Page I IN THE COURT OF COMMON PLEAS 1 CUYAHOGA COUNTY, OHIO 2 MICHELLE KASCHAK, et al., 3 4 Plaintiffs, 5 Case No. CV 3551360 vs UHHS BEDFORD MEDICAL 6 CENTER, et al., 7 Defendants. 8 9 10 11 DEPOSITION OF ARTHUR B. ZINN, M.D., Ph.D. 12 THURSDAY, AUGUST 17, 2000 13 The deposition of ARTHUR B. ZINN, M.D., 14 15 Ph.D., the Witness herein, called by counsel on behalf of the Plaintiff for examination under the 16 statute, taken before me, Vivian L. Gordon, a 17 Registered Diplomate Reporter and Notary Public 18 in and for the State of Ohio, pursuant to 19 agreement of counsel, at the offices of Reminger 20 & Reminger, The 113 St. Clair Building, 21 22 Cleveland, Ohio, commencing at 1:30 o'clock p.m. on the day and date above set forth. 23 24 25

Page 2 **APPEARANCES:** 1 2 3 On behalf of the Plaintiff Becker & Mishkind BY: HOWARD D. MISHKIND, ESQ. 4 Skylight Office Tower Suite 660 Cleveland, Ohio 44113 5 6 On behalf of the Defendant University Hospitals Moscarino & Treu BY: KEVIN M. NORCHI, ESQ. 7 630 Hanna Building Cleveland, Ohio 44115 8 On behalf of the Witness: 9 Reminger & Reminger BY: P.J. MALNAR, ESQ. 10 The 113 St. Clair Building Cleveland, Ohio 44113 11 ALSO PRESENT: 12 Michelle Kaschak Michael Kaschak 13 14 15 16 17 18 19 20 21 22 23 24 25

Page 3 1 ARTHUR B. ZINN, M.D., Ph.D., a witness 2 herein, called for examination, as provided by 3 the Ohio Rules of Civil Procedure, being by me first duly sworn, as hereinafter certified, was 4 5 deposed and said as follows: 6 EXAMINATION OF ARTHUR B. ZINN, M.D., Ph.D. BY MR. MISHKIND: 7 MR. MISHKIND: The record should 8 9 reflect that we are here today on August 17th, The purpose of this deposition is to 10 2000. obtain the facts and information known to Dr. 11 12 Arthur Zinn who was involved in certain aspects of Megan Kaschak's care at or around the time of 13 her demise. 14 15 The deposition is being taken by agreement. Ms. Malnar has arranged to have 16 17 Dr. Zinn here. Dr. Zinn has made himself available without the issuance of a subpoena to 18 answer questions. The Kaschaks are also here. 19 Certainly no issues of waiver of 20 21 privilege -- there are no privileges that are being withheld. 22 23 MS. MALNAR: Fair enough. Q. 24 Let's start out with an easy question 25 for you. Tell us who you are.

Page 4 1 Α. My name is Arthur Zinn. Q. 2 And what is your occupation? Α. I'm a medical geneticist. 3 Ο. 4 Could you trace briefly for me your educational background. 5 I grew up in New York City, went to 6 Α. 7 public schools, went to college and got a bachelor of arts in chemistry from Brandeis. 8 Ι came to Case Western Reserve University where I 9 got my M.D. and Ph.D. in biochemistry. 10 I then went to the University of Minnesota where I 11 trained in pediatrics. I then went to Yale 12 University where I trained in human genetics. 13 Q. 14 Where is your practice situated currently? 15 My main office is at University 16 Α. Hospitals of Cleveland. 17 Q, And just for housekeeping purposes, 18 19 are you an employee of University Hospitals or are you employed by some other entity? 20 My paycheck comes from Case Western 21 Α. Reserve University. 22 Q. Are you affiliated with some practice 23 24 group? I think I am now. I think it's called 25 Α.

Page 5 University Genetics, Inc. I'm not sure there was 1 2 such an Inc. when I spoke with the Kaschaks. Q. 3 Your M.D. degree from Case was 4 obtained in what year, approximately? Α. 76 5 Ο. Did you then immediately pursue your 6 7 Ph.D.? They were done concurrently. 8 Α. Q. 9 Dual degrees? I was awarded the Ph.D. in January 10 Α. 11 '77. Q. Your training then at University of 12 Minnesota was in what? 13 14 Α. Pediatrics. Q. Was it a straight residency in 15 pediatrics? 16 It was a residency in pediatrics. 17 Α. 18 0. How many years were you there? 19 Α. About two and a half. And then at Yale, it was in human 20 0. genetics? 21 22 Α. Correct. How many years were you there, sir? 23 Q. A little over two. Α. 24 At the beginning of the deposition, 25 Q.

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Page 6 1 you said that you are a medical geneticist. 2 Α. Correct. Q. Is there a board certification in that 3 subspecialty? 4 Α. 5 Yes. 0. Are you board certified? 6 Α. I'm certified by the American Board of 7 Medical Genetics in both clinical genetics and 8 clinica biochemical genetics. 9 Q, Do *you* have any other board 10 11 certification outside of the area of genetics? Α. No. 12 MS. MALNAR: One thing I should have 13 mentioned to you, if you let Mr. Mishkind finish 14 his questions before you start your answers, that 15 will make things easier. 16 Q. Do **you** have a relatively recent or 17 current curriculum vitae? 18 Α. 19 Yes. Q. Do you have it with you? 20 21 Α. No. Q. Would it be difficult for you to make 22 it available to Ms. Malnar, assuming she doesn't 23 already have it in her hands? 24 I can do that. 25 Α.

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Page 7 1 Ο. If you would, please, I would 2 appreciate it. 3 Your employer then today is Case 4 Western Reserve University or is it University Genetics, Inc.? 5 6 Case Western Reserve. Α. 0. 7 Your office is located where? 8 In University Hospitals of Cleveland. Α. Q. 9 In what department or division is the office located at the hospital? 10 11 The Center for Human Genetics. Α. 12 Ο. After you finished your training at 13 Yale, did you then come to University Hospitals? 14 Α. I then came to Cleveland to my present position. 15 Ο. You have been in this present position 16 for how many years, sir? 17 18 Α. Since 1982. Ο. The Center for Human Genetics consists 19 20 of you and how many other geneticists are in that 21 division? 22 Α. It's not a division, it's a 23 department. And it's a department, I think, of 24 about 25 faculty, not all of whom have clinical 25 responsibility.

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Page 8 1 0. Can you tell me the names of those that do have clinical responsibility? 2 3 Α. When? 4 Q. Good question. At or around the time 5 of the events that concern Megan back in January, February, '97. 6 7 Suzanne Cassidy, Nathaniel Robin, Α. 8 Matt Warman, and Georgia Wiesner are the other 9 physicians. There are other people who have clinical responsibility, but they are primarily 10 laboratory based. 11 12 Ο. In terms of the genetic issues that 13 were involved in Megan's case, were you the only medical geneticist that was involved or were 14 there others? 15 First of all, I was not involved. Α. 16 Second, the patient was seen by Nat Robin in 17 consultation. 18 Q. 19 Is Dr. Robin still in the department? 20 Α. Yes. Q. 21 When did you become involved in one way or another in terms of disseminating or 22 23 communicating any information? 24 After Megan's death. Α. Q. 25 How is it that you were chosen to be

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1	involved postmortem?
2	A. The question was whether or not Megan
3	might have had a metabolic genetic disorder, and
4	there is a series of people who had made various
5	recommendations, and the thought was I could help
6	expedite the postmortem studies so that those
7	questions could be addressed appropriately.
8	Q. Do you have a recollection as to who
9	those people were that thought you might be able
10	to expedite some of the answers to those
11	questions?
12	A. I believe it was the neonatologist who
13	called me.
14	Q. Do you remember which neonatologist
15	that was?
16	A. Michele Walsh-Sukys.
17	Q. Is Dr. Walsh-Sukys still at RB&C, to
18	your knowledge?
19	A. Yes.
20	Q. Some time ago when we first attempted
21	to schedule your deposition, there had been a
22	request made that you bring with you a complete
23	copy of your office chart or records regarding
24	Megan that would include autopsy results and the
25	results from any metabolic tests that had been

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Page 10 performed. 1 2 Have you brought any such documents with you today? 3 The documents that he 4 MS. MALNAR: brought with him are copies of the documents that 5 I faxed to you today. 6 7 MR. MISHKIND: I didn't get a fax from you today. When did you fax that? 8 MS. MALNAR: Did you get a fax from me 9 today? 10 MR. NORCHI: It was around 11:30, 11 12:00. 12 MR. MISHKIND: I was out of the 13 14 office. MS, MALNAR: I can make another copy 15 of it now for you, if you like. 16 (Thereupon, a recess was taken.) 17 Q. The fax that came over from Ms. Malnar 18 19 at 11:30, 12:00 o'clock, I had already left my office, so I have not seen that, but we are in 20 the process of photocopying it and I will have 21 questions for you relative to that. 22 But are these results of certain 23 24 metabolic tests that had been performed on Megan? They are the results of either 25 Α.

Page 11 enzymatic or DNA based molecular tests, yes. 1 Ο. Are these results of tests that were 2 done in-house or that were sent out elsewhere? 3 Α. The tests were done either at a 4 5 laboratory that physically is located in Rainbow, a laboratory that's part of a Rainbow Center 6 center that's at the V.A. Hospital in Cleveland, 7 a laboratory that's -- I can't remember where it 8 was back then, but it is part of the genetics 9 department -- and a laboratory that's in 10 California. 11 Ο. Where are these records maintained, 12 the originals of what you have brought with you 13 today? 14 They are put in an envelope and put 15 Α. somewhere on my bookshelf. 16 And what you brought with you today, 17 Ο. does that constitute all of the test results that 18 you have obtained back relative to Megan? 19 Α. Those are the test results that I had 20 some knowledge or some involvement with. There 21 are laboratory tests that were done prior to 22 Megan's death that **I** did not keep records of and 23 don't have, didn't seek to. 24 Q. 25 **In** a moment, doctor, I'm going to have

Page 12 1 these documents marked as an exhibit for purposes 2 of the transcript, but before I do that, let me ask you, there is reference, in addition to 3 yourself, there is reference to Dr. Kerr and 4 there is also reference to Stuart Schwartz that 5 was somehow involved in the interpretation of 6 some of this information; is that correct? 7 8 Α. I'm sorry, do that again, sir. Q. 9 Dr. Kerr was involved in apparently the interpretation of some of the test results? 10 Oh, I see. Yes, Dr. Kerr signed, I 11 Α. think, two of the reports. Dr. Schwartz signed, 12 I think, two of the reports. I signed none of 13 them since I didn't perform them. 14 I didn't interpret any of the primary results. 15 Q. Were there any tests that you had 16 recommended that were never done in this case? 17 Not to my knowledge. 18 Α. 19 Ο, **So** that all of the tests that you believe were reasonably appropriate to do to rule 20 out or to confirm some type of a metabolic or 21 genetic disorder were, in fact, done; is that a 22 fair statement? 23 24 Α. The list that was provided to me from the assessment prior to Megan's death, I saw to 25

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1	it that that list got executed appropriately.
2	Q. And as you look back at that list,
3	would there have been any additional tests,
4	metabolic or genetic tests, that in your
5	professional opinion should have been performed
6	in order to rule out any loose ends, if you will?
7	A. At the time \mathbf{I} thought what was done
8	was what should have been done.
9	Q. And as you sit here now, do <i>you</i> still
10	hold that same opinion?
11	A. Actually I haven't thought about that.
12	Q. Is there anything that comes to mind
13	as you are reflecting on it now that it would
14	have been nice to have done to have ruled out or
15	to confirm things?
16	A. I have to remember. You are asking me
17	if there is something nice to have done. The
18	answer to that is always yes. You always want
19	more information, but the answer to the question
20	you want, I think, is should more studies have
21	been done, given the evidence at that time, and I
22	think the answer is no.
23	Q. In a moment I'm going to hand these
24	documents to Vivian. Let me ask you one
25	question.

