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SUPERIOR COURT OF THE STATE OF CALIFORNIA
FOR THE COUNTY OF LOS ANGELES

STEVEN AND KATHY WORKMAN,	)	
	)	
Plaintiffs,	)	
	)	
vs.	)	No. BC 250793
	)	
POMA DISTRIBUTING COMPANY, INC., a	)	
California corporation; ADVANTAGE	)	
TANK LINES, INC., a California	)	
corporation; ULTRAMAR, INC., a	)	
Nevada corporation; MILLARD	)	
REFRIGERATED SERVICES, INC., a	)	
Nebraska corporation; MILLARD	)	
REFRIGERATED SERVICES - ATLANTA,	)	
INC., a Georgia corporation;	)	
MILLARD REFRIGERATED SERVICES -	)	
ATLANTA II, INC., a Georgia	)	
corporation; and DOES 1 through	)	
100, inclusive,	)	
	)	
Defendants.	)	
	)	

DEPOSITION OF PHILIP COLE, M.D.

Monday, February 9, 2004

Long Beach, California

REPORTED BY: Lyn Corrin Aaker, CSR No. 6228

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ATLANTA II, INC., a Georgia)
corporation; and DOES 1 through)
100, inclusive,)
)
Defendants.)
)

No. BC 250793

Deposition of PHILIP COLE, M.D., an Expert
Witness, taken on behalf of the Plaintiffs, at
401 East Ocean Boulevard, Suite 800, Long Beach,
California 90802, commencing at the hour of
10:00 a.m., Monday, February 9, 2004, before
Lyn Corrin Aaker, CSR No. 6228, pursuant to
Notice of Taking Deposition.

1 APPEARANCES OF COUNSEL:

2

3 For Plaintiffs:

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6 Attorney at Law
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Long Beach, California 90802
(562) 437-4499

8 For Defendants POMA DISTRIBUTING COMPANY, INC., and
9 ADVANTAGE TANK LINES, INC.:

10 WOOD SMITH HENNING & BERMAN, LLP
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I N D E X

WITNESS	EXAMINATION	PAGE
Philip Cole, M.D.	By Mr. Metzger	5
	By Mr. Metzger (continued)	74

EXHIBITS FOR IDENTIFICATION

Plaintiffs' A	Copy of "Reviews Epidemiologic Perspectives on Myelodysplastic Syndromes and Leukemia"; 5 pages	33
Plaintiffs' B	Copy of Draft "Table A-5"	54
Plaintiffs' C	Copy of "Does Diesel Exhaust Cause Human Lung Cancer?"; 23 pages	55
Plaintiffs' D	Copy of "Tobacco and Cancer: Recent Epidemiological Evidence; 8 pages	56
Plaintiffs' E	Copy of 12/16/03 correspondence with attachments; 6 pages	57
Plaintiffs' F	Copy of "Curriculum Vitae"; 16 pages	67
Plaintiffs' G	Copy of "Cases in Which Testimony Has Been Provided by Philip Cole, MD, DrPH"; 3 pages	67

WITNESS INSTRUCTED NOT TO ANSWER

PAGE	LINE
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1 MONDAY, FEBRUARY 9, 2004, LONG BEACH, CALIFORNIA

2 10:00 A.M.

3 * * *

4

5 PHILIP COLE, M.D.,

6 the witness herein, having been first duly sworn, was

7 examined and testified as follows:

8

9 EXAMINATION +

10 BY MR. METZGER:

11 Q. Would you introduce yourself for the record,
12 Doctor.

13 A. My name is Philip Cole.

14 Q. And you are an esteemed epidemiologist but not a
15 medical doctor. True?

16 A. I think you may be wrong on both counts there. I
17 am a medical doctor, and I make no claim to be esteemed.

18 Q. Oh, okay. Yes, I do see you actually do have an
19 M.D. Okay. Well, Dr. Cole, you've testified a number of
20 times in the past?

21 A. Yes, I have.

22 Q. Let's talk about that briefly. Could you tell me
23 approximately how many depositions you have given in your
24 career?

25 A. I'd say about 70.

1 Q. And about how many trials have you testified in?

2 A. Mr. Metzger, when you say "trial," would that
3 include things that occurred in court, but I'm not sure --
4 some of them aren't actually trials. They may be hearings
5 of a sort.

6 Q. Sure. Any court hearing or trial.

7 A. Maybe 10 or 15.

8 Q. And have you ever testified before any
9 governmental agencies?

10 A. Again, I need some clarification. Would that be,
11 for example, a scientific body or only regulatory body?

12 Q. Are you suggesting that regulatory bodies don't
13 follow science?

14 A. Well, no. I'm sorry if I did. I meant where the
15 purpose was to get information pertinent to a regulation
16 or get information pertinent to some kind of a treatise of
17 a learned nature.

18 Q. Well, let me try clarifying it this way. What
19 I'm referring to is any time you spoke before a
20 governmental agency, that's governmental, not private, and
21 where you took an oath and swore to tell the truth.

22 A. Okay. That's helpful. Yes, I have done that
23 perhaps six or eight times in my life.

24 Q. Okay. Of the approximately 70 depositions that
25 you've given, could you tell me how many of them related

1 to benzene or some form of blood dyscrasia or hematologic
2 malignancy?

3 A. The benzene ones would be about 10 or 12, and the
4 blood dyscrasia would be maybe 15 or so, a few more.

5 Q. Okay. And that would include hematologic
6 malignancies?

7 A. Yes. Hematologic malignancies where benzene was
8 not the issue.

9 Q. Well, then, let's see. Regarding those, I assume
10 some were regarding 1,3 butadiene. Is that true?

11 A. No. I've never testified regarding butadiene.

12 Q. Oh, what were the subjects of the blood dyscrasia
13 depositions?

14 A. They were all the same. They were
15 electromagnetic fields, or EMF for short.

16 Q. Other than EMFs and benzene, have you given
17 depositions regarding any other actual reputative causes
18 of hematologic diseases or malignancies?

19 A. I don't believe so.

20 Q. Could you tell me what the last deposition that
21 you gave regarding a benzene case was?

22 A. I really don't recall now. It's been quite a
23 while.

24 Q. Do you have a list of litigations in which you've
25 testified?

1 A. Yes, I do.

2 Q. Could I have that, please.

3 A. I don't have it with me. I mean, I will be glad
4 to provide it.

5 Q. Okay. It was requested. Could we possibly get
6 that faxed here?

7 A. I think we could do that. It's possible. If we
8 take a break and I call my secretary and if she can find
9 it because it's not on my university file, but I have --
10 she may have access to it.

11 Q. Okay. Let's see. She is on Alabama time?

12 A. Two hours.

13 Q. Two hours. Would she be in now, or this is the
14 noon hour there?

15 A. No. It's quarter to 12:00, so this would be a
16 good time.

17 Q. Okay. Well, let's see. Perhaps we should call
18 for that. I'm trying to think if there's anything else
19 that we will need to call on. You have your CV with you?

20 A. No, I don't. I just assumed that Mr. Amundson
21 would be bringing it.

22 MR. METZGER: Do you have it, Greg?

23 MR. AMUNDSON: No, but I think it's attached to
24 our expert designation. I will look.

25 THE WITNESS: That's easy to have her send, but I

1 could have her e-mail it.

2 BY MR. METZGER:

3 Q. Actually, e-mail both would be fine. We could
4 just print them out. That might be quickest. And does
5 that have your list of publications?

6 A. Yes. Do you want to give me a fax number? I'm
7 sorry. E-mail address.

8 Q. Sure. If she could e-mail it to
9 rmetzger@toxictorts.com.

10 THE WITNESS: Greg, I assume that you are okaying
11 all of this.

12 MR. AMUNDSON: Of course.

13 THE WITNESS: Are we off the record?

14 MR. METZGER: We'll go off the record. Sure.

15 (Discussion held off the record.)

16 BY MR. METZGER:

17 Q. Well, I can ask you a few questions without the
18 list here. Regarding the approximately 10 to 12
19 depositions that you gave regarding benzene, were any of
20 those on behalf of a worker claiming to be injured?

21 A. No.

22 Q. Were they all on behalf of the defense?

23 A. Yes.

24 Q. And did the defense constitute oil companies
25 and manufacturers or distributors of benzene or

1 benzene-containing products?

2 MR. AMUNDSON: Objection; compound. Go ahead and
3 answer.

4 THE WITNESS: The answer is yes.

5 BY MR. METZGER:

6 Q. Okay. And in the EMF cases that you've testified
7 in, were any of those on behalf of someone claiming to
8 have been injured, or were those on behalf of the defense?

9 MR. AMUNDSON: Same objections. Go ahead.

10 THE WITNESS: Sorry. They were all on behalf of
11 defense.

12 BY MR. METZGER:

13 Q. Of the 70 depositions that you've given, how
14 many, if any, of them were where you were engaged as an
15 expert or testified as an expert on behalf of a person
16 claiming to have been injured?

