Todd JK. Comparison of two candida antigen detection systems for identifying patients with disseminated candidiasis. Presented at the 25th Interscience Conference on Antimicrobial Agents and Chemotherapy. Minneapolis, October 1985. Abstract 760.