Page 14 1 Given the fact that I have not had a 2 chance to read through these documents and 3 without taking probably more time than any of us 4 have today to study them, can you tell me whether 5 these test results, which I have not seen before, 6 Whether or not they permit you to opine that there was any type of a metabolic or genetic 7 disorder that caused Megan to experience her 8 9 cardiopulmonary arrest? My assessment of reviewing these 10 Α. results when they became available was that I 11 could not make a diagnosis based on these 12 results. 13 Ο. Do you use, from a professional 14 standpoint, the term that the results are 15 inconclusive as a descriptive term? 16 17 Α. I sometimes use that term. Ο, 18 And might that have been a term that you used on one or more occasions when you talked 19 with Mrs. Kaschak? 20 21 Α. Again, we had multiple conversations. I cannot remember all the conversations in 22 My feeling at the time was that rather detail. 23 24 than saying inconclusive, what I would say, they are not consistent within themselves and do not 25

Page 15 allow me to make a specific diagnosis. 1 2 Ο, And again, the diagnosis that you were looking to make was to determine whether or not 3 4 there was some type of a metabolic or genetic 5 abnormality, if you will, or error, inborn error 6 that caused Megan on her first day of life to experience this cardiopulmonary arrest; correct? 7 8 Α. Yes. 9 Q. And then, carrying that further, whether there was any metabolic or genetic 10 disorder that prevented Megan from surviving the 11 results of this cardiopulmonary arrest? 12 I probably wasn't trying to address 13 Α. 14 that question. 15 Q. So you were focusing --16 I was trying to see with the results Α. of completing the studies of the people who 17 evaluated her, and **I** considered on their list 18 whether those things were explanations of what 19 was going on and if any of the results, if they 20 came back one way or another, would suggest 21 further testing that I should do to either prove 22 or disprove those initial hypotheses. 23 24 Q. The focus then -- and I don't mean to 25 repeat --

Page 16 I'm trying to make sure I answer the 1 Α. question in a way that \mathbf{I} am comfortable with. 2 Q, And **I** am trying to understand 3 Sure. 4 it in a way that **I** am comfortable. T understand. 5 Α. Q. **So** I am not trying to be difficult. 6 The evaluation was to determine 7 whether or not there was some metabolic or 8 9 genetic disorder that Megan had that is the more probable or likely explanation for why she 10 11 experienced a cardiopulmonary arrest; true? 12 Α. Yes. a. And based upon all of the tests, there 13 is insufficient evidence for you to provide an 14 opinion that her cardiopulmonary arrest was 15 caused by a metabolic or genetic error or genetic 16 abnormality? 17 18 Α. That's correct. MR. MISHKIND: Let's go ahead and mark 19 this as an exhibit. I'm not going to go through 20 it; perhaps Mr. Norchi might, since he has the 21 benefit of hours of reading it over better than I 22 have. And what I may do is defer to Mr. Norchi 23 24 and then read it while he is asking you questions and then maybe come back afterwards to try to 25

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Page 17 save some time. 1 2 MR. MISHKIND: Let's mark this as Plaintiff's Exhibit 1. 3 4 (Thereupon, Plaintiff's Deposition 5 Exhibit 1 was marked for 6 purposes of identification.) 7 8 9 Ο. Just for purposes of bookkeeping, Plaintiff's Exhibit 1 is a copy of the various 10 11 interpretations and test results that you have provided today and that were within your 12 possession from the Center for Human Genetics 13 14 relative to Megan; is that correct? Α. They were in my possession because 15 they were mailed to me. 16 Q. 17 Are there other tests -- strike that. 18 Are these all postmortem tests? 19 Α. Yes. Q. As I understand it, the autopsy was 20 done within 40 minutes of Megan's death? 21 I don't know the details. I know it 22 Α. was done rapidly because of the requirements to 23 24 try to preserve tissue. Q. What we have here are interpretations 25

Page 18 1 and reports relative to what was done on a 2 postmortem basis on these various tests; true? 3 Α. Right. They were done not within that 4 40 minute period. The tissue was collected, 5 frozen, and then --6 THE WITNESS: I'm sorry to be so 7 graphic. 8 MR. MISHKIND: They realize. 9 THE WITNESS: I feel bad. 10 MR. MISHKIND: I appreciate that, 11 doctor. 12 I will finish my answer. Yes, the Α. 13 tissues were collected and frozen, and as the 14 tests proceeded, samples were analyzed. As you look at what we have here in 15 Q. Plaintiff's Exhibit 1, which now has an exhibit 16 17 sticker and a staple preserving all the documents, are there other reports or 18 interpretations that, for whatever reason, aren't 19 contained in Exhibit 1 that relate to tests that 20 21 were ordered postmortem? 22 Not to my knowledge. Α. Q. I want to ask you about certain 23 24 communications that you may or may not have had with Michelle during this order. 25

Page 19 1 Α. Mrs. Kaschak? Q. 2 Yes. According to information that I have 3 4 from Michelle, you spoke to her in August of '97 indicating that you expected results from 5 6 biochemical tests to be available in about a 7 month or sometime in September of 1997. 8 I don't expect that you can remember 9 the exact date, but during the summer of 1997, were there some biochemical tests that you were 10 11 waiting the results on? I believe so. I don't know the 12 Α. interval. 13 What was the date of Megan's death? 14 Q. February of 1997. 15 That sounds correct. Α. 16 Q. 17 And the biochemical tests, are they contained within the packet that is Exhibit 1? 18 19 Α. Yes. Q. Can you refer to what the document is 20 that show the test results? 21 22 Α. Do you want specifically which test I was waiting for on that particular phone call? 23 Q. 24 At that particular time that would refer to biochemical tests that you were waiting 25

Page 20 1 for in August of '97. 2 Then I would say the packet refers to, Α. 3 in some broad way, what **I** was waiting for. Q. All of those are biochemical tests? 4 Α. Maybe I should define the terms. 5 Ο. All right. 6 7 Many **of** these reports are enzyme Α. analyses, which is a form of biochemical 8 measurement. Other tests are DNA based molecular 9 testing, which some consider biochemical and 10 others don't. But they are all metabolic studies 11 12 designed to elaborate the cause of a metabolic disease. These are all metabolic tests. 13 Q . 14 Do you recall ever indicating to Michelle the results on the phone as to the 15 results of the biochemical tests, whether enzyme 16 17 or otherwise? Yes. 18 Α. Q. And what is your recollection of what 19 20 you told her? **I** told her what I had received and 21 Α. 22 what we had found. Q. And the bottom line in terms of what 23 you found and what you would have told her would 24 be what? 25

Page 21 1 Α. No doubt, it changed somewhat as we 2 went through the conversations, but initially I said I don't have an answer, and I probably said 3 I don't have an answer in each of our phone 4 calls. 5 I would also say that we have a 6 7 result; I'm not sure what it means, or it shows this, but we need to wait for something else to 8 see if that's been confirmed by the other 9 studies. 10 I would say that was the content of 11 12 most of the conversations. Ο. Do you recall ever having a 13 14 conversation with Michelle where you gave her the final report on all of the tests, after having 15 all of the information back over the many months 16 17 that this process took place? To the extent that I was uncomfortable Α. 18 19 doing everything on the phone, I would say I gave a final report. And I said that we don't have an 20 21 answer for what happened to your baby. Q. 22 Do you recall indicating to Mrs. Kaschak at one time or another that the 23 24 neurological changes that occurred to Megan appeared to be most likely secondary to the 25

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Page 22 1 hypoxic episode that she experienced on her first day of life? 2 I don't think I would have commented 3 Α. on that. 4 Q. 5 Tell me why is that. One, I wasn't a witness to those 6 Α. 7 events, and two, that's a neuropathology 8 interpretation, and I don't think I could say more or less likely to that. 9 Q. 10 Have you had a chance to review the 11 autopsy? 12 Α. Yes. Ο. 13 Do you have a copy of the autopsy with 14 you today? 15 Α. No. Q. To the extent that you need it in 16 responding then to any of my questions, I'm going 17 to go ahead and mark a copy of the autopsy as 18 Plaintiff's Exhibit 2, and feel free to reference 19 20 it as necessary. 21 (Thereupon, Plaintiff's Deposition 22 Exhibit 2 was marked for 23 24 purposes of identification.) 25

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1	MS. MALNAR: May I?
2	MR. MISHKIND: Sure.
3	A. Can you repeat your last question, the
4	one about the hypoxic changes?
5	Q. Sure.
6	What ${f I}$ asked you was whether you
7	recalled indicating to Mrs. Kaschak that the
8	neurological changes that occurred to Megan
9	appeared to be most likely secondary to the
10	hypoxic episode that she experienced on her first
11	day of life.
12	A. Okay. Let me amplify my answer. Is
13	that permissible?
14	Q. Go right ahead.
15	A. I hope what I said I don't think I
16	would've said more likely, because I don't think
17	that would be my position to do that, and I think
18	it would have been presumptuous.
19	I think at one time ${\tt I}$ might have said
20	could be consistent with, but I don't think I
21	would have said more likely.
22	Q. Were you able to provide her with any
23	explanation that was equally likely as being the
24	cause of the neurological changes, other than the
25	hypoxic episode she experienced?