17 A. Three or four.

18 Q. Can you identify what they were?

19 A. When you say "they," do you mean the case or the
20 agents?

21 Q. However you can.

22 A. The agents were two. The first one is DES, which
23 is diethylstilbestrol, and the outcome was clear cell
24 adenocarcinoma of the genital tract in young women. And
25 the other agent was menopausal estrogens, specifically

1 Premarin, and the outcome was breast cancer.

2 Q. With respect to the 70 depositions that --
3 approximately 70 depositions that you've given, excluding
4 pharmaceuticals or medical devices, were all of the
5 depositions on behalf of the defense?

6 A. Yes.

7 Q. Of the approximately 10 to 15 trials or hearings
8 that you've testified in in court, how many of those
9 concerned benzene?

10 A. Perhaps six or so, maybe a little less than half
11 of the total.

12 Q. And in those cases, at those trials did you
13 always testify on behalf of the defense?

14 A. Yes.

15 Q. Of the 10 to 15 trials that you've testified in,
16 have all of them been for the defense?

17 A. No.

18 Q. Excluding pharmaceuticals and medical devices,
19 have they all been on behalf of the defense?

20 A. I'll need some clarification. The agent in
21 question is DES. It was administered to one person as a
22 pharmaceutical, but the person who was injured was not the
23 person who received it as a pharmaceutical. So I'm not
24 sure what the answer to your question is.

25 Q. The injury was from a pharmaceutical agent. It

1 was an in utero exposure, and the daughter was a DES
2 daughter. Is that what we're talking about?

3 A. Yes.

4 Q. Other than that, were all the trials for the
5 defense?

6 A. Yes. And the other one -- no. My answer is yes.

7 Q. Okay. Now, of the approximately six to eight
8 times you've testified before governmental agencies, have
9 any of those concerned benzene?

10 A. No.

11 Q. What have been the subjects of those testimonies?

12 A. The electromagnetic fields, formaldehyde, many
13 years ago saccharine, chlorpyrifos, and an agent whose
14 chemical name I have forgotten, but it was the principal
15 ingredient in a shampoo that is used to rid children of
16 lice.

17 Q. Lindane?

18 A. I think that's correct. I think Lindane was
19 either the or one of the principal ingredients.

20 Q. All right. When you testified before the
21 governmental agency regarding EMF, what agency was that?

22 A. Well, there were several state agencies. Shall I
23 mention which ones they were?

24 Q. Sure.

25 A. State of Maine, State of New Jersey, and

1 Illinois. There was a federal agency. It was actually a
2 congressional committee. I can't remember now what
3 committee it was. Oh, and, also, State of Florida.

4 Q. Approximately when was this testimony before the
5 congressional committee?

6 A. That would have been quite a while ago, about
7 1987 or '88.

8 Q. On whose behalf were you testifying?

9 A. When you say "on whose behalf," you mean who was
10 sponsoring the appearance there?

11 Q. Yes. Who paid you?

12 A. I believe that was paid -- well, I should say
13 that I'm not sure, but my best recollection is that it was
14 on behalf of EPRI, E-P-R-I.

15 Q. And what is EPRI?

16 A. That stands for the Electric Power Research
17 Institute.

18 Q. That's an industry organization, according to
19 your understanding?

20 A. Well, the financing of the institute comes from
21 industry, yes.

22 Q. And when you testified regarding formaldehyde,
23 what agencies was that before?

24 A. I don't recall.

25 Q. And on whose behalf did you testify for

1 formaldehyde?

2 A. I'm sorry. Isn't that what you just asked?

3 Q. I asked you what agency.

4 A. In front of what agency? I'm sorry. It was in
5 front of EPA.

6 Q. About when was that?

7 A. Mid 1980s.

8 Q. And was it some industry trade organization on
9 whose behalf you testified?

10 A. No. It was on -- I think different people were
11 represented, sponsored by different corporations. I
12 believe I was sponsored by Shell Oil Company.

13 Q. Chlorpyrifos, what agencies did you testify
14 before regarding that?

15 A. I believe it was a scientific advisory board of
16 NIOSH.

17 Q. About what year was that?

18 A. About 1995 or so. I'll just say that there is a
19 publication that will be listed on my CV which will give
20 us specific information on that.

21 Q. Okay. And who sponsored your testimony?

22 A. That was sponsored by a chemical company. I
23 can't remember now whether it was Dow or Dupont. I think
24 it was one or the other of those two.

25 Q. Was it the register of chlorpyrifos?

1 A. I really don't know.

2 Q. Do you recall generally the thrust of your
3 testimony at that hearing, what it was about?

4 A. Chlorpyrifos?

5 Q. Yes.

6 A. It was actually as a participant in a roundtable
7 discussion, and I don't recall my specific assignment or,
8 indeed, even if I had one. But I can tell you that the
9 purpose of the roundtable was to review the available
10 epidemiologic information and to make a recommendation to
11 the agency as to what, if any, action they might take.

12 Q. And what recommendation, if any, was made?

13 A. I believe there were two. I think one was that
14 the available evidence did not support the idea that
15 chlorpyrifos was a cause of any form of cancer in human
16 beings, and that the research that the agency had done and
17 which they were considering either updating or upgrading
18 in some way should, in fact, have those improvements made
19 if possible.

20 Q. What improvements?

21 A. Well, that was not clear to us either. It was
22 simply stated that they were considering upgrading the
23 study that they had done.

24 Q. Okay.

25 A. Whether it meant extending it in time or size or

1 what, either was not known at that time or was known at
2 that time but I have forgotten.

3 Q. And when you testified regarding Lindane, what
4 agency was that before?

5 A. That was in front of a committee of the Food and
6 Drug Administration.

7 Q. About when was that?

8 A. That would have been about 1995.

9 Q. And who sponsored your testimony?

10 A. The manufacturer of the shampoo.

11 Q. Do you recall who that was?

12 A. No. I just remember that it was a relatively
13 small pharmaceutical company.

14 Q. Okay. Now, have you in the past done work for
15 industry apart from the testimony of the cases that you've
16 worked on?

17 A. Yes.

18 Q. Have you done work for the American Petroleum
19 Institute?

20 A. I have done two things that I can recall for API.

21 Q. What are they?

22 A. One was I wrote or I coauthored a chapter in a
23 book or monograph that they produced, and the other was at
24 their request I attended a meeting of their executive body
25 to discuss certain issues with them regarding time trends

1 in cancer.

2 Q. Okay. We'll talk about these. What was the
3 subject of the chapter in the book?

4 A. The chapter was in effect what is the current
5 state of knowledge regarding, and based on the
6 epidemiologic information, the status of gasoline as a
7 human carcinogen. And, in fact, particularly it might
8 have been limited. I'm not sure now. It might have been
9 limited to unleaded gasoline.

10 Q. And in what book was that published? Do you
11 recall?

12 A. Well, again, if we had my CV, it would be cited
13 there.

14 Q. Very well. We'll wait for your CV. Okay. Now,
15 regarding the meeting of the executive body regarding time
16 trends in cancer, what was this executive body, according
17 to your understanding?

18 A. Mr. Metzger, I was about to say my understanding
19 is a little fuzzy now, but certainly the CEO of the
20 institute was there. There were perhaps seven or eight
21 other people. As I recall, three or four of these people
22 were credentialed in science, and the other three or four
23 were, let's say, using the term generally, policy makers
24 and advisors.

25 Q. Who were they?

1 A. You mean their names?

2 Q. Yes.

3 A. I don't remember the names.

4 Q. Any of them, the scientists?

5 A. I don't remember any of the names, not even the

6 CEO.

7 Q. Was any paper prepared regarding this meeting?

8 A. No.

9 Q. What was the general thrust of the advice that

10 you gave the committee?

11 A. I didn't give them any advice.

12 Q. Did they ask you anything?

13 A. Did they ask me anything? Yes. They had asked

14 me beforehand to come and make a presentation along

15 certain lines.

16 Q. And you did that?

17 A. Yes.

18 Q. What was the subject of your presentation?

19 A. They had heard that I had said -- in fact, I had

20 said it at a meeting of the National Cancer Advisory

21 Board, and I'm sorry if I didn't mention that in response

22 to your previous questions. I'm not sure whether that

23 would have qualified or not.

24 Q. That's okay.

25 A. That I had said that we were soon to experience a

1 downturn in the overall cancer mortality rate and various
2 consequences of that and the reasons for it, and they
3 wanted to hear more about that.

4 Q. What are the consequences you're referring to?

5 A. The extension of life among adults.

6 Q. Okay. Anything else?

7 A. No.

8 Q. And the reasons for it?

9 A. The reason, there being two principal reasons.
10 One is the plateauing -- it hadn't happened yet at the
11 time, you understand, but the prediction of the plateau in
12 mortality rates from lung cancer and the --

13 (Telephone interruption.)

14 BY MR. METZGER:

15 Q. You were saying plateauing of morality from lung
16 cancer?

17 A. And advances in treatment and diagnosis and
18 treatment or let's say medical care in general.

19 Q. Is there anything else that you have done over
20 the years for the American Petroleum Institute that you
21 can think of?

22 A. No.

23 Q. Have you ever --

24 A. Well, wait a minute. I do remember a third
25 thing. Unfortunately, I don't remember what it was. I

1 just remember that it existed. I did go to their
2 headquarters in New York and did attend a roundtable
3 discussion of some sort. I don't remember what the issue
4 was.

5 Q. Did you ever receive any contracts or have any
6 contracts with the American Petroleum Institute?

7 A. I certainly don't have any now, and for the three
8 instances that I have mentioned I was paid. I don't
9 believe it was under contract but, rather, that I simply
10 bill them after the fact.

11 Q. So your best recollection is you've never had any
12 contracts with the American Petroleum Institute?

13 A. That is correct.

14 Q. Approximately how many contracts have you had
15 with oil companies?

16 A. I need to ask a question of clarification. When
17 you say how many have I had, do you mean that were
18 personal with me in a private capacity or with the
19 university and related to work in which I was either the
20 principal or a significant participant?

21 Q. Either of those two and, also, of any contracts
22 with any consulting organizations with which you have been
23 affiliated.

24 MR. AMUNDSON: The latter might be a little
25 vague, but go ahead if you can answer the question.

1 THE WITNESS: Let me try to be systematic about
2 it. As far as contracts with me, I actually recall only
3 one. That was with Texaco perhaps some 15 or so years
4 ago.

5 BY MR. METZGER:

6 Q. Regarding what?

7 A. It was regarding a risk assessment for benzene
8 and leukemia, and it resulted in a publication that
9 appears on my CV and which acknowledges that support.

10 With regard to the university, there were three
11 or four contracts for our research in which the university
12 was actually the contracting party, not me. And two or
13 three of these were with the Shell Oil Company. I'm
14 sorry. Did the question relate to oil companies?

15 Q. Oil companies or petrochemical companies. I'm
16 not limiting it to Shell Oil. It could also include
17 Shell Chemical, for example.

18 A. Well, actually, when I said Shell Oil, I perhaps
19 should have said Shell Oil or Chemical. It's a
20 distinction I don't keep very clear in my mind.

21 Q. Nor do I.

22 A. There was at least one with a company which at
23 the time was called Ciba-Geigy, now known as Novartis.
24 There was one with an entity whose name I cannot recall,
25 but I can tell you the spirit of it, and that is it's a

1 manufacturers' trade association of the synthetic rubber
2 producers. That's all that I recall.

3 Q. Okay. Could you tell me approximately regarding
4 the contract that you personally had with Texaco what the
5 funding of that was?

6 A. I really can't remember.

7 Q. Can you estimate? Was it five figures or six
8 figures or seven figures?

9 A. By "five figures" you mean under \$100,000?

10 Q. Yes.

11 A. Yes. It was under \$100,000. In fact, it was
12 probably, as I recall, in the vicinity of \$10- to \$15,000.
13 But I would point out that that was not all paid to me.
14 There were two other participants in that.

15 Q. And the contracts that you had with Shell Oil,
16 what was the funding for those?

17 A. Again, I want to point out that I did not have
18 these contracts.

19 Q. The university.

20 A. Those contracts would have been for six figures,
21 that is in excess of 100,000 and less than one million,
22 and, in fact, much closer to 100,000 than to one million.
23 But I was not the principal investigator in those, and I
24 really don't recall the budget figures.

25 Q. Okay.

1 A. I'd like to mention another one just so that it
2 doesn't appear later that I was not forthcoming. There
3 was also one with Amoco Company, which was I believe in
4 the vicinity of \$75,000.

5 Q. Were the contracts with Shell Oil regarding
6 benzene?

7 A. Two are them were, in fact, primarily focused on
8 benzene and were for the conduct of a retrospective
9 follow-up study or case control study of leukemia in one
10 of their refineries, specifically the Wood River refinery.
11 And that work led to two or three publications which are
12 included in my CV.

13 Q. And the other one?

14 A. The other Shell?

15 Q. Yes.

16 A. That one was not focused on benzene. As I
17 recall, it was primarily concerned with pesticides which
18 had been manufactured at a facility that Shell -- that was
19 actually owned by the United States Government but
20 operated for the United States Government by Shell. And I
21 don't think it was focused on any particular chemical, but
22 on the health of the work force.

23 Q. Did that result in a publication?

24 A. Yes.

25 Q. And that's on your CV?

1 A. Yes.

2 Q. The Amoco consultation, what was that regarding?

3 A. That related to a perceived outbreak of brain
4 cancer at a research and development facility that Amoco
5 ran in a city that's a suburb of Chicago. Its name
6 escapes me right now.

7 Q. Naperville or something.

8 A. It is Naperville, yes.

9 Q. And the synthetic rubber producers, what was that
10 about?

11 A. That was primarily focused on butadiene.

12 Q. And did that result in a publication?

13 A. That work which has been renewed now several
14 times, so the span of time is probably close to 12 years,
15 has resulted in a number of publications, perhaps four or
16 five. I think I am included in the authorship of only one
17 or two of those, and yes, they would be on the CV.

18 Q. Did these publications explore hematopoietic
19 malignancies?

20 A. When you say "these publications," I'll take it
21 to mean all of the ones I've mentioned, and I can answer
22 yes with the exception of the Amoco.

23 Q. I'm talking about the synthetic rubber ones.

24 A. The principal focus was on what I'll call LHC,
25 lymphatic hematopoietic cancer.

1 Q. And was that also true of the Ciba-Geigy?

2 A. I'll need to think for a minute about that one.

3 The answer is no.

4 Q. Okay. Do you recall what that was about
5 generally?

6 A. It was primarily focused on two things. One was
7 adverse outcomes of pregnancy in the female employees, and
8 secondly on the health of the work force in general.

9 Q. Was there a suspect agent that you recall?

10 A. The suspect agent was a category of compounds
11 called triazenes.

12 Q. Could you estimate for me the funding for the
13 projects that you consulted on, that the university had
14 and on which you consulted for the synthetic rubber
15 manufacturers?

16 A. I really don't know what the budget has become.
17 I don't even recall what it was in the first few years. I
18 was never the principal investigator on it.

19 Q. Do you know whether it was a six- or seven-figure
20 contract?

21 A. Anything I say would be a guesstimate, but I
22 would say it would probably have run somewhere in the
23 vicinity of \$75,000 to \$100,000 per year, each of the
24 contracts running for several years. And I want to just
25 make it clear that you used the expression in that

1 question that I had consulted for. I was not a
2 consultant. I was a collaborator.

3 Q. Fair enough.

4 A. And I just want to make it clear that I was not
5 paid. That is my university was reimbursed for my salary.

6 Q. When you've done these projects and the
7 university had the contract, has your salary reflected any
8 type of compensation, remuneration, or bonus due to the
9 amounts that these contracts brought into the university?

10 A. Well, certainly not a bonus. There is no bonus
11 in the university setting, at least not that I've ever
12 received or known of. Whether or not my salary was
13 influenced -- let me phrase it this way.

14 Whether or not the person who decided whether or
15 not I would receive increases in pay took this into
16 consideration, I can only say the following: For many of
17 these years I was the head of the department, so the
18 circumstance was that I was in a position to recommend my
19 own salary. And all I can say is that the recommendation
20 that I made at least in those years, which was about 15 of
21 the 20 or so years that I was on faculty at UAB, reflected
22 my perception of my worth to the department, which
23 included teaching, mentoring, particularly young faculty,
24 and also research and, yes, the amount of research funding
25 that I thought I was at least in part instrumental in

1 bringing into the department. So I don't recall that my
2 recommendation to my superior, who would have been the
3 dean of the school, was ever not accepted, so I guess the
4 short -- that's the long answer. Maybe the short answer
5 is yes.

6 Q. Fine. I understand. Thank you. What about have
7 you ever been a private consultant through a consulting
8 firm?

9 A. I have I guess the word would be helped out or
10 participated in projects that were being done by
11 consulting firms, yes.

12 Q. And have you had any ownership interest in any of
13 these consulting firms?

14 A. I had an ownership interest in one of these firms
15 for some time, yes.

16 Q. What was that firm?

17 A. That firm, which is no longer in existence, was
18 called the New England Epidemiology Institute.

19 Q. Okay. What types of research or projects did
20 that organization do?

21 A. Just for clarity, do you want to know what they
22 did, or do you want to know in which ones I participated?

23 Q. We'll take the ones you participated in.

24 A. I participated in one -- let me start over again.
25 I participated in many in which they asked me to review

1 the report or work product that had been prepared by their
2 staff, but I had no direct involvement in the actual
3 procurement of the contract or the conduct of the
4 research. I did have an active role in one that related
5 to the projection of the number of cases that would be
6 filed against the Johns Manville Company as a result of
7 asbestos exposure.

8 I also participated -- well, no. I have to
9 retract that. I don't recall another one. The company as
10 a whole did epidemiologic research of all sorts.

11 Q. I'm sorry?

12 A. The company in a larger sense -- you asked what
13 the company did as well as what I did. The company
14 provided epidemiologic services of many types.