Page 24 1 To answer that, then, I think I would Α. 2 have focused on the metabolic potential cause, and at no time did I have an explanation 3 4 sufficient to provide her with that explanation. Ο. So that you considered the hypoxic 5 episode as being one of a number of potential 6 7 explanations for the neurological changes and you also considered metabolic abnormalities as a 8 potential explanation for the neurological 9 changes? 10 Objection. 11 MR. NORCHI: 12 The neuropathologic? Α. Q. 13 Yes. **I** would leave that to the 14 Α. neuropathologist. 15 See, the answer -- I'm not sure what 16 your objection is. 17 Q. Don't worry about the objection. 18 19 Α. Sorry. Q. Sometimes lawyers object because they 20 21 know why they are and sometimes they object 22 because they feel they have to. MR. NORCHI: I know why I am 23 objecting, Howard. 24 THE WITNESS: I'm sorry, **I** am trying 25

Page 25 to answer people's questions. 1 2 Go ahead, doctor. MR. NORCHI: 3 Α. The question I would have, if I was trying to discuss that with Mrs. Kaschak, would 4 be, do I know the cause or the effect. I mean, 5 if there was a metabolic disorder that caused the 6 7 child to have a cardiorespiratory arrest, then 8 you can have problems with the arrest itself. 9 It's a question of what caused the And so that's why I couldn't tell cause 10 arrest. 11 and effect and why I wouldn't comment further. 12 Q. As you sit here now, do you have a 13 basis to say to a reasonable degree of medical probability that you know what the cause of the 14 15 cardiopulmonary arrest was? 16 I do not know the cause of that. Α. 17 Q. **Do** you have a basis to a reasonable degree of medical probability to say that you 18 know what the cause of the hypoxic ischemic 19 20 encephalopathy was that the baby suffered? 21 I don't have a comment on that. Α. Ι 22 don't have a position on that. 23 Q. Is there a reason that you don't have a position on that? 24 25 Α. Because I think the neuropathologists

Page 26 1 have to comment on -- I mean, it's a neuropathology diagnosis of what they see and I 2 3 didn't focus on that. Q. 4 In looking at the autopsy for a moment, if you could turn to page four. In the 5 6 summary and comments section --7 Α. Yes. Q. 8 -- the second paragraph, just so that 9 we are referencing it, it says abnormal 10 significant autopsy results fell into two major 11 categories. 12 Α. Yes. Ο. One being severe hypotensive/ischemic 13 encephalopathy, which is attributable to cardiac 14 arrest on the first day of life, and two, 15 excessive glycogen deposition in liver, skeletal 16 17 muscle and cardiac muscle. 18 Did I read that correctly? 19 Α. Correct. 0. 20 Do you have an opinion as to the significance of the excessive glycogen 21 depositions that were found in the liver, the 22 23 skeletal muscle and the cardiac muscle? 24 I don't have an explanation for it. Α. Q. 25 Are there factors that you considered

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1	as possible explanations for that?
2	A. I thought it was either related to the
3	events experienced prior to Megan's death or it
4	could have been secondary to a metabolic
5	disorder.
6	Q. Were you ever able to attribute the
7	glycogen deposition in the liver, skeletal muscle
8	and cardiac muscle to a metabolic disorder?
9	A. No.
10	Q. You said the other possibility was to
11	events that occurred prior to her death. What do
12	you mean by that?
13	A. That as you experience the severe
14	problems that Megan had, you can have tissue
15	damage that effects mitochondrial function. You
16	may not be able to use glucose, break down
17	glycogen and use the glucose; especially when you
18	place the child on large amounts of intravenous
19	sugar, that sugar has no place to go and the
20	cells tend to store it.
21	Q. Who prepared the autopsy report
22	itself?
23	A. I presume the pediatric pathologist.
24	${\tt I}$ believe Dr. Beverly Dahms signed it and I don't
25	know if a resident helped her or not. I don't

Page 28 remember the names on it. Maybe it says. 1 Q. 2 I believe Beverly Dahms is at least the major person involved. 3 She is the staff pathologist. 4 Α. Ο. Did you have any ongoing discussion or 5 conversation with Dr. Dahms as the tests were 6 7 being ordered and the results were coming back postmortem? 8 We spoke on a number of occasions. 9 Α. **a** . 10 Tell me to the best of your recollection, either specifically or in general, 11 12 what was discussed and what information you likely imparted to her or she to you. 13 The initial conversation had to do 14 Α. with what she would like me to do to help her, 15 and what I told her I would need from her. 16 So I asked her basically to do as best 17 a job as she could and provide me with as much 18 information as she could and then to let me know. 19 And she had the results, and on the second visit, 20 third, or whatever it was, I went over and 21 22 discussed those results. We spoke about the glycogen deposition 23 at some length. We tried to see if that was a 24 good clue or that might help me establish a 25

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course for further study. We looked at all the 1 2 related causes of glycogen deposition in inborns and we could not match those with the pattern we 3 could see. 4 I could not do that alone. Dr. Dahms 5 6 had to look at the cells. She had to tell me where they were in the cell, if they had 7 8 membranes around it. 9 I don't mean to go through the particulars, but that's what she could tell me. 10 11 And based on that, I thought it was likely that the glycogen was secondary, if anything, to a 12 13 mitochondria, as people call it, or oxidative phosphorylation defect. 14 The decision was made to focus on that 15 as previously suggested prior to Megan's death by 16 her attending physicians and some of the 17 consultations. 18 Q. Did you ultimately rule out or confirm 19 the existence of that condition? 20 We could not establish that she had an 21 Α. electron chain transport defect. 22 & • So that from a probable --23 I should probably still 24 Α. finish answering your previous question. 25

Page 30 Q. Go ahead. 1 2 Α. Then I also got back to Dr. Dahms and told her what we had found and hadn't found and I 3 informed her that we couldn't come up with a 4 consistent metabolic pattern that would allow me 5 to give her a final diagnosis. 6 I'm not sure how many conversations 7 that was, but I would say that would be the three 8 stages of our conversations. 9 10 Q. Did there ever come a time that you 11 attempted to set up a meeting to talk with the Kaschaks about the final results? 12 We spoke a number of times about that. 13 Α. Q. Did you ever have an in-person 14 meeting? 15 No, we didn't. 16 Α. 17 Q. Do you know why that is? I just know it wasn't possible to set 18 Α. it up with Mrs. Kaschak on the phone. 19 I mean, my 20 impression was it was just too difficult for her. Q. 21 Fair enough. But I don't know that for sure. 22 Α. Q. Again, referring to the autopsy, 23 doctor, to my knowledge in reading the deposition 24 25 (SIC), the placenta was normal on review.

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Page 31 1 Was that the information that you had, as well? 2 MS. MALNAR: You are reading the 3 4 autopsy? 5 MR, MISHKIND: Yes. MS. MALNAR: You said deposition. 6 I'm sorry, where am I looking? 7 Α. Q. 8 On page four under the summary and comments, the third paragraph, about five lines 9 down. 10 11 Α. The placenta was normal on subsequent review, and what is your question? 12 Q. 13 If the placenta was abnormal, would that cause you to have --14 The placenta was normal, however. 15 Α. 16 Q. If it had been abnormal, would that have caused you to consider other factors being 17 18 contributory to the baby's cardiopulmonary 19 arrest? 20 One, I would have spoken with the Α. 21 pathologist and asked him what the abnormality was and what they thought it could be. 22 That would generally be something they would tell me 23 24 as opposed to the opposite. Q. 25 Now, there is a note --

Page 32 1 Α. I'm slower than you. Q. 2 Okay, that's all right. I don't mean to cut you off. 3 The fact is, if the pathologist would 4 Α. have reviewed the placenta and said they found 5 6 something unusual or something suggestive, we often would get involved in that, because part of 7 medical genetics, it deals with things like 8 that. So I might have done something else I 9 think is what you are really asking me. 10 Q, 11 Correct. But that was not said to me. 12 Α. Ο. **I** am looking to see whether you can 13 14 help me with certain terms in the autopsy. Ι recognize you did not prepare it, but certainly 15 you are familiar with the terminology as it 16 relates to this baby, and hopefully you can help 17 18 me. 19 The bottom of that page where it says certainly this prolonged cardiorespiratory arrest 20 could explain the severe hypotensive/ischemic 21 encephalopathy observed at autopsy --22 23 Yes. Α. 24 Q. -- can you explain what that means? The brain is absolutely dependent upon 2**5** Α.

Page 33 1 oxygen to function. In the absence of oxygen, 2 the brain cannot function and undergoes destructive changes. 3 4 If you do not have adequate blood flow when you are hypotensive and your blood pressure 5 is low, you get ischemia, which is the б 7 insufficiency of oxygen, and you subsequently experience damage to the brain, which is called 8 9 encephalopathy. 10 Q. In this particular case, do you know how prolonged the cardiorespiratory arrest was 11 12 before there was adequate resuscitative efforts given to the baby? 13 14 Α. I may have known that at some point, but I don't know it now. And the most accurate 15 answer to your question is my assumption was that 16 17 it was rather long and it was a terrible event, so that's where I left it when people gave me 18 that information. 19 Q. Again, from a general ethics 20 standpoint, when one looks at a 21 hypotensive/ischemic encephalopathy, is the 22 length of time where the resuscitative efforts 23 24 are not adequate after a cardiopulmonary arrest, the longer the length of inadequate resuscitative 25

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Fage 34 effort, does that increase the potential for more 1 2 damage to the brain? 3 I am probably not the one to ask, but Α. in my opinion --4 MR. NORCHI: Objection. 5 Q. Go right ahead, doctor. 6 7 MR. NORCHI: Go ahead. From a biochemist's point of view, the 8 Α. longer the brain is without oxygen, the more at 9 risk it is for damage. 10 The part I am having trouble 11 12 answering, just to be clear, you made the statement of inadequate efforts to resuscitate 13 and I didn't make any effort to assess whether 14 efforts were adequate or not. I just knew the 15 duration, okay? 16 17 Q. There was some suggestion in another deposition about some suggestion of a dysmorphic 18 finding in the University Hospital records. 19 Did you see any suggestion that there was any 20 dysmorphic characteristics or dysmorphic 21 22 findings? I haven't reviewed that for a long 23 Α. time, and my recollection is that it was not the 24 feeling that this child had significant 25

Page 35 1 dysmorphism. 2 Q. And to the extent that there is some reference --3 4 Α. Actually, I am slower than you. Q, 5 I'm going to slow down. As a rule, what the pathologist does 6 Α. on the first page, they would summarize that, and 7 if there was a suggestion prior to death that 8 there was significant dysmorphism, that would 9 generally be included in the clinical diagnosis 10 or summary statement. I don't see that. 11 Τ']] 12 look carefully, but I don't see a statement to that effect. 13 Q, If there was significant dysmorphic 14 features, would that cause you as a medical 15 geneticist to look to other metabolic or genetic 16 17 disorders as an explanation? Depending on the nature of the 18 Α. dysmorphism, yes. 19 Q. But absent any such evidence in this 20 case, there is no reason to have looked to that 21 22 level; correct? I can't remember what was done, 23 Α. whether the chromosome analysis was done, for 24 example, or something else. My feeling is that 25

Page 36 the neonatal intensive care unit with their 1 consultants would have looked for such things. 2 3 I would not have done any more than 4 that unless the pathologist had told me after 5 summarizing everything that there was still a question about that. Is that an answer? 6 7 Ο. Not only is it an answer, but it's 8 responsive too. 9 I want to jump back for a second, 10 because I am looking at my notes. You said that 11 you are a clinical geneticist? 12 Α. T am. That's my board. Ο. 13 There are medical geneticists that are 14 not clinically based? I am trying to think. The American 15 Α. Board of Medical Genetics first started 16 17 certifying people the year I graduated from my fellowship, so I took my boards in 1981 and 18 19 that's when I took clinical genetics. I thought T was done. 20 21 There are actually five subboards on that board. My recent boss wanted me to take 22 another subboard, so I sat in '96 for the 23 subboard of clinical biochemical genetics. 24 25 So that the direct answer to your
Page 37 1 question, there are people who are certified by the American Board of Medical Genetics who are 2 not physicians, in fact, and are boarded in other 3 activities. 4 Q. Going back to the autopsy now -- thank 5 you for clarifying that. 6 7 In the autopsy, there is a reference, page five, about halfway down in the first 8 9 paragraph of the page, there was no evidence of structural congenital anomalies in the central 10 nervous system to explain the infant's course. 11 12 Α. Yes. Q. 13 Do you see that? 14 Α. Yes. Q. What type of structural congenital 15 anomalies in the central nervous system would 16 17 typically be looked at to explain the course that 18 Megan experienced? I can answer this only from a 19 Α. 20 geneticist's perspective, not a pathologist's. From a geneticist's perspective, if a child stops 21 22 breathing, you want to look at the portion of the brain that controls breathing, and so they would 23 look in the mid brain, for example. 24 25 Similarly, they would look for

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Page 38 evidence of increased pressure, hydrocephalus, 1 2 things like that. Q. These would be things --3 4 Α. They would look for a whole range of 5 things. Q. They would be looking for things that 6 essentially would be incompatible with life? 7 Not everything they would find would 8 Α. 9 be incompatible with life. Q. 10 Were there any findings, either from a genetic standpoint, or from a neuropathology 11 12 standpoint, as you understand them, to suggest that Megan had some characteristic that made her 13 14 incompatible with life? 15 MS. MALNAR: Objection. Go ahead. From this sentence alone, it would Α. 16 seem that Dr. Dahms didn't see any such thing. 17 Ι don't know where the neuropathology report is. 18 That's different. It's done by a different 19 pathologist, and they see things that Dr. Dahms 20 21 would not see. They usually have an addition. Ι don't know where that is. 22 23 Q, There is a neuropathology diagnosis on 24 page two. 25 Α. Okay.