15 Q. That was this New England Institute?

16 A. Yes.

17 Q. Are there any other consulting firms that you've
18 been affiliated with?

19 A. I don't like to use the word "affiliated with"
20 when the relationship was essentially a strictly ad hoc
21 relationship. But the answer is yes, and the company is
22 Exponent.

23 Q. And what type of work have you done for Exponent?

24 A. I have done two things for them. One is that I
25 participated in a committee which they convened on the

1 subject of the appropriate classification of
2 carcinogenicity to humans of dioxin. That also resulted
3 in a publication that is listed in my CV. And the other
4 project was the participation in an advisory group which
5 was pulled together for the purpose of evaluating research
6 that had been conducted by a governmental agency. I'm
7 sorry I've forgotten the exact name of it, but it relates
8 to occupational health in Scotland.

9 Q. Have you ever done any consulting for the
10 Chemical Manufacturers' Association or the Manufacturing
11 Chemists' Association or I think it's now called the
12 American Chemistry Council? They don't like the name
13 "Manufacture" in there any more, I guess.

14 A. And the first part of the question?

15 Q. Have you ever done any consulting for that
16 organization?

17 A. I don't know that it would be called consulting,
18 but I'll tell you what the activity was. I have on
19 several occasions given talks to either their own
20 membership or to more general audiences at meetings that
21 they had convened.

22 Q. Talks regarding what?

23 A. I really don't recall.

24 Q. About how many talks?

25 A. Two that I recall.

1 Q. Were they given before any certain committees of
2 the CMA?

3 A. One was given to a group of people, all of whom
4 were either members of the CMA, if there is such a thing
5 as that, or scientists or executives of the companies that
6 support the CMA. But I don't recall the subject.

7 Q. All right. Well, let's start talking about this
8 case, and we'll get back to some of these other things
9 when we have your CV and the case list. You understand
10 that you've been engaged as an expert in this case, and
11 you've been asked to form opinions on certain topics.
12 True?

13 A. I've been asked to form opinions?

14 Q. Have you not been asked to form opinions on
15 certain topics?

16 A. Yes.

17 Q. Can you tell me, please, what the opinions are
18 that you've formed regarding the present case.

19 A. If I recall correctly, there is a document that
20 states what I will be or may be asked to testify about.
21 Is that the same thing that we're talking about?

22 Q. Well, not really. I assume that was prepared by
23 counsel. I'd like to know what your opinions are as
24 opposed to what counsel says you might talk about.

25 MR. AMUNDSON: I think what he's really asking

1 for is if you could give him kind of a highlight of your
2 table of contents of the various subjects on which you
3 will be opining at trial. Just kind of a quick hit list,
4 and he will go through each of your opinions and ask you
5 more definitive opinions.

6 BY MR. METZGER:

7 Q. That's true. I'll ask you for the foundation,
8 the basis, and the reasons. But first I'd like to get the
9 landscape of what you're going to be testifying about.

10 A. I think some are fairly obvious. Some are not at
11 all obvious to me. But I suppose I will be asked what I
12 consider to be the causes of the myelodysplastic
13 syndromes. I suppose I would be asked and would venture
14 an opinion as to whether or not the MDS's are caused by
15 benzene. That might incorporate any of a number of
16 subopinions regarding dose response and the like.

17 I might be asked to opine as to what kind or
18 which type of MDS Mr. Workman had. I might be asked and
19 would opine whether or not he had acute myelocytic
20 leukemia, hereafter AML, and I might be asked and try to
21 be responsive with an opinion as to whether or not his
22 particular condition was caused by his particular exposure
23 to benzene or, possibly as a variant of that opinion,
24 whether or not it was caused by the benzene that would
25 have been supplied by Mr. Amundson's client. I don't know

1 whether I will be asked about the possible role of smoking
2 in Mr. Workman's illness.

3 Q. Okay. Any other general topics, or should I
4 start exploring these?

5 A. I can't think of any more general topics.

6 Q. Okay. Well, let me ask you, then: What do you
7 consider to be the known causes of myelodysplastic
8 syndromes?

9 A. I don't know of any.

10 Q. Well, have you conducted any literature searches
11 to ascertain the literature that pertains to the etiology
12 of the myelodysplastic syndromes?

13 A. Yes. In fact, about six or seven years ago,
14 granted that may be a little out of date, I actually
15 published a review of the epidemiology of the MDS's. On
16 that particular one we wouldn't necessarily have to wait
17 for the CV if you're interested in it because I brought a
18 copy along. Since then I have maintained a familiarity
19 with the literature on the epidemiology and, hence, I
20 think the causes or suspect causes of MDS.

21 Q. Well, could I see the review article that you're
22 speaking of?

23 A. Sure. It was actually an invited talk that I
24 published.

25 Q. Okay. And you're referring to the article in

1 "Leukemia Research" which we'll mark as Exhibit A.

2 (A copy of the aforementioned document was
3 marked by the court reporter as Plaintiffs'
4 Exhibit+ A for identification; attached hereto.)

5 BY MR. METZGER:

6 Q. And who invited you to give this talk?

7 A. Dr. John Bennett.

8 Q. And did you know Dr. Bennett before this?

9 A. Yes.

10 Q. How did you know him?

11 A. I had met him previously at several scientific
12 meetings. I also met him on at least one occasion at
13 Wood River.

14 Q. And that was in connection with Shell's
15 investigation of the leukemias at that facility on which
16 you were engaged as a consultant, and Dr. Bennett was
17 engaged as a consultant?

18 A. No, it wasn't.

19 Q. Please tell me about it.

20 A. I think that that particular meeting where
21 John Bennett and I -- in which John Bennett and I both
22 participated, and there were other people as well, was
23 actually on the issue of MDS.

24 Q. And who participated in that meeting?

25 A. The participants were Dr. Sally Cowles,

1 Dr. Elizabeth Delzell, Dr. John Bennett, me. There were
2 several technical people who were employed at the
3 Wood River refinery and one or two management people as
4 well.

5 Q. Did Dr. Bennett, to your knowledge, review any of
6 the slides of the MDS and AML cases out of Wood River?

7 A. I have no knowledge of that.

8 Q. Was that your first meeting with John Bennett?

9 A. No. I had met him several times previously.

10 Q. At what?

11 A. At scientific meetings.

12 Q. Such as?

13 A. Well, we're talking about a 20- or 25-year time
14 span, and I just don't recall. He was a person who when I
15 met him at this meeting I recognized and walked directly
16 up to him and said, "Hello, John. How are you?" So I had
17 met him before several times.

18 Q. Had you met him at the meetings of the American
19 Society of Hematology?

20 A. No.

21 Q. Had you met him at any other medical meetings?

22 A. I believe I met him on at least one occasion at a
23 meeting of the Society for Epidemiologic Research.

24 Q. Any other meetings that you recall meeting him
25 at?

1 A. No. I mean, there were other meetings, but I
2 don't recall which ones they were.

3 Q. Before you met with him at Wood River, had you
4 met with him in connection with any other benzene leukemia
5 clusters or investigations?

6 A. No.

7 Q. John Bennett, I believe, had once prepared a
8 report to the American Petroleum Institute arguing against
9 medical monitoring of benzene-exposed workers. Did you
10 have any involvement in that?

11 A. The meeting at Wood River was for the purpose of
12 discussing that issue. I don't know -- I don't recall
13 whether his report -- and, by the way, I don't have a
14 recall, a recollection of that report. I don't know
15 whether that report preceded or followed that meeting.

16 Q. I'm speaking actually of a report that he
17 prepared for the American Petroleum Institute many years
18 earlier than Wood River.

19 A. May I ask in what year or about what year that
20 report was?

21 Q. I'd have to dig it out. It was around the time
22 of the Benzene Emergency Temporary Standard, so late '70s.

23 A. Late '70s. Yes. That would be many years before
24 the Wood River meeting. And you're asking whether or not
25 I influenced him or had any discussions with him about

1 that report?

2 Q. Yes.

3 A. No.

4 Q. You've never seen or heard of that report?

5 A. I have no recall of ever seeing it.

6 Q. Okay. Well, let's see. When did you first
7 conduct a search of the literature to investigate causal
8 associations for myelodysplastic syndromes?

9 MR. AMUNDSON: Just for the record, if I can have
10 a continuing line on this, the term "causal associations"
11 is objected to as vague and ambiguous and contradictory.
12 You can answer or respond to that question in your own
13 words.

14 MR. METZGER: He is an epidemiologist. He can
15 certainly deal with it.

16 MR. AMUNDSON: I'm sure he can.

17 THE WITNESS: I would say that I did not do ever
18 a literature search that I would have characterized as
19 oriented towards -- in any direct sense towards the causes
20 or causes of MDS but, rather, was directed towards the
21 epidemiology of MDS, which I intend to mean as a broader
22 subject.

23 BY MR. METZGER:

24 Q. When did you first do literature search regarding
25 the epidemiology of MDS?

1 A. May I see that review article? The first search
2 probably would have been done in about 1994, this paper
3 being published in 1995 but actually is an elaborated
4 version of a talk that I gave in 1944.

5 Q. '94?

6 A. What did I say?

7 Q. '44?

8 A. 1994.

9 Q. Now, incidentally, did you receive any funding or
10 compensation for this publication or the presentation that
11 you gave?