Page 39 1 Q. Are there any findings that would 2 suggest or be consistent with some state that would cause Megan to be incompatible with life? 3 I think here I have to defer. 4 Α. Ο. 5 Fair enough. What is it that causes 6 you the inability? 7 Α. I'm not smart enough. I'm not trying to be cute. I just don't know enough about this. 8 Q. 9 The electron transport chain enzymes that were done at University Hospitals by the 10 11 CIDEM unit --12 Α. Would you like to know what that stands for? 13 Q. I will in a moment. 14 MS. MALNAR: I would like to know 15 where you are. 16 MR. MISHKIND: Page five, second last 17 paragraph. 18 Thank you. MS. MALNAR: 19 Ο. 20 Are the results of this CIDEM unit, 21 are they part of the documents that we have in Exhibit 1? 22 Α. 23 Yes. Q. And were they normal? 24 No. 25 Α.

Page 40 Q. 1 Of what significance was the 2 abnormality? 3 Α. I can't explain the significance. 4 They formed a pattern that I could not interpret. Q. Of what significance would this 5 6 pattern have had, if any, on Megan, had she not experienced the cardiorespiratory arrest on day 7 one of her life? 8 MR. NORCHI: Objection. Go ahead, 9 10 doctor. 11 I can't answer the question as Α. My difficulty is if I can't interpret 12 phrased. 13 the significance of the results, I can't tell you either biochemically -- I can't turn around and 14 tell you the clinical significance to judge the 15 past or what might have happened in the future. 16 Q. **So** in other words, even though there 17 is something abnormal in the CIDEM unit, the 18 electron transport chain enzyme test, in terms of 19 20 how that might have or would have manifested itself in terms of disability, if any, to Megan, 21 you can't provide any type of an opinion on that; 22 23 is that correct? 24 Α. Yes. Q. And let me ask you this, just to try 25

Page 41 to save some time. 1 2 I understand that you have indicated to me thus far that all of the tests that were 3 done, the metabolic and genetic tests that were 4 done, did not permit you to conclude more likely 5 6 than not that any of the conditions that you were 7 looking for were likely the cause of her cardiopulmonary arrest; correct? 8 9 Α. I'm sorry, I wouldn't phrase it that I couldn't find a diagnosis, that's what I 10 way. 11 was doing. I was looking for a diagnosis. Q. 12 From a metabolic and genetic standpoint? 13 14 I couldn't find a diagnosis. Α. So in other words, in lay terms, there 15 Q., 16 is nothing in the metabolic or genetic tests that explained why she experienced a cardiopulmonary 17 arrest on the first day of her life? 18 19 MR. NORCHI: I'm going to object to 20 what explained means. It doesn't take into 21 content the probability, which is what you were 22 looking at before, and possibilities, but go ahead and answer the question. 23 24 If I'm not sure what the answer was, I Α. 25 couldn't then turn around when I was speaking to

Page 42 1 Mrs. Kaschak and say I know what happened. Ι 2 just couldn't do that. And that's my answer to 3 your question. Q. And basically, what you wanted to be 4 able to say to Mrs. Kaschak is that I found a 5 6 metabolic or a genetic condition -- strike that. 7 There are several things that you wanted to arrive at. One was being able to say 8 to her, I found a metabolic or genetic 9 10 abnormality that explains more likely than not 11 why she experienced a cardiopulmonary arrest; 12 correct? 13 I probably would have phrased it why Α. she died, but the answer is yes. 14 15 Q. And you weren't able to, after the exhaustive testing that was done, you were not 16 able to arrive at a test result that from a 17 causal relationship standpoint explained why she 18 experienced the cardiopulmonary arrest? 19 MS. MALNAR: Objection. 20 Α. 21 Correct. **a** . Now, had Megan not experienced the 22 cardiopulmonary arrest on day one of her life, is 23 24 there anything from any of the test results that you have that I haven't studied or that I have 25

Page 43 studied that would portend a poor prognosis or 1 2 some adverse consequences in terms of Megan's development as she went through her infancy and 3 her childhood and the rest of her life? 4 MR. NORCHI: Objection. We have 5 already been over this, but go ahead. 6 7 Α. I didn't find a consistent pattern that would have allowed me to make any 8 predictions. 9 Ο. So is there anything in any of the 1.0test results that would cause you to say that she 11 12 more likely than not would have had neurological or motor deficits or cognitive deficits as a 13 consequence of any of these test results? 14 As the tests stand, the answer is no. Α. 15 Fair enough. Q. 16 I'm curious more than anything else, 17 so I will ask it. There is a saying about 18 curiosity. 19 My office had requested back into 1998 2021 that the results of these tests performed 22relative to the autopsy on Megan Kaschak be provided. I have a letter to you back in 23 February of '98 and June of '98, and those 24 results never were provided to my office, and 25

Page 44 certainly they were not provided to the Kaschaks 1 either. 2 3 Is there a reason why that information wasn't provided? And I am not criticizing, I am 4 5 just trying to understand why. One, in my mind, I had never seen 6 Α. 7 Megan and I had never seen her parents, so that 8 my view is she wasn't my patient and I actually had no records that were unique to me. 9 And my thought was all the records 10 11 that were available were either available through Dr. Walsh-Suyks, who would be involved in this 12 13 and could send you the whole package, or the pathology department, which is actually 14 responsible, and not me, for postmortem studies. 15 I have actually gotten in trouble with 16 the pathology department because I tended to do 17 18 things in the past independent of them, and they 19 pointed out that the law requires that they deliver all that themselves, so I thought you had 20 21 it. 22 I'm actually surprised you didn't. None of this stuff was sent to me uniquely. 23 Ι was always given copies of things and I just 24 thought, I actually thought you had it because I 25

Page 45 1 had nothing unique. Q. 2 I am going to show you a letter that I have marked as Plaintiff's Exhibit 3. I will 3 show it to your counsel first and then I just 4 have one question for you on this point. 5 6 (Thereupon, Plaintiff's Deposition 7 Exhibit 3 was marked for 8 purposes of identification.) 9 10 Q. First, is Plaintiff's Exhibit 3 a copy 11 of a letter sent by my office to you back in June 12 13 of '98? 14 Okay. Α. Q. 15 Do you recall receiving that letter? 16 Α. No. Q. You don't. Are you saying that you 17 didn't or you just don't remember one way or 18 another? 19 20 Α. I am saying I don't recall. Q. 21 And the only reason **I** am taking the 22 time on that is that this letter was pointing out 23 to you that we had tried to get the records and were told and were directed back basically to 24 you. We requested the records from the hospital 25

Page 46 and they indicated that there were no such 1 2 results. 3 Do you ever recall having conversations with my office where we were saying 4 that we couldn't get these records from anywhere 5 else and we needed your assistance in terms of 6 7 providing them to us? I don't recall the conversation 8 Α. specifically. I truly thought all these records 9 would be sent by pathology and sometimes, I mean, 10 I will tell you probably on my end, it takes 11 forever to get things, and **I** just figured it was 12 taking forever. But I was told to have all 13 records go through the hospital and that's what I 14 tried to do. 15 Q. Who was it that advised you **of** that? 16 17 Α. I mean, it's no one in particular. The medical records department has asked people, 18 various administrators have asked people, I am 19 sure the lawyers have asked people to ask us to 20 do this. 21 Ο. When is the last time you had an 22 occasion to talk to Dr. Walsh-Sukys about this 23 particular baby? 24 I can't tell you in detail when that 25 Α.

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1	might be. But I have not talked about the
2	details of the testing for some time.
3	Q. To your knowledge, has Dr. Walsh-Sukys
4	arrived at or expressed to you any opinions as to
5	the cause of the cardiopulmonary arrest that she
6	had shared with you?
7	MS. MALNAR: Objection. Go ahead.
8	A. I don't think so. The longer answer
9	is that since I was uninvolved in Megan's care, ${\tt I}$
10	think one of the reasons I was asked to do this
11	is I had no prior knowledge of events, and so I
12	didn't want to hear about anything else.
13	I just wanted to do what they had
14	asked me to do. I wanted to do the lab tests
15	without prejudice from anything else that had
16	gone on clinically. So if someone had a
17	conversation, I would've said, okay, <i>go</i> back and
18	do what ${\tt I}$ have to do and that's what ${\tt I}$ would have
19	done.
20	Q. You clearly did not want to know what
21	the precipitant was or how responsive the
22	resuscitative efforts were or were not following
23	the cardiopulmonary arrest?
24	A. I mean, the story I heard was that
25	MR. NORCHI: Objection. Go ahead.

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Page 48 1 0. Go ahead. Α. -- was that Megan had had a bad time 2 of it and it was a long time. 3 Ο. And who shared that with you? 4 Α. Michelle probably did and I think 5 Dr. Dahms when she went over the clinical story. 6 Anyone else that you recall besides 7 0. Dr. Dahms and Michelle that shared with you that 8 9 history? 10 Α. Not directly with me. 11 How about indirectly? 0. 12 I mean, I am trying to think if in the Α. course of things people said something, but no 13 one had a conversation with me about it. 14 15 Q. Is there something in general that you are referring to? 16 There are a lot of people in 17 Α. No, no. the hospital. People talk. I am just trying to 18 say no one spoke to me specifically about Megan's 19 20 case that **I** stayed long enough to hear the end of 21 it. Ο. I take it from that -- and if I am 22 wrong, set me straight -- that you overheard 23 excerpts of some conversations? 24 25 Α. No, I am saying I may have. I just

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1	know this was a case that there was some
2	discussion about and people would have talked,.
3	And I know, I might have been around the unit or
4	talking to Michelle and someone else said
5	something. I just don't remember details. But
6	my information, which I think I am not sure
7	what your question pertinence is but to me the
8	information was given when Dr. Walsh-Suyks asked
9	me to get involved or talking to Dr. Dahms to try
10	to help out there.
11	Q. Did you ever talk with Dr. Bruce Cohn
12	from The Cleveland Clinic?
13	A. No.
14	Q. Did you know that he was brought over
15	to do a neurological evaluation?
16	A. I specifically knew that, yes. I have
17	spoken with Dr. Cohn on numerous occasions, but I
18	don't think I have ever spoken about any of
19	details of Megan's care.
20	Q. When you say you have spoken to him on
21	numerous occasions, did it have anything to do
22	with any aspect of this case?
23	A. What I am saying is, Dr. Cohn used to
24	come to meetings where I was at and he might
25	we might have said we saw this or that patient in

Page 50 the course of things, but I did not go over the 1 details of this with Dr. Cohn. 2 One of the potential causes of a Ο. 3 cardiopulmonary arrest was obviously potential 4 inborn errors of metabolism; correct? 5 6 Α. Start again. One of the potential causes of Q. 7 cardiopulmonary arrest was the possibility of 8 inborn error of metabolism; correct? 9 10 Α. Correct. Q. And such things as galactosemia and 11 phenylalanine as the other classic inborn errors 12 of metabolism were looked at and ruled out; 13 14 correct? MR. NORCHI: Objection. If you know. 15 MS. MALNAR: If you know, you can 16 17 answer. My assumption was that the state 18 Α. screening looked at those. On the other hand, I 19 would not have thought to look again or more 20 21 carefully at those possibilities, because they do not produce the kind of problems that Megan had. 22 Q. Okay, fair enough. 23 I'm going to quote as best as I can 24 what I'm told as it relates to a conversation 25

Page 51 1 that you had in October of '98, probably one of 2 the last conversations that you might have had with Mrs. Kaschak. And I'll give it to you 3 4 entirely and then I'm going to ask you whether or not you remember that conversation or something 5 similar to that taking place. And if not, tell 6 7 me why. Okay? 8 Α. Okay. Q. You understand the context under which 9 T'm about to --10 11 Α. I'm not sure. Ο. 12 In October of '98, I am advised by Mrs. Kaschak that you had a conversation with her 13 and told her that all of the genetic tests done 14 on Megan were normal, except for an inconclusive 15 borderline electron transport chain assay test 16 17 which Mrs. Kaschak told me you told her could be many things or nothing at all. 18 Does that sound consistent with what 19 you likely discussed with her? 20 21 Α. I discussed that kind of thing with Mrs. Kaschak. I'm not sure I would've said it 22 that way, though, but I discussed those kinds of 23 things with Mrs. Kaschak. 24 Q. When one says that the borderline 25

Page 52 1 electron transport chain assay test -- which 2 could be many things or nothing at all -- in simple terms that a lawyer or a jury can 3 understand, what does that mean? 4 Α. Again, I'm not sure that that's what I 5 said. Second, what the results show is an 6 7 inconclusive pattern of abnormalities. Q. 8 Can you be any more specific as to what the pattern of abnormalities are? 9 10 Α. I can be as specific as you want if you want to go over the labs, but if you don't 11 12 want me to do that, I can tell you that the results from different tissues did not agree 13 within themselves, so that an abnormality that 14 was found in one tissue was not normal, and was 15 normal, for example, in another, and in one case 16 17 was actually increased; in other words, certainly not deficient. 18 Ο. Have you ever seen that kind of a 19 20 pattern before? 21 Α. Yes. 22 Q. And what does it mean to you as a geneticist where you have these inconclusive or 23 inconsistent findings? 24 It means I can't interpret the results 25 Α.

Page 53 in a consistent and helpful way. And I think 1 2 that was the gist of my conversation with Mrs. 3 Kaschak. Q. Were there any additional tests after 4 the electron transport chain assay test in 5 6 October of '98 that you indicated to Mrs. Kaschak 7 were being done that you were going to report back to her or that she would learn about? 8 9 I don't remember the date of our last Α. conversation, but in our last conversation, I 10 11 think I said that there was no more testing I had planned. 12 Ο. And the CIDEM unit stands for what? 13 14 The Center for Inherited Disorders of Α. Energy Metabolism. 15 16 Ο. That's at UH? 17 Α. You know, I actually don't know where under house auspices. It's physically located in 18 19 Rainbow and at the V.A. Beside the V.A. and Rainbow, were Ο. 20 21 there tests that were sent out of state? As I said before, there was one 22 Α. Yes. test that was sent to Los Angeles and some tests 23 sent to another building for the genetic tests. 24 25 Q. The ones to Los Angeles, was that sent

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Page 54 1 to Dr. Ng's office? 2 Α. Dr. Ng? Ο. 3 Yes 4 I don't see the name on it. I think Α. 5 Dr. Ng is the one that actually did them. They 6 were signed by somebody else. She subsequently 7 left Los Angeles, so I can't remember. 8 (Thereupon, a discussion was had off 9 the record.) 10 0. What is an oxidative metabolic energy metabolism defect? 11 12 It's a defect where you're burning up Α. 13 some food product with oxygen to get energy from 14 it. Ο, 15 Was there any evidence that Megan had that condition? 16 No conclusive evidence. 17 Α. Q. 18 The glycogen deposition found on 19 autopsy raised an issue of glycogen storage 20 disease; correct? 21 It raised that possibility. Α. 22 Ο. But that ultimately was not established? 23 24 I thought it was more likely since we Α. couldn't fit it into a recognized pattern of one 25

Page 55 1 of the glycogen diseases to think that it was related to an oxidative metabolic disorder for 2 which there was precedent. 3 4 Ο. And what causes such a condition? 5 Α. I think we discussed that before, but 6 basically if the cell can't mobilize energy, it can't use glycogen effectively. 7 Q, 8 But what can cause the inability of 9 the cell to mobilize or to metabolize? 10 If there is a defect in mitochondrial Α. function where the mitochondria is the carburetor 11 12 of the cells. It takes different cells and fuels 13 and burns them. 14 If the mitochondria cannot do that, various fuels will accumulate because it has no 15 16 place to go, so you can get deposits of glycogen. 17 Q. Are there any test results that establish a mitochondrial defect? 18 19 Α. No. Ο. 20 Can you state to a probability then 21 that there was some type of a glycogen storage 22 disease? T found no evidence in the biochemist 23 Α. school studies directly confirming that --24 Q. 25 So this is just --

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Page 5 1 Α. -- or disproving that. Q. 2 -- just a possibility, but not a probability? 3 4 Α. Correct. Ο. There is some reference in the autopsy 5 to Megan's death falling into the category of a 6 near missed SIDS. 7 Are you qualified to provide any 8 opinions on whether or not her age and the 9 circumstances surrounding her cardiopulmonary 10 11 arrest and her death fit the stigmata or the description of a near SIDS death? 12 13 I'm not an expert on SIDS. But I can Α. 14 answer the question to the extent that SIDS-like episodes mean that they don't necessarily fit 15 into the standard criteria and a subset of those 16 can have an underlying metabolic cause. 17 Q. But again, we don't have anything that 18 you could point to that establishes an underlying 19 metabolic cause in this case? 20 Correct. 21 Α. Q. There were muscle biopsies done; 22 23 correct? 24 Α. Postmortem? Q. Yes. 25

Page 57 1 I wouldn't call it a biopsy then. Α. Ι 2 would say muscle samples were obtained 3 postmortem. Q. 4 Were there any findings that were significant in terms of leading to any definitive 5 conclusions? 6 The pathology report would answer 7 Α. that. I don't remember any mention that they 8 found such. 9 Q. The tests that you were performing 10 wouldn't have any implication on any --11 12 Α. No, if you were asking me morphologic changes that the pathologist pursues. 13 I am answering the question, some of the reports that 14 I provided that you now have in your possession 15 are done on skeletal muscle and that's included 16 17 in the comments we have made before. Q. Any abnormalities on the skeletal 18 muscles that you describe? 19 The skeletal muscle does have 20 Α. Yes. abnormalities in some of the electron transport 21 chain. 22 Q. And of what significance are those 23 abnormalities? 24 25 I cannot tell you the significance Α.

Page 58 because they are not consistent within different 1 tissues, so by sharing the results in the 2 skeletal muscle with the heart muscle with the 3 liver, it's the whole picture that I can't put 4 together. 5 Q. Did you take into account in trying to 6 7 put these findings together and arrive at some explanation for her death the fact that she had 8 good apgars at the time of birth? In other 9 words, the clinical picture during the first 24 10 hours, did you take any of that into account? 11 12 I took it into account, yes. Α. Ο. And in what respect did that help or 13 hinder your conclusions? 14 15 Α. It didn't help. Q. Why is that? 16 Because it's not helpful. You often 17 Α. have children who have perfectly fine apgar 18 scores, who, until they have a requirement for 19 20 energy production, do not get into trouble. Q. And those children that have perfect 21 apgar scores that then have metabolic studies or 22 genetic studies done, more often than not, you 23 are able to arrive at some explanation from a 24 metabolic or genetic standpoint? 25

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Page 59 1 More often, I am not able to arrive at Α. 2 an answer. **So** then you ultimately wind up saying, 3 Q. I have no explanation? 4 You haven't asked me, but -- I 5 Α. Yes. know I'm not supposed to do this -- but the apgar 6 scores, you can have normal appar scores, 7 abnormal apgar scores. They are not a singular 8 useful thing and often I find that I cannot 9 10 provide an explanation independent of the initial 11 apgar scores. Ο. This is a search for the truth, so if 12 you happen to blurt out something --13 I am not blurting, I am trying to 14 Α. 15 help. Q. I appreciate that. 16 The existence of severe 17 hypotensive/ischemic encephalopathy that's found 18 on autopsy, is that in any way explained by any 19 of the results from a metabolic or genetic 20 21 standpoint? 22 Α. No. Q. Are you aware of any studies, 23 articles, that are contained within the medical 24 literature that are similar in any way to Megan's 25

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1 case where there were concerns about genetic or 2 metabolic abnormalities being the explanation for 3 a cardiopulmonary arrest, where all of the tests 4 were done and basically they were inconclusive 5 from a genetic standpoint, a series of tests or 6 articles?

A. It's well recognized that you don't
always find the answers. It's in the literature
in various forms.

10 Q. Where would you refer someone for, 11 perhaps, something in the 1990s, if you will, 12 that talks about the various metabolic tests that 13 are performed and the conclusive or inconclusive 14 nature of the tests as it relates to trying to 15 come up with an explanation for why such an event 16 would occur?