12 A. I recall with regard to this -- may I look at it
13 to refresh my memory? I recall certainly receiving
14 reimbursement for my expenses in going to Chicago to make
15 the presentation. I don't recall -- I don't believe I was
16 compensated in any other way.

17 Q. Was the university, to your knowledge?

18 A. No.

19 Q. Now, would you happen to have -- when you did
20 this search in 1994, did you do that using on-line data
21 bases?

22 A. No.

23 Q. How did you do that research, that search?

24 A. It actually was done by a graduate student
25 working for me.

1 Q. Who is the graduate student?

2 A. He is one of the co-authors on the report. His
3 name is Warren Sateren.

4 Q. Do you know how Mr. Sateren did the literature
5 search?

6 A. He did not use any kind of computerized data
7 base. He essentially worked from a recent textbook and
8 worked back from that.

9 Q. What textbook did he use?

10 A. I think the one that he used was the one called
11 "Blood."

12 Q. By Jandl?

13 A. Yes.

14 Q. And that was the 1990s? No, it couldn't have
15 been. Which edition of Jandl did he use?

16 A. I don't remember. It would have been whichever
17 one was current at the time.

18 Q. I think the one that I have is '96, which would
19 post date this. Okay. And did Mr. Sateren come up with a
20 list of articles from that method?

21 A. Yes. He came up with a fairly extensive list.

22 Q. Of approximately how many articles?

23 A. I don't remember.

24 Q. Do you have that list?

25 A. No. The publication includes a list of the

1 references that would actually have been included in it.

2 Q. Okay. Were there any other articles on that list
3 other than those included in this publication?

4 A. Oh, yes.

5 Q. Could you tell me what they are?

6 A. No.

7 Q. Have you ever done an on-line search?

8 A. You mean on this subject or in general?

9 Q. In general.

10 A. Yes.

11 Q. And do you find the on-line search to be helpful
12 in being sure to obtain the literature existing on topics?

13 A. Yes.

14 Q. Did you tell Mr. Sateren that he should do an
15 on-line search when he did this?

16 A. No.

17 Q. Why not?

18 A. I felt that the first thing was that I wanted to
19 see what he would do. And when I got the list that he
20 developed, I looked at it, and I looked in a textbook, and
21 I wanted to see whether or not there was likely to be any
22 significant omissions from it. And I decided there was
23 none, and so I selected from his list the papers that I
24 thought would be helpful to our writing this review.

25 Q. And what was the textbook that you referred to?

1 A. It was a textbook by Schattenfeld and Fraumeni
2 called "Cancer Epidemiology and Prevention." I might have
3 also looked --

4 Q. That was the first edition of that?

5 A. No. But let me finish, if I may.

6 Q. I'm sorry.

7 A. I might have also looked in both "Blood" and in
8 this book, "Clinical Hematology."

9 Q. Wintrobe?

10 A. Yes.

11 Q. And the Schattenfeld and Fraumeni book, was that
12 the first edition?

13 A. No. It would have been the edition prior to the
14 current one. And I can't tell you what number that would
15 be. I think the current one is perhaps Edition 5, so that
16 would have been Edition 4.

17 Q. What is the title of this textbook?

18 A. It's called "Cancer Epidemiology and Prevention."

19 Q. Right. Okay. And first of all, is MDS a cancer?

20 A. No.

21 Q. And can you tell me what you would expect to find
22 regarding myelodysplastic syndrome in Schattenfeld and
23 Fraumeni's textbook?

24 A. Well, now, wait a minute. When you ask me what I
25 would expect to find, do you mean as if I had never gone

1 there before?

2 Q. No. Specifically regarding MDS, since it's not a
3 cancer.

4 A. The textbook includes a number of conditions
5 which might be labeled preneoplastic or paraneoplastic
6 conditions, just as does the book "Clinical Hematology,"
7 for example. So I'm not sure. When you ask me that
8 question what I would expect to find, I guess the answer
9 is that I would expect to find some level of discussion
10 and description of the epidemiology of the conditions
11 known as the MDS's, and I did.

12 Q. Okay. Do you recall specifically whether you
13 consulted Wintrobe's "Clinical Hematology" in doing this
14 work?

15 A. Well, unfortunately, my memory of "Clinical
16 Hematology" would be very difficult to isolate from my
17 current perceptions of what's in the book, which is a
18 different edition. So I think the answer to your question
19 is no.

20 Q. Do you recall specifically whether you consulted
21 Jandl's "Blood" in the course of this literature search?

22 A. I can't say for sure that I did, but I believe I
23 did.

24 Q. Do you recall any other textbook or any other
25 means by which you familiarized yourself with the extent

1 literature regarding the epidemiology of myelodysplastic
2 syndromes at the time in 1994?

3 A. No.

4 Q. Since 1994, have you ever done a computerized
5 data base literature search regarding epidemiology of MDS?

6 A. No.

7 Q. Have you ever done any other noncomputerized data
8 base search for literature of the epidemiology of MDS?

9 A. I'm not sure whether the answer to the question
10 is yes or no. Maybe I should describe what I have done.

11 Q. Sure.

12 A. I do try to stay current with the literature on
13 the epidemiology of LHCs, and as part of that one of the
14 things that I do is look at the references that accompany
15 the papers that I get. And if one came to my attention
16 which I did not have and which I thought might be of
17 interest to me, either regarding the MDS's or one of the
18 LHCs, then I would simply obtain that.

19 Q. And do you have a list of those articles that you
20 obtained?

21 A. I don't keep such list, no.

22 Q. Can you identify them for me?

23 A. When you say "identify" them to you, do you mean
24 can I provide you with a list of the papers?

25 Q. Identify them by author and year, somehow

1 identify them.

2 A. It's not the nature of the identification. It's
3 the category. What is it that we're referring to here;
4 the literature on leukemia or the literature on MDS or the
5 literature on MDS that I got in that particular way?

6 Q. We were talking specifically about the
7 epidemiology of MDS. That's what I'm talking about.

8 A. And you want me to tell you, if I can, the
9 citations that are relevant?

10 Q. I'm not asking you to determine what's relevant.
11 I'm asking if you can identify for me citations simply by
12 author, preferably author and year, of epidemiology
13 studies concerning myelodysplastic syndrome that you've
14 become familiar with since the publication of your article
15 in "Blood" and the research that you did for that in 1994.

16 A. There are only two papers that come to my mind.
17 There are other papers, but I don't -- I'm not in a
18 position to be able to tell you who authored them.

19 Q. Fine.

20 (Telephone interruption.)

21 MR. METZGER: Let's go back on.

22 Q. Dr. Cole, you were about to tell me of two papers
23 that you recall regarding the epidemiology of MDS which
24 you reviewed since you prepared your publication in
25 "Blood" in 1995.

1 A. Actually, I think it's twice you mentioned the
2 journal as "Blood." It's actually "Leukemia Research."

3 Q. I'm sorry. It is "Leukemia Research." You're
4 right.

5 A. I don't recall whether either of these papers was
6 specifically or even primarily oriented towards MDS, but I
7 do recall that there was information in them pertaining to
8 MDS. And one is a paper by Hayes and others, and the
9 other one is by Yin and others, and both reflect
10 epidemiologic research done in China.

11 Q. Are there any other epidemiology studies
12 concerning MDS that you have reviewed since the research
13 that you did for your article in "Leukemia Research" in
14 1995?

15 A. There are others, but I don't recall them at
16 present.

17 Q. And have you reviewed any of those for purposes
18 of this case?

19 A. May I take a minute to look through my file?

20 Q. Certainly.

21 A. (Witness peruses document.)

22 There is one other paper for a fact I reviewed in
23 connection with this case. It stands out only because it
24 was provided to me yesterday in a package of materials by
25 Mr. Amundson. Since I got it, I looked at it, and I

1 believe there is another one in here.

2 Q. Are you referring to the Albin paper?

3 A. Yes. Do you want me to cite it for the record?

4 Q. Perhaps -- I'll do that. You're referring to the
5 Albin paper from 2003 "Cytogenetic and Morphologic
6 Subgroups of Myelodysplastic Syndromes in Relation to
7 Occupational and Hobby Exposures." Correct?

8 A. Yes. That's the only other one -- well, it's
9 actually the only one that I specifically reviewed for
10 this case. I think I mentioned the other two pages in
11 connection with literature that I have reviewed since my
12 own review.

13 Q. Let's just take a moment now and deal with some
14 of the documents. There is a collection of papers
15 including the Albin paper which you just pulled out of an
16 envelope. The envelope says Philip Cole, M.D., as the
17 recipient, and the sender is R.G. Amundson, Wood Smith
18 Henning & Berman. Correct?

19 A. Yes.

20 Q. And are these all papers which Mr. Amundson gave
21 you yesterday to review before your deposition?

22 A. I don't know. I have to look through it and see
23 whether or not some things that I brought with me have
24 gotten mixed in or things that he did send have gotten
25 pulled out.

1 Q. Please take a look and answer that.

2 A. (Witness peruses document.)

3 Let me begin my answer to your question by saying
4 that everything that I'm holding in my hand was provided
5 to me yesterday by Mr. Amundson. However, a number of
6 things here were previously available to me.

7 Q. Let's talk about them, if I might.

8 A. Okay. Could I -- it might help if I finish by
9 going through this to see whether or not there are any
10 things in here that actually came with this pile.

11 Q. Oh, sure. Please do that as well.

12 A. (Witness peruses document.)

13 Okay. There is not.

14 Q. Okay. Then let me just ask you these questions.
15 Is yesterday the first time that you saw the article by
16 Albin?

17 A. No.

18 Q. You had reviewed that previously?

19 A. I hadn't reviewed it. I had seen it.

20 Q. You had seen it, but you had not reviewed it
21 previously?

22 A. That's correct.

23 Q. So the first time you reviewed it, actually
24 studied it, was yesterday or today after Mr. Amundson
25 provided it to you. Correct?

1 A. I wouldn't say I have studied it. I have
2 reviewed it.

3 Q. You have reviewed it. Fair enough. The article
4 by Sandler "Cigarette Smoking and Risk of Acute Leukemia,"
5 JNCI '93, had you seen that before yesterday?

6 A. Seen it, reviewed it, and studied it.

7 Q. You did that at about the time that it was
8 published?

9 A. I really don't recall.

10 Q. The article by Korte, K-o-r-t-e, in
11 "Environmental Health Perspectives" entitled "The
12 Contribution of Benzene to Smoking-Induced Leukemia," had
13 you seen that before yesterday?

14 A. Yes.

15 Q. And had you studied that before yesterday?

16 A. Yes.

17 Q. There is an abstract by Marsh, "Mortality
18 Patterns Among Petroleum Refinery and Chemical Plant
19 Workers," "American Journal of Industrial Medicine 1991."
20 Had you seen that before yesterday?

21 A. Well, I have not actually seen the abstract in
22 isolation before. I have that paper.

23 Q. And have you studied that paper before?

24 A. Yes, yes.

25 Q. There is an excerpt from the report to the

1 scientific review panel on benzene prepared by the
2 California Department of Health Services in 1984. Had you
3 seen that before yesterday?

4 A. I have seen the whole report.

5 Q. There is something printed off the Internet on
6 Polycythemia Vera. Does that have anything to do with
7 this case in your view?

8 A. Not that I could tell.

9 Q. Nor could I. There is an article by Madl and
10 Paustenbach, M-a-d-l, "Airborne Concentrations of Benzene
11 Due to Diesel Locomotive Exhaust in a Roundhouse." Had
12 you seen that article before yesterday?

13 A. Yes.

14 Q. And had you studied that before yesterday?

15 A. Yes.

16 Q. What was the connection that you studied that
17 article? It's not an epidemiology article, is it?

18 A. Yeah, I think it is. It is not epidemiology per
19 se, but it is highly relevant to epidemiology, I believe.

20 Q. What was the context that you came across that
21 article?

22 A. I have been involved as an expert witness in a
23 number of cases in which the putative cause of a disease
24 is diesel exhaust or diesel fuel exposure, and it is in
25 that context. And let me just say -- excuse me -- I

1 should have acknowledged that in response to one or more
2 of your previous questions. I have never actually been
3 deposed in a diesel case.

4 Q. I see. How many such cases have you consulted
5 on?

6 A. Four that I can recall.

7 Q. Are all of these consultations on behalf of
8 industry?

9 A. Yes.

10 Q. Do you recall the names of the cases in which you
11 consulted?

12 A. When you say "the name," you mean the last name
13 of the first plaintiff?

14 Q. Yes. Some way of identifying it.

15 A. I do recall some of them.

16 Q. Please identify them.

17 A. Okay. One I know as the case of Alexander. I
18 think it's versus Norfolk Southern Railroad. That case
19 has recently been settled. The next one is Dampier,
20 D-a-m-p-i-e-r, also versus Norfolk Southern. The next
21 one -- and I'm actually going back in time -- is Shapona,
22 S-h-a-p-o-n-a. I can't remember who the defendant was. I
23 think it might have been a city. He was a firefighter.

24 Q. Can you remember the fourth?

25 A. The fourth I don't remember the name, but the

1 defendant I do recall was also Norfolk Southern.

2 Q. In these cases did you prepare written reports?

3 A. In the last one no. Now I'm coming forward. In
4 Shapona, yes; in Dampier, yes; in Alexander, no.

5 Q. Do you know whether -- strike that. Were these
6 reports given to anyone other than the defense attorney
7 who had engaged you?

8 A. I have no idea.

9 Q. Have you prepared any publications regarding --
10 which derive from any of these reports?

11 A. No.

12 Q. With respect to these four cases, were they all
13 lung cancer cases?

14 A. No.

15 Q. Were any of them leukemia or MDS cases?

16 A. No.

17 Q. The next article in this stack by Pogoda "Smoking
18 and Risk of Acute Myeloid Leukemia," did you see that
19 article before yesterday?

20 A. Yes.

21 Q. And did you study that before yesterday?

22 A. Yes.

23 Q. There is an excerpt from --

24 A. It's a textbook.

25 Q. Yes, it is. What's the name of the author?

1 A. The principal editor is DeVita.

2 Q. Right. Of course. This is the 5th edition from
3 1997 regarding myeloid leukemias. Had you read this
4 excerpt from DeVita before yesterday?

5 A. Excuse me. May I look at it just a moment?

6 Q. Certainly.

7 A. The reason I asked to look at it is I think this
8 is the 6th edition, and you said 5th. But the answer is
9 no, I have not looked at that material in it before. It
10 does not say 6th edition.

11 Q. I'm sorry. I must have misspoken. There is
12 attached to that stapled to it a copy of Schnatter's
13 article "Determination of Leukemogenic Benzene Exposure
14 Concentrations." Have you seen that article before?

15 A. Oh, yes.

16 Q. And did you study that before yesterday?

17 A. Yes.

18 Q. There is an abstract by Nagean, N-a-g-e-a-n, on
19 Polycythemia Vera. We'll put that in an applicable pile.

20 There is an excerpt from the toxicological
21 profile for benzene, 1997, by the ATSDR. Have you seen
22 that publication before yesterday?

23 A. Yes. The whole publication.

24 Q. And did you study it?

25 A. Yes, I have studied it.

1 Q. There are some photographs here of the facility.
2 I'm sure you didn't see those before yesterday. Right?

3 A. That is true.

4 Q. There is an article by Fedoruk, et al.,
5 "Measurement of Volatile Organic Compounds Inside
6 Automobiles." Had you seen that article before yesterday?

7 A. No.

8 Q. Have you studied that article?

9 A. No.

10 Q. There is the ACGIH. I assume this is the TLV
11 documentation for benzene. Have you seen that before?

12 A. May I have a look at it when you're through?

13 Q. This is actually copyright 2001. I will see
14 where it ends here.

15 A. (Witness peruses document.)

16 I can't say that I have seen this particular
17 version before.

18 Q. All right. There is a deposition summary of
19 Jimmy Morrison, M.D., and also one of Don Kuchenbecker.
20 Have you read those?

21 A. No.

22 Q. There is a health hazard evaluation report by
23 NIOSH with the Costa Mesa Fire Department. Have you seen
24 that before?

25 A. No.

1 Q. There is another copy of Madl's.

2 A. May I see that, please?

3 (Witness peruses document.)

4 It is a copy.

5 Q. There is a copy of the declaration of
6 Dr. Brautbar and apparently his -- oh, no. Yes, and a
7 minuscrite of his deposition. Have you read either of
8 those?

9 A. Both.

10 Q. And there is a minuscrite, a rough version of
11 Fedoruk's deposition. Have you read that?

12 A. Okay. Is that a minuscrite?

13 Q. It's a rough version. It's not a minuscrite.
14 It's a rough version.

15 A. And is your question have I read that?

16 Q. Yes.

17 A. The answer is yes.

18 Q. Could I see the other materials that you've
19 brought with you?

20 A. You want me to just give you the whole batch?

21 Q. Yes. And we'll talk about them. You have here a
22 table of "Tabular Values of 95 Percent Confidence Limit
23 Factors for Estimates of a Poisson-distributed Variable."
24 This is something that -- well, what is this?

25 A. This is a page from a textbook that I bring with

1 me to depositions because I have found that it often comes
2 in handy when a question of whether or not something is
3 statistically significant comes up, and it is a shortcut
4 way of getting confidence intervals.

5 MR. METZGER: Okay. Let's mark that as
6 Exhibit B.

7 (A copy of the aforementioned document was
8 marked by the court reporter as Plaintiffs'
9 Exhibit+ B for identification; attached hereto.)

10 BY MR. METZGER:

11 Q. At this point let me sign this one and give you a
12 check for \$1,000 at least so you're not -- I'm not in debt
13 to you as of the present moment.

14 A. Are we on the record?

15 Q. I guess we are. I apologize.

16 A. May we go off the record?

17 Q. Surely.

18 (Discussion held off the record.)