17 A. I don't know the specific reference to18 refer you to.

But having said that, none of this Q. 19 explains the nature of the -- none of the 20 21 inconclusive test results explain why Megan had the ischemic encephalopathy; correct? 22 23 Α. I have already stated, no. Q. That's a separate clinical issue which 24 is not explained away or by what you did in this 25

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Page 61 1 case? I don't know if it's separate or not. 2 Α. I don't know what that means. 3 There is nothing that you have come up 4 Ο. with that indicated why there was such a profound 5 ischemic encephalopathy in this case; correct? 6 7 Α. Correct. MR. MISHKIND: Let's take a couple 8 minute break 9 (Thereupon, a recess was taken.) 10 Q. Doctor, **I** just have one other question 11 12 for you and then **I** am done. There was some testing done of one or 13 14 both of the parents, I believe, after Megan died to try to determine something. 15 My question to you is -- and I believe 16 17 it was requested by Dr. Robin -- are you aware of that? 18 Α. No, I'm not actually. 19 Ο. I lied. Actually have one more 20 21 question. Do you know a reason that blood tests 22 would be done on the parents in this context? 23 MS. MALNAR: Objection. Go ahead. 24 25 Α. Not in the context that I was

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Page 62 considering. I don't know what Dr. Robin was 1 2 thinking. MR. MISHKIND: I have no further 3 questions. 4 5 Α. If you tell me the tests, I will tell 6 you what it's indicated for. Ο. I would if I knew, but I can't, so I 7 won't. 8 MR. MISHKIND: I have no further 9 10 questions. EXAMINATION OF ARTHUR B. ZINN, M.D., Ph.D. 11 BY MR. NORCHI: 12 Ο. I have some, doctor. I represent the 13 defendant in this case. Dr. Zinn, my name is 14 15 Kevin Norchi. You mentioned earlier that you were 16 given a list of tests to perform or have 17 performed; is that correct? 18 I was given a list of differential 19 Α. 20 diagnoses -- not given a list. In reviewing the chart, there was a list prepared by the 21 neonatologist as well as one in Dr. Cohn's 22 23 consultation. Ο. So there was a differential diagnosis 24 in the discharge summary from the chart at 25

Page 63 1 Rainbow Baby and Children's Hospital. Is that what you are talking about? 2 Α. No. I don't know. I wasn't given a 3 4 list. In the discussions either with Dr. Walsh-Sukys or Dr. Dahms in going over 5 things, I reviewed Dr. Cohn's set of 6 7 recommendations, as well as any that they had advised me that their own people had considered. 8 Q. 9 It wasn't clear from your earlier testimony. It sounds as though somebody gave you 10 a list of potential diagnoses. 11 12 Α. It may have felt that way, but it wasn't actually being given a list. 13 I generated 14 a list, I suppose. Q. Do you have a copy of that list you 15 qenerated? 16 17 Α. No. Q. Did you actually write out or type out 18 or have printed out this list? 19 I didn't. 20 Α. 21 Q. Did anybody? The list of tests are the ones that Α. 22 are listed through Dr. Dahms' note. That's what 23 that list was generated to use for. So in her 24 summary, she goes through the various 25

Page 64 considerations that were made prior to and after 1 death 2 Q. Were you involved in discussions with 3 Dr. Dahms when she prepared that list or did you 4 get involved after she prepared her list? 5 6 Α. Dr. Dahms got involved after the 7 death, so at some point Dr. Dahms and I spoke and generated a list together. 8 Q, 9 So that's a list you and Dr. Dahms 10 generated? 11 Α. As I indicated before, I had discussion with Dr. Dahms. 12 Ο. I know that. I am wondering where the 13 list came from; that's all. 14 From our discussions of our 15 Α. considerations before death and what the autopsy 16 might have suggested to amend that list. 17 And then your responsibility after Q. 18 receiving this list was to coordinate certain 19 20 genetic and metabolic tests? 21 Α. I didn't receive it, I generated the 22 list. 23 Q. After you and --24 See, the list that was in the chart Α. was a list of differential diagnostic 25

Page 65 1 considerations. The reason they wanted my help, 2 they wanted to know how to translate that efficiently and effectively into a list of tests 3 that could answer whether any of these 4 possibilities are the case or not. 5 6 So Dr. Dahms is a fine pathologist, 7 but she needs help. In actually saying if you are going to look for pyruvate dehydrogenase 8 deficiency, then what tissues do you need and 9 where do you send it. And so that's what we 10 generated together. 11 Q. 12 So if I understand you correctly, Dr. Dahms provided the identity of particular 13 deficiencies that she thought might possibly 14 exist and then your involvement was to identify 15 the tests? 16 We review the records together, if you 17 Α. will. 18 Q. Well, you can clarify. I appreciate 19 your help. But then you identify tests that 20 could be done to identify whether or not each of 21 those or any of those conditions existed? 22 23 Α. Correct. 24 Q. And then from there, then it was your 25 obligation to coordinate these tests?

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1	A. To have pathology send the appropriate
2	samples to the appropriate laboratory. I never
3	physically carried samples and ${\tt I}$ never have
4	responsibility for performing tests.
5	Q. But it was your job, if you will, your
6	obligation, to direct certain specimens to
7	certain pathologists or laboratories?
8	A. Correct.
9	Q. And as you mentioned, you did not
10	perform any of these tests yourself; correct?
11	A. Correct.
12	Q. The test results came back to you,
13	though; is that correct?
14	A. Copies came back to me.
15	Q. Was the purpose of you receiving those
16	copies of those test results to interpret the
17	test results?
18	A. No. The interpretations were provided
19	on the reports.
20	The other reason ${\tt I}$ was asked to get
21	involved, and the reason ${\tt I}$ actually agreed to get
22	involved was that someone had to communicate with
23	the Kaschaks. And since ${\tt I}$ had been previously
24	uninvolved, part of my neutrality, people
25	thought, would be useful to communicate with the

Page 67 1 Kaschaks. Q. 2 Okay. Α. So what I needed to do was, my hope 3 was that when everything was done, I would sit 4 and meet with the Kaschaks and help them deal 5 with and understand the information and use it 6 7 appropriately. You testified earlier that there were Q, а certain inconsistent test results; correct? 9 10 Α. Correct. Q. Was it your responsibility in this 11 setting to look at all the test results and 12 13 determine whether they were consistent or inconsistent? 14 That was part of my responsibility. Ι 15 Α. was ultimately going to sit down with the 16 Kaschaks and do that, correct. 17 Did anybody else, any other physician, 0 -18 have the responsibility of looking at all the 19 test results, determining whether they were 20 consistent or inconsistent with a particular 21 22 diagnosis? 23 Α. None of the physicians who had cared for Megan had that responsibility. Michele 24 Walsh-Suyks had asked me to do that for her, 25

Page 68 1 thinking I was more qualified, so I would talk with her about the results in that context. 2 And I know, I think on at least one 3 occasion, she got back with the Kaschaks and 4 tried to communicate what I would say to her, 5 because she knew them already. 6 Ο. So if I understand, the test results 7 8 came back to you and because of your 9 qualifications as a medical geneticist, you 10 interpreted those test results, arrived at some conclusion or inconclusion, as it were? 11 Right. 12 Α. Ο. And communicated that to the Kaschaks 13 and to Dr. Walsh-Suyks; correct? 14 Α. Correct. I also spoke with the 15 laboratory directors if I had questions about 16 interpretation, and on at least, I think, a 17 18 couple of occasions where there was ambiguities, the laboratory actually redid certain tests and 19 tried to refine their assays, so I had 20 21 communication with the laboratory. Q. So you did speak with the lab 22 23 directors or physicians at the labs and perform 24 these tests? The lab directors, yes. 25 Α.

Page 69 1 0. Did you keep any notes regarding your 2 discussions with any of those individuals? 3 Α No. Q. 4 Do you have any notes regarding any 5 discussions you had with the Kaschak family or Dr. Walsh-Sukys or anybody? 6 7 Α. Probably not. Q. Why do you say probably not? 8 9 Because the computer systems have been Α. 10 changed and I don't know if I wrote anything on the old computer. As a habit, I generally 11 12 didn't. I would generally wait until I had met with the family and summarize our discussions 13 14 that way. 15 Q. You mentioned earlier that there were abnormalities found on some of the test reports; 16 17 correct? 18 Α. Yes. Ο. I would like to ask you some questions 19 20 about them. On the first page -- I believe it's 21 Exhibit 1 that was marked -- there is an electron 22 transport chain report for specimen type skeletal 23 muscle? 24 25 Α. Correct.

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Page 70 1 Ο. And that's an ETC assay dated September 29, 1997? 2 3 Α. Yes. Q. Just to make sure we have the same 4 5 page. 6 Α. Yes. 7 Q, There are abnormalities noted in the interpretation below, the bottom part; correct? 8 9 There are abnormalities noted in words А below, and to the right of the last column there 10 are H's and L's, which means H is high and L is 11 So that's the first designation of an 12 low. abnormality. 13 Ο. 14 What is a complex I, complex II and complex III? 15 16 If that requires an hour long 17 dissertation, I will probably abandon the 18 question. 19 If you can imagine that the Α. 20 mitochondria is like a carburetor and that it mixes the fuel, and the oxygen that you 2 1 contribute, the mitochondria's job is to burn 22 that and make energy. There is different parts 23 24 in the mitochondria -- you **call** those parts 25 complexes -- so there that laboratory measures

	Page 71
1	four different parts of the carburetor.
2	Q. On the second page of Exhibit 1, do
3	you see under interpretation by Dr. Kerr, next to
4	that is a date, October 14, 1997, he identifies
5	that in the assay of the electron transport
6	chain, the postmortem skeletal muscle showed a
7	defect in the later components of complex II;
8	correct?
9	A. You are reading correctly.
10	Q. First of all, if we can, is it
11	possible to explain for me what a later component
12	of complex 11 is?
13	A. Sure. If you think about the
14	carburetor having four major parts, part II, if
15	you take it out, you can sort of, it has four
16	subpart components to it. So the first two parts
17	are the early part and the last two parts are the
18	late part.
19	Q. What is the effect of those later
20	parts in the oxidative phosphorylation mechanism
21	or OxPhos?
22	A. Let me go back to the parts. If you
23	look above the line on the first page, there are
24	A. So complex II_{ℓ} its job is to take the
25	energy intrinsic to a substance called succinate

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and get the energy out of it. So the first two 1 2 components of that complex are together called succinate dehydrogenase and those sort of take 3 succinate and gives the order over to cytochrome 4 5 с. The second two parts in turn take the 6 7 energy from the components and give it to cytochrome c, so the way these things are named 8 9 is complex II is a bridge between the energy and succinate and the energy that will be transferred 10 to cytochrome c. So that's it. 11 Ο. 12 Are there any particular syndromes that you are familiar with that are related to a 13 defect in the later components of complex II? 14 There have been reports of patients Α. 15 who have defects in complex II. They have a 16 range of medical problems. 17 Q. Have you ever done any research in 18 19 that particular area; that is, specifically focused on defects in the later components of 20 21 complex II? 22 Α. Have I, personally? Q. Yes, sir. 23

24 A. No.

25 Q. Is this a relatively new area of
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Page 73 inquiry in the field of medical genetics focusing 1 2 on defects in a particular portion of a complex? 3 Α. Yes. 4 Ο. Would it be fair to assume that this 5 is an evolving area of medical genetics? 6 Α. Yes. 7 Ο. I will assume most areas of medical 8 genetics are evolving? 9 Α. Many are. Let me go back to two 10 questions ago. 11 Q. That's fine. 12 Α. I have not done research, but I have 13 done all these assays. I worked in Dr. Hoppel's lab for several years and I know what they are. 14 I often interpret these results for other 15 patients and I have some idea what electron 16 17 transport chain is about. I am defining research narrowly. 18 19 Ο. I didn't mean to suggest that you don't understand the electron transport chain. 20 21 Α. But you asked a specific question. Q. 22 It was specific. 23 And I am trying to explain what I mean Α. 24 by research. 25 Q., I think we understood each other.