19 MR. METZGER: Back on the record.

20 Q. There is an article here "Does Diesel Exhaust
21 Cause Human Lung Cancer?" By Tony Cox. Have you seen that
22 before yesterday?

23 A. Oh, yes. That was mine. I brought that.

24 Q. That was yours.

25 A. Yes.

1 MR. METZGER: Let's mark that as Exhibit C.

2 (A copy of the aforementioned document was
3 marked by the court reporter as Plaintiffs'
4 Exhibit+ C for identification; attached hereto.)

5 BY MR. METZGER:

6 Q. Okay. And you've also brought with you the 9th
7 edition, Volume 2, of Wintrobe's "Clinical Hematology,"
8 which we're not going to mark as an exhibit.

9 A. Good.

10 Q. Are there any other materials that you have
11 brought with you regarding this case?

12 A. Yes. I think you didn't finish going through the
13 pile of stuff that I gave you.

14 Q. I'm sorry. No. I think I did. Oh, there's more
15 here. I apologize. You're right. There is an article by
16 Paolo Vineis. Is that how you pronounce his name?

17 A. Vineis.

18 Q. V-i-n-e-i-s. "Tobacco and Cancer: Recent
19 Epidemiological Evidence," JNCI current January 21, 2004.
20 And is this something that you just recently obtained or
21 that Mr. Amundson gave you.

22 A. No. He did not give it to me. I obtained it
23 myself about -- actually, I think it was faxed to me by
24 another attorney perhaps two weeks ago or so.

25 Q. It says Brown, McCarrol. Who are they?

1 A. That's a law firm in Texas.

2 Q. Have you done work with them before?

3 A. I've done work with them over the years, yes.

4 Q. And did you ask them if they had anything
5 regarding that might be helpful for this case?

6 A. No.

7 Q. How did it come to you?

8 A. The attorney who sent it is a Mr. George Butts,
9 and I have worked with him on a number of benzene and
10 leukemia cases, and he sent it because he considered it
11 relevant to that issue.

12 Q. Unsolicited?

13 A. Correct. Well, I don't know if solicited or
14 unsolicited is correct. He had sent me an e-mail asking
15 me if I had seen the paper, and, if not, if I would like
16 him to send it to me, and I answered no and yes to those
17 two questions.

18 MR. METZGER: We'll mark that paper as
19 Exhibit D.

20 (A copy of the aforementioned document was
21 marked by the court reporter as Plaintiffs'
22 Exhibit+ D for identification; attached hereto.)

23 BY MR. METZGER:

24 Q. And then there is a yellow sheet on top of it.
25 It says "NTPB Particulates and IARC 2A Limited in Humans,"

1 and that's referring to the classification of diesel
2 exhaust as a carcinogen. Is that correct?

3 A. I would say not as a carcinogen, but it refers to
4 the classification assigned by those two agencies.

5 Q. Regarding diesel exhaust carcinogenicity?

6 A. Yeah, that's okay.

7 Q. Sure. Okay. Let me just ask you: Do you
8 consider diesel exhaust to be a human carcinogen?

9 A. No.

10 Q. Okay. Do you consider benzene to be a human
11 carcinogen?

12 A. I consider it to be a human leukemogen.

13 Q. Fair enough. And then there is also a letter
14 from Paul Lee with some notes, handwritten notes, and a
15 pathology report for Mr. Workman. Have you reviewed this?

16 A. Yes.

17 MR. METZGER: We'll mark that as Exhibit E.

18 (A copy of the aforementioned document was
19 marked by the court reporter as Plaintiffs'
20 Exhibit+ E for identification; attached hereto.)

21 BY MR. METZGER:

22 Q. Are there any other materials that you've been
23 provided by counsel for this case?

24 A. Yes.

25 Q. What are they?

1 A. They are a number of medical records and several
2 depositions including the deposition of Mrs. Workman.
3 Those are the only two categories of information that I
4 recall.

5 Q. Are you relying on any of those records for any
6 of your opinions in this case?

7 A. I rely upon the medical records. I rely to some
8 degree on the deposition of Mrs. Workman.

9 Q. Other than the diagnosis of MDS and the other
10 hematologic conditions reported in the medical records,
11 are you relying on the medical records for anything else?

12 MR. AMUNDSON: Let me just object that the brief
13 statement of diagnosis of MDS and other hematological
14 conditions is vague and ambiguous, misleading, and a
15 misstatement. But go ahead and answer the question.

16 THE WITNESS: May I ask what your last word was?

17 MR. AMUNDSON: And a misstatement.

18 THE WITNESS: And a misstatement. Okay. May I
19 ask to have the question read back?

20 BY MR. METZGER:

21 Q. Let me ask it this way: What are you relying on
22 the medical records for?

23 A. I rely upon the medical records primarily for two
24 things: One is the virtual certainty that Mr. Workman had
25 one of the myelodysplastic syndromes, the one usually

1 referred to as RARS. And I rely upon it, on them, in a
2 way which is a little difficult because it's the absence
3 of something. But I rely upon them for the failure of the
4 documentation of the existence of AML.

5 Q. So you've reviewed the medical records you were
6 provided. Correct?

7 A. Yes.

8 Q. And from your review of the records, Mr. Workman
9 never received a diagnosis of AML. Correct?

10 A. I could not -- although I found allusions to that
11 diagnosis and reference to it, I never found the
12 documentation of it.

13 Q. Well, do you have an opinion as to whether
14 Mr. Workman had AML?

15 A. The only opinion I have is that I have not seen
16 any documentation of that diagnosis.

17 Q. I'm asking you if you personally -- since you are
18 a medical doctor, I'm asking you if you personally have an
19 opinion whether he had AML or whether he did not have AML.

20 MR. AMUNDSON: Within reasonable medical
21 probability?

22 MR. METZGER: Sure. He is a medical doctor.

23 Q. I'm asking you whether you have an opinion, not
24 what your opinion is.

25 A. I believe I've answered this question.

1 Q. I don't know that you have.

2 MR. AMUNDSON: Go ahead and answer it again.

3 THE WITNESS: Let me answer it again. And if the
4 answer is not satisfactory, maybe we can work on it.

5 BY MR. METZGER:

6 Q. Let me interrupt you because I heard your answer.
7 What you said was you had an opinion that certain things
8 were referenced. I'm not asking you what's referenced in
9 the records. I'm asking you whether you have an opinion
10 as to whether Mr. Workman had AML.

11 A. Yes.

12 Q. What is your opinion?

13 A. No.

14 Q. Upon what do you base your opinion that
15 Mr. Workman did not have AML?

16 A. I would base it on two things -- three things.
17 First, I reviewed a very large amount of medical records,
18 and I found no documentation of this condition; and I feel
19 it is likely that if he were to have had this condition,
20 it would have been documented in those records. That's
21 No. 1.

22 No. 2, he had a condition known as RARS, which
23 very infrequently converts to a blast crisis, and there is
24 no evidence that that conversion occurred in this case.
25 Furthermore, I think that the use of the language AML to

1 describe the blast crisis of MDS is a shorthand commonly
2 used in the medical profession because the blast crisis
3 has the morphologic appearance of AML, but I am not
4 persuaded that the blast crisis of MDS is, in fact, AML.

5 The third reason, Dr. -- is it Upadhyaya.

6 MR. AMUNDSON: Upadhyaya I believe is how she
7 pronounces it.

8 THE WITNESS: This is a lady who gave a
9 deposition in this case.

10 -- commented, and I believe she was a treating
11 physician for Mr. Workman and was familiar with the
12 circumstances, his circumstances close to even if not
13 right at the end of his life. And she mentions that there
14 was no blast crisis and that, in her opinion, there was
15 not a so-called conversion to AML. So those are the three
16 reasons why I think that Mr. Workman did not have AML.

17 BY MR. METZGER:

18 Q. Okay. Now, have you ever diagnosed AML in a
19 patient?

20 A. No.

21 Q. Have you ever diagnosed MDS in a patient?

22 A. No.

23 Q. Have you ever treated any patients with
24 hematologic conditions?

25 A. Yes.

1 Q. How long ago?

2 A. Well, I think the best way to answer it is from
3 about 1968 to about 1975.

4 Q. How many patients did you treat with hematologic
5 conditions?

6 A. And by "hematologic" may I understand the LHCs?

7 Q. Not limited to cancers including blood
8 dyscrasias.

9 A. Well, anything that I treated would have been a
10 malignancy.

11 Q. Okay.

12 A. But I just wanted to know if I could include the
13 lymphatic series.

14 Q. Sure.

15 A. And the question was how many?

16 Q. Yes.

17 A. Something in the vicinity of 50 or 60. I want to
18 say I was not the sole provider of their care.

19 Q. Are you an oncologist?

20 A. No. Not insofar as that term is used to refer to
21 the practice of clinical oncology.

22 Q. Are you a hematologist?

23 A. No.

24 Q. What was the context in which you treated these
25 50 to 60 patients?

1 A. I was a fellow in the hematology and oncology
2 clinics of the Massachusetts General Hospital over this
3 period of time.

4 Q. And how many patients -- well, during that time
5 period, did myelodysplastic syndrome exist as a disease
6 entity?

7 A. I'm not sure it exists as a clearly defined
8 disease entity even today, but the answer to your question
9 is that the word "myelodysplastic syndromes" was in use.
10 However, the classification and terminology was not well
11 developed at that time.

12 Q. Did you ever make a diagnosis of myelodysplastic
13 syndrome in any patient during that period?

14 A. Perhaps I could save us some time by mentioning
15 that this was a treatment clinic and that the patients who
16 were seen at that clinic had been diagnosed before coming
17 there.

18 Q. I'm asking --

19 A. No. I never actually made, myself, the diagnosis
20 of any of the myelodysplastic syndromes or any of the
21 lymphatic hematopoietic cancers.

22 Q. Including AML?

23 A. Including AML.

24 Q. Okay. Now, what is the percentage of blasts --
25 strike that. Are you familiar with the different

1 classification systems for MDS?

2 A. When you say the different systems, do you mean,
3 in fact, the different systems or the different categories
4 of MDS?

5 Q. No. I'm talking about the different
6 classification systems; World Health Organization, FAB,
7 those types of systems.

8 A. I'm only familiar with the FAB.

9 Q. And under the FAB, what is the percentage of
10 blasts that is required for an MDS to be deemed to have
11 converted to an AML?

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

13 Q. And you are not familiar with that classification
14 under the World Health Organization?

15 A. That is correct.

16 Q. Okay. Would you define blast crisis, please, as
17 you use it.

18 A. No, I can't define it. I use it when a physician
19 who is involved in the care of a patient says that such a
20 phenomenon has occurred.

21 Q. Is it your belief that there is a blast crisis of
22 MDS?

23 A. Certainly there can be in some cases, yes.

24 Q. Do you have any source to support that?

25 A. I want to be sure I understand the question. Do

1 I have a source to support my representation that
2 something described as a blast crisis can occur --

3 Q. In MDS?

4 A. -- as part of the natural history of some cases
5 of MDS?

6 Q. Yes.

7 A. Yes.

8 Q. What's that source?

9 A. It's here in this book (indicating).

10 Q. Wintrobe?

11 A. Yes.

12 Q. All right. Now, have you ever made a study of
13 the epidemiologic studies concerning alkylating agent
14 therapy and secondary MDS?

15 A. I will need a little clarification here. When
16 you say have I ever made a study of this phenomenon, do
17 you mean have I done any of the original data generating
18 research on it?

19 Q. We can start with that. I am familiar with your
20 background. I assume the answer is no. Is that correct?

21 A. That is correct.

22 Q. Have you ever made a literature search for the
23 epidemiologic studies concerning alkylating agent therapy
24 and the development of MDS?

25 A. I have never done a literature search that was

1 specifically oriented to that question.

2 Q. And have you ever evaluated causality for
3 alkylating agent therapy and MDS?

4 MR. AMUNDSON: Objection; the term "causality" is
5 a legal term. If you want to answer within the field of
6 epidemiology, go ahead, Doctor.

7 THE WITNESS: I'd like to have the question read
8 back.

9 MR. METZGER: Sure. Please read it back.

10 (Question read.)

11 THE WITNESS: Only in the context that I have
12 attempted to maintain a familiarity with the causes of the
13 disease. I have not done anything specifically to explore
14 that particular possible relationship.

15 BY MR. METZGER:

16 Q. Okay. Have you ever done any literature search
17 to ascertain the available epidemiologic literature
18 concerning alkylating agent therapy and secondary AML?

19 A. I need some clarification of this expression
20 which you have used several times and perhaps will use
21 again, and that is have I made a literature search of.
22 And I'd like to understand whether or not you mean that I
23 have specifically through one means or another attempted
24 in some systematic way to identify the literature that
25 pertained or that might pertain to that specific issue.

1 Q. That's exactly what I mean.

2 A. The answer is no.

3 Q. And without having done that, have you ever made
4 an assessment of causality for alkylating agent therapy
5 and secondary AML?

6 MR. AMUNDSON: Objection; calls for a legal
7 conclusion. If you can answer from an epidemiological
8 standpoint, go ahead.

9 THE WITNESS: Again, I'd like to ask for the
10 question to be read back.

11 (Question read.)

12 THE WITNESS: Yes.

13 BY MR. METZGER:

14 Q. How did you do that?

15 A. Well, first I'd like to say not only did I do it,
16 but I published it. And has my CV come?

17 Q. Yes. Let's mark your CV as Exhibit F.

18 (A copy of the aforementioned document was
19 marked by the court reporter as Plaintiffs'
20 Exhibit+ F for identification; attached hereto.)

21 MR. METZGER: And your case list we'll mark as
22 Exhibit G.

23 (A copy of the aforementioned document was
24 marked by the court reporter as Plaintiff's
25 Exhibit+ G for identification; attached hereto.)

1 THE WITNESS: Allow me just a minute.

2 BY MR. METZGER:

3 Q. Sure.

4 A. May I respond now?

5 Q. Certainly.

6 A. My publication No. 182 is a book chapter which
7 lists the causes of the various forms of cancer, and it
8 describes alkylating agents as a cause of acute
9 myelogenous leukemia. That book chapter is now about
10 almost four years old.

11 Q. Okay. And when you wrote that book chapter, how
12 did you determine that alkylating agent therapy causes
13 AML?

14 A. There are two ways that I determine it. But the
15 way that I determined it for this particular book chapter
16 is that that book chapter is a recapitulation of the IARC
17 Group 1 agents. If there were a cause-effect relationship
18 according to IARC that I disagreed with, then I would have
19 noted that in that chapter, not necessarily by saying I
20 disagree but, rather, that the nature of the evidence was
21 less than persuasive or something along those lines.

22 Q. Is benzene an alkylating agent?

23 A. No.

24 Q. Does it have any alkylating properties?

25 A. I don't know.

1 Q. Is benzene a topoisomerase 2 inhibitor?

2 A. I don't know.

3 Q. Are you familiar with a condition known as the
4 5q- syndrome?

5 A. Well, I call it the 5q deletion syndrome. But
6 when you say am I familiar with it, I am aware of its
7 existence.

8 Q. How did you become aware of its existence?

9 A. I don't know. I've been aware of it for quite a
10 while.

11 Q. And what is that condition?

12 A. I can't describe it.

13 Q. Have you made a systematic search of the
14 literature to ascertain epidemiologic studies concerning
15 that condition?

16 A. No.

17 Q. Have you ever attempted to investigate the causes
18 of that condition?

19 A. No.

20 Q. Are you familiar with a condition known as
21 monosomy 7?

22 A. Yes.

23 Q. And could you define that for us?

24 A. It's the absence of one of the No. 7 chromosomes.

25 Q. How did you become familiar with that condition?

1 A. I really don't know how to answer that other than
2 to say I have been aware of the existence of that
3 condition for many years.

4 Q. Have you ever conducted a systematic search of
5 the literature to ascertain epidemiologic studies which
6 bear upon that condition?

7 A. Well, the answer to your question is no. But I
8 just want to state, because this question about systematic
9 searches of the literature has come up several times, that
10 this is not the way in which I do my work. It would be a
11 rare thing for me to go to the computer and do what I
12 think you mean by a systematic review of the literature.
13 In fact, I would never do it. I have a person who would
14 do it for me. I conduct my work by finding the most
15 recent paper and then going back from that. And then when
16 I get to the point where I'm starting to get -- paths are
17 starting to cross, I will usually stop. I'll stop at that
18 point.

19 Q. Well, then, let me ask you: Have you ever done
20 that process that you just described for the 5q- syndrome?

21 A. No.

22 Q. Have you ever done that process for the
23 monosomy 7?

24 A. No.

25 Q. Other than the papers which we've identified here

1 today and which are referenced in your article in
2 "Leukemia Research" regarding myelodysplastic syndromes,
3 can you identify for me any other articles which assess
4 the epidemiology of myelodysplastic syndromes?

5 A. I think I have mentioned the papers by Hayes and
6 Yin. I don't know if that was included in your question.

7 Q. Yes.

8 A. The answer is no, I cannot, from my memory.

9 Q. And can you identify for me any articles which
10 assess the epidemiology of the 5q- syndrome?

11 A. No.

12 Q. Or monosomy 7?

13 A. No.

14 Q. Or the 5q- syndrome in MDS?

15 A. No.

16 Q. Or the monosomy 7 in MDS?

17 A. No.

18 Q. Have you ever studied the benzene leukemia
19 literature to ascertain the frequency that a
20 benzene-induced AML is preceded by a myelodysplastic
21 process?

22 MR. AMUNDSON: Can I hear that question again.

23 (Question read.)

24 THE WITNESS: Could I ask you to read it one more
25 time a little bit more slowly.

1 (Question reread.)

2 THE WITNESS: May I ask a question of
3 clarification?

4 BY MR. METZGER:

5 Q. Certainly.

6 A. I believe the question uses the term a
7 benzene-induced case of AML.

8 Q. Yes.

9 A. Is that correct?

10 Q. That's correct.

11 A. I don't know of any way of knowing