Page ?4 You had an opportunity to review these 1 reports today before your deposition testimony? 2 Yes. 3 Α. Q. The defect in the late component of 4 complex II is one of the defects identified in 5 these reports; correct? 6 It's one of the laboratory Α. 7 8 abnormalities identified in these reports. Q. 9 Are there any other lab abnormalities that are identified by these reports? 10 11 Α. I can turn the pages with you. Q, 12 Could you, please? 13 So the first page refers to the Α. 14 analysis on skeletal muscle done on October lst, 15 '97. At that point, there are actually two 16 abnormalities. One is the assay that measures 17 complexes I and III together was actually, let's 18 see -- the one that measures I and III together 19 was higher than the range. Whereas the assay 20 that measures II and III together was lower than 21 the range. 22 Q. 23 In and of themselves, do those abnormalities direct you towards any particular 24 diagnosis or syndrome? 25

Page 75 Not a particular syndrome. 1 Α. The 2 question is does this patient, does this girl have complex II deficiency. That would be the 3 question. 4 Q., And, of course, this report doesn't 5 give you a definitive answer to that question, 6 though, correct? 7 8 Α. Correct. Q. But certainly leads you towards 9 inquiry in that direction? 10 11 Α. Correct. Ο. 12 Are there any other --Not on that page. On the next page is 13 Α. Dr. Kerr's interpretation, his analysis of three 14 different -- of two different enzymes and we will 15 just do initials. PDC and dihydrolipoamide 16 dehydrogenase, he found those both to be normal. 17 The next -- those are enzymes that 18 live in the mitochondria that are primarily 19 20 active in the liver as opposed to other tissues, but they are both present in other tissues. 21 On the next page, Dr. Kerr also 22 measured in the liver two other enzymes, pyruvate 23 carboxylase and phosphoenolpyruvate carboxykinase 24 and he found those to be normal, as well. 25

	Page 76					
1	So that is, all the pyruvate related					
2	enzymes live in the mitochondria, but they are					
3	not directly a part of the oxidative phosphor					
4	system. They feed energy into it.					
5	Q. So there may be an abnormality in the					
6	OxPhos mechanism?					
7	A. But you wouldn't say it by assaying					
8	these enzymes.					
9	Q. Could you see an abnormality in the					
10	OxPhos mechanism by assaying skeletal muscle?					
11	A. That was done. That's what the first					
12	page is doing.					
13	Q. And that's looking for an abnormality					
14	in the actual OxPhos mechanism?					
15	A. That's correct. And this was yes.					
16	On the next page what happens and					
17	these next four, I think, reports are from 4th of					
18	November of '97.					
19	Q. Charles Hoppel, Dr. Hoppel?					
20	A. Yes. And these were done after he					
21	actually redesigned some of the analyses. So					
22	they represent remeasurements with what Dr.					
23	Hoppel considered a more updated set of analyses					
24	for the same complexes that we discussed reported					
25	on October 1st, okay?					

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	Fage 77
1	Q. Okay.
2	A. So he basically said that we found the
3	stuff, we are going to change some of the ways we
4	do the stuff in the lab, let me go back and do it
5	over again, as well as do the other tissues.
6	Q. Who asked him to do this again; do you
7	know?
8	A. Knowing Dr. Hoppel, he decided to
9	change A s own assays.
10	Q. Okay.
11	A. But I'm sure I had discussions with
12	him about the difference and what he was doing.
13	The results of that skeletal muscle,
14	again, the analysis for the succinate-cytochrome
15	c reductase was on the low side, just on the
16	border.
17	Q. Yes.
18	A. Whereas on this occasion, as opposed
19	to the first, the analysis for the fourth
20	components cytochrome c oxidase was below the
21	range of normal and below the prime mean for
22	controls.
23	On this occasion, though, the citrate
24	synthase, which is the last enzyme on this page,
25	which is used, an intramitrochondrial enzyme,

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1	that's not a part of the electron transport
2	chain, but is sometimes used as a reflection and
3	our lab uses it as that of the integrity of the
4	mitochondria. In other words, if you're
5	measuring it, you want to make sure what you are
6	measuring is at least, you know, healthy.
7	Q. What does the low result tell us about
8	the integrity?
9	A. It's on the board and it let's see
10	if the report comments.
11	The activity of citrate synthase, a
12	mitochondrial marker enzyme is just below the
13	control range.
14	Now, so that's Dr. Hoppel's
15	interpretation.
16	Q. Does the low number for the citrate
17	synthase suggest that we may have lost integrity
18	of the mitochondria?
19	A. That's the concern. But from reading
20	this report, Dr. Hoppel's view is borderline. He
2 1	doesn't come out one way or the other saying
22	whether it was healthy or not.
23	Q. Would that affect the reliability of
24	the test in any way?
25	A. Certainly can.

Page 79 I just don't know how complicated to 1 2 make it. Wait until he asks the MS. MALNAR: 3 4 question and then go ahead and answer that. Q. In this report of November 4, 1997, 5 6 Dr. Hoppel in the second to last sentence, he states that the activities measured on this new 7 homogenate are similar in comparison to the assay 8 9 done on September 29, 1997. 10 Α. Correct. 11 Q, Do you see the results of any assay 12 done on September 29, 1997? That's page one. When you read the 13 Α. date, to me that's the date, not his signature 14 date. 15 Q. Okay. That clears that up. 16 Do you interpret his findings to state 17 that the assay done on November 4, 1997 is 18 19 consistent with or the same as the findings on his report dated October 1, 1997? 20 21 MS. MALNAR: Objection. Go ahead. 22 I interpret just the way he said. He Α. said they are similar. They are not identical. 23 Similar? Q, 24 Okay. That's the word Dr. Hoppel uses. 25 Α.

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Page 80 The next page then is a report of the 1 2 same assays used on the liver, and here what you see is a different pattern than what you saw for 3 4 the skeletal muscle. And the pattern is, I mean -- you can look at it for yourself. 5 Q, Well, for example, the citrate 6 synthase on the liver sample is low; correct? 7 8 Α. Correct. 9 Q. Does that, again, refer to the integrity of a mitochondria that is being 10 assayed, tested? 11 Again, Dr. Hoppel's comment is that 12 Α. the activity of citrate synthase a mitochondrial 13 14 marker enzyme is below control range. Q. What does that mean? 15 That means that the marker for 16 Α. mitochondrial integrity is not within the range 17 that you would like to see. 18 Ο. 19 Does that mean that these tests are less reliable now? 20 The way I can answer that is that my 21 Α. experience with Dr. Hoppel -- and obviously you 22 need to talk to him directly if you want to have 23 24 him interpret his own sentences -- but in my experience, what that means is what Dr. Hoppel 25

Page 81 observed; that it was low, but thought that some 1 2 of the other assays on balance were interpretable and that's why he phrased it this way. 3 If Dr. Hoppel's results are such that 4 he does not think that he can do an 5 interpretation, he will state that. 6 7 Q. Doctor, you were --Α. Should I keep going through the 8 assays? 9 Q. Let me ask you about this one so we 10 don't have to keep coming back. There are some 11 abnormal results here; right? 12 Α. Right. 13 And you attempted to interpret the 14 0. results so you could communicate with the 15 Kaschaks? 16 17 Α. Right. 18 Q. What interpretation can you make of these test results relative to Megan's condition? 19 20 MS. MALNAR: Just this page? Just this page or can't that be done? 21 Q. Α. It can be done. It's just not very 22 useful, I mean, to interpret this page, I mean, 23 I can read Dr. Hoppel's report and the question 24 about the citrate synthase, sure, it makes me 25

Page 82 1 worry whether these were intact and that's why they were being used in this test and that's what 2 3 Hoppel is saying. But my experience is Hoppel is not cavalier, and if he thought he couldn't do it 4 at all, he would've said so. 5 But if you allow me, the real 6 importance is comparison of this with the other 7 results. 8 Q. 9 Okay. And for example, on the skeletal 10 Α. muscle, what you found on two occasions was a 11 succinate cytrochrome c reductase, that was 12 borderline low. On this case, you have a normal 13 activity of succinate cytrochrome c reductase; 14 even in the context in which you are not sure 15 what the integrity of that liver is. 16 So what I concluded was that there is 17 not a consistent deficiency for the components 18 assayed by this complex II, III analysis. 19 20 Therefore I couldn't say that I think there is error in complex II and 111. 21 There is always provisos. 22 The 23 provisos are that there are tissue specific defects for mitochondrial disorders. So that 24 underscores the difficulty in interpreting these 25

Page 83 1 things. 2 The cleanest picture is if you have 3 one tissue that's completely abnormal or selectively abnormal and one tissue that's 4 completely normal and then you say maybe if that 5 corresponds to the clinical course then maybe 6 what you have is a tissue specific defect, 7 because the genes and the proteins of these 8 complexes can be different in different tissues. 9 I'm sorry to go on. 10 Q. 11 No. 12 That's my interpretation here so far, Α. 13 okay? Q. 14 Okay. And if you then turn to the next page, 15 Α. you see the results in the heart, which again, 16 you can go through this, again the same proviso 17 with the citrate synthase, but you see a pattern 18 different here than you saw for the first two 19 20 tissues. So what I say is, there seems to be 21 some areas of abnormality in all the tissues. 22 Some of them are marginal and some less marginal 23 or borderline, but there is no consistency 24 25 between or amongst the tissues.

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	Page 84					
1	Q. Did you try to determine whether there					
2	was any tissue specific mitochondrial					
3	abnormality?					
4	A. Other than by trying to look at these					
5	and see if there was a pattern that became					
6	evident.					
7	Q. From just looking at the skeletal					
8	muscle well, let me ask you, is the tissue in					
9	skeletal muscle similar to tissue in other parts					
10	of the body, like heart muscle, for example?					
11	A. There are some similarities and some					
12	differences, and there are instances in which we					
13	evaluated patients and they have the same defect					
14	in both. There are other instances where they					
15	have a defect in one and not the other, depending					
16	on the particular component.					
17	I don't need to make this a lecture,					
18	but there are five different electron transport					
19	chain complexes. This lab has four of them.					
20	Within those four components there are about 70					
21	different enzymes, 70 different proteins, which					
22	means there is at least 70 different genes that					
23	do all this. And those different genes often are					
24	expressed differently in different tissues. We					
25	do not have the ability to look directly at most					
1						

Page 85 of those tissue differences. 1 Q. Do you find any similarity between the 2 heart electron transport chain report and the 3 skeletal transport chain report that permits you 4 to come to some conclusion about any 5 abnormalities in any of the ETC complexes? 6 Ruling out or ruling in, 7 MS. MALNAR: or either, in general? 8 MR. NORCHI: General. I am not asking 9 for a specific diagnosis. I understand he can't 10 do that. So it's a general. 11 I can't reach a conclusion. 12 Α. Q. And what would the conclusion be? 13 No, I can't. 14 Α. Q . You can't? 15 Α. I cannot. 16 Q. 17 I thought you said you could. The next report from December 11, 1997 18 fibroblasts, there are no abnormal results on 19 20 that page; correct? 21 Α. Correct. The comment on that page, though, which you didn't ask me, but the comment 22 23 on that page, one, you cannot measure complex I in skin fibroblast, so that's not included in 24 there. 25

	Page 86
1	Two, the purpose of looking at
2	fibroblasts is that whatever might have happened
3	with Megan during her life would not be reflected
4	in the skin fibroblast study. You take a skin
5	fibroblast, take a biopsy, grow it in tissue
6	culture under as optimal conditions as you can so
7	that it's relatively independent of the medical
8	status, because it's done so much later and
9	outside the body.
10	On the other hand, what takes away
11	from the merit to try to interpret the other
12	results is that the tissue specificity of the
13	other tissue is often least reflected in skin
14	fibroblast.
15	Q. Meaning?
16	A. So that skin is less like heart,
17	perhaps, than skeletal muscle is like heart.
18	This is not always the case. Geneticists
19	traditionally try and do fibroblast studies for
20	the reasons I just said. They are not always a
21	reflection of what is present in other tissues.
22	Q. For example, if you wanted to do a
23	tissue specific test for skeletal muscle or heart
24	or that type of tissue, doing a skin fibroblast
25	assay

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Page 87 Not if the test you are going to do is 1 Α. a functional test measuring a particular 2 If you wanted to do a DNA base test, 3 protein. 4 the genes are the same in each tissue, but how those genes make their proteins that does the 5 work is different. 6 Q. 7 Okay. That wasn't done here; That last -correct? 8 The DNA based studies I believe. 9 Α. Q. There were some done? 10 They were done on the skeletal muscle. Α. 11 Q, Okay. And these are the last set of 12 DNA reports? 13 Let me go through those just to make 14 Α. 15 sure. The next page is testing for MELAS, 16 three genetic misspellings in the mitochondrial 17 chromosome and those were not found. 18 The next two pages later we looked for 19 20 the common genetic misspelling and something called MERRF. And we looked for the common 21 misspelling that occurs in patients with a 22 particular fatty acid oxidation disorder which is 23 abbreviated as MCAD. That was not found. 24 25 The last page, as we looked at muscle

1 for, **I** don't know, I think they looked for almost 2 50 misspellings in the mitochondrial DNA in Los Angeles, and they did not find any. And they 3 also looked not just for misspellings but torn 4 out pages or extra pages in the genetic 5 instructions. So the extent of our DNA based 6 testing was for a subset of misspellings in the 7 8 mitochondrial gene.

9 Q. From your earlier testimony, I
10 understand that you did not arrive at a
11 definitive diagnosis that Megan Kaschak had a
12 specific metabolic or genetic abnormality;
13 correct?

14

A. Correct.

Q. Based on your analysis here that was
done, were you able to rule out an underlying
genetic or metabolic abnormality in Megan Kaschak
as being the cause for her respiratory arrest and
subsequent death?

20 MR. MISHKIND: Objection. Go ahead. 21 A. I could rule out only those things 22 that I specifically tested for and found to be 23 consistently normal. And the enzyme assays were 24 found to be normal on the DNA based testing. I 25 could not rule out other possible metabolic

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Page 89 1 diseases. Ο. 2 Were you able to rule out a 3 deficiency, an abnormality related to the complex II abnormalities found in the electron transport 4 chain reports? 5 6 Α. I didn't rule it out. Ο. 7 Okay. 8 Α. I quess I will tell you, it was not ruled out. 9 Q. Is there any evidence in any of these 10 reports or the autopsy reports or any information 11 12 that you have suggestive of any OxPhos metabolic disorder? 13 14 MS. MALNAR: Did you say in the autopsy report, because **I** don't think he reviewed 15 the whole thing. 16 17 MR. NORCHI: I did say that. Go 18 ahead. The concern is that some of those 19 Α. enzyme enzymatic abnormalities in the electron 20 21 transport chain assays do, in fact, reflect an underlying problem. I just didn't say I saw a 22 consistent pattern so I could say that's the 23 24 case. 25 Q. I have had genetics issues arise

Page 90 before in cases like this and T have often had 1 the medical geneticists tell me who I have had 2 look at the case that they can get me to the 3 right state, and get me to the right city; they 4 5 can tell me, in fact, what part of the city and maybe even get me to the street, but they 6 couldn't tell me what the address was that would 7 8 tell me what the exact disorder was for the problem. 9 Is that an explanation or is that --10 let me back up. Is that similar to what we have 11 in this case? 12 Let me object to the 13 MR. MISHKIND: question. It's artful, but I think 14 15 inappropriate. MR. NORCHI: It may be inartful too, 16 but we will find out. 17 Q, Do you understand what I am saying? 18 Α. I know why geneticists tell you that. 19 The answer to your question, though, is we were 20 looking for a possible electron transport chain 21 defect and we were looking as hard as we were 22 able to do given the limitations of what's 23 available. We did not find something. 24 Do I worry or other people who do this 25

Page 91 1 worry that there is something else there? Sure. 2 But I can't say we missed it. I can't say I was 3 just next door to it. Q. The results on these reports show what 4 I think you have termed an inconclusive pattern 5 6 of abnormalities; correct? 7 Α. Yes. Q. 8 And you also indicated that they are suggestive of an OxPhos mechanism disorder? 9 10 No, I didn't say they are suggestive. Α. 11 Q. Are they consistent with? 12 MR, MISHKIND: Objection. Ο. 13 Or did I just use the wrong word? 14 They are not internally consistent to Α. allow me to make a specific diagnosis is what I 15 think I said. 16 I think we are saying the same thing. 17 Q. It's an inconclusive pattern of abnormalities? 18 19 Α. I agree with that statement. Q. There are abnormalities reflected in 20 21 these test results? 22 Α. Correct. 23 Q. But as you review the abnormalities in 24 each of the tests, they don't identify for you a 25 particular pattern that's consistent with a

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Page 92 1 specific disorder? 2 Α. Either for me or when I spoke with Dr. Hoppel about some of these, correct. 3 Q. 4 Was there a neuropathologist who didn't -- I believe there are electron 5 microscopies done of tissue. Is that your 6 7 understanding also? I thought there were, but I didn't 8 Α. specifically review that. 9 10 MS, MALNAR: I believe they are 11 attached to the autopsy report. 12 MR. NORCHI: Microscopies, have you seen them, the actual films? 13 MR. MISHKIND: No, the films are not 14 15 attached. Q. You didn't look at the microscopies; 16 17 correct? 18 Α. I don't remember doing so. Q. 19 I am almost finished here, doctor. 20 MR. NORCHI: Thank you. 21 EXAMINATION OF ARTHUR B. ZINN, M.D., Ph.D. 22 BY MR. MISHKIND: Q. 23 Just a couple questions. Bottom line, the abnormalities that we 24 25 talked about when you put all of the different

Page 93 specimens together, the different reports 1 2 together, and you compare the abnormalities, 3 whether it be the skeletal muscle to the liver, to the heart, et cetera, these results in plain 4 terms do not explain to a reasonable degree of 5 medical certainty why Megan died; correct? 6 7 They do not provide me with a Α. 8 diagnosis that I can use to explain the events of 9 Megan's life. Ο. 10 The complex II syndrome that you said 11 you could not rule out, certainly looking at the flip side, there was insufficient evidence for 12 you to confirm to a probability the existence of 13 a complex II syndrome; correct? 14 15 Α. Correct. Q. If there were such a complex II 16 17 syndrome, of what significance would that be from 18 the standpoint of mortality and morbidity? 19 As I said before, complex II Α. deficiencies are, one, relatively rare, and 20 associated with actually a quite broad range of 21 22 medical problems. 23 Q. Are you qualified to talk to what those problems are? 24 25 I believe so. Α.

	Page 94
1	Q. What are they?
2	A. They can be anywhere from the severest
3	form I think there has been one case of
4	complex II deficiency that's been implicated as a
5	cause of a particular necrotizing encephalopathy,
6	although ${\tt I}$ am not sure how strong that case is.
7	I don't know that there is any other. I think
8	there may be two cases in the literature, which
9	means that the child would have it and die within
10	the first couple years of life.
11	At the other range, there is adults
12	who had no problems until 30s or 40s, walk around
13	and suddenly get exercise intolerance and you
14	can't run the same two miles as fast as they used
15	to. Just based on the enzyme assay ${\tt I}$ could not
16	say more than that.
17	Q. Taking the adult scenario and I
18	understand you can't say to a reasonable degree
19	of medical probability that Megan had complex II
20	syndrome. I ask you in terms of if, in fact, she
2 1	did have that, what are the manifestations you
22	have given me some of them from extreme to the
23	adult situation. The adult situation that you
24	are aware of, besides exercise intolerance, are
25	they associated with any degree of increased

	Page 95				
1	mortality				
2	A. No.				
3	Q in the adult?				
4	A. I stated it correctly. It is simply				
5	an exercise intolerance, clinical problem.				
6	Q. Fair enough.				
7	MR. MISHKIND: No further questions.				
8	Thanks.				
9	MR. NORCHI: Thank you.				
10	MS. MALNAR: You can read this				
11	deposition transcript if you would like to check				
12	for basically typographical errors. It's up to				
13	you. Would you like to read it?				
14	THE WITNESS: If I don't have to, I				
15	think I'm okay.				
16	MS, MALNAR: If someone orders it, I				
17	would like to have a copy of it.				
18	-				
19	(Deposition concluded at 4:00				
20	p.m.; signature waived.)				
2 1					
22					
23					
24					
25					

	Page 96
1	CERTIFICATE
2	
3	State of Ohio,
4	<i>SS</i> :
5	County of Cuyahoga.
6	
7	
8	
	I, Vivian L. Gordon, a Notary Public within
9	and for the State of Ohio, duly commissioned and qualified, do hereby certify that the within
10	named ARTHUR ZINN, M.D., Ph.D., was by me first
	duly sworn to testify to the truth, the whole
11	truth and nothing but the truth in the cause
	aforesaid; that the testimony as above set forth
12	was by me reduced to stenotypy, afterwards
	transcribed, and that the foregoing is a true and
13	correct transcription of the testimony.
14	I do further certify that this deposition
	was taken at the time and place specified and was
15	completed without adjournment; that I am not a
	relative or attorney for either party or
16	otherwise interested in the event of this action.
17	IN WITNESS WHEREOF, I have hereunto set my hand and affixed my seal of office at Cleveland,
1 0	Ohio, on this 28th day of August, 2000.
18 19	· · · · · · · · · · · · · · · · · · ·
20	Vivian L. Hordon
20	Vivian L. Gordon, Notary Public
2 1	Within and for the State of Ohio
22	My commission expires June 8, 2004.
23	
24	
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	BY MR, MISHKIND:	92	22
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4	Exhibit 2 was marked	22	23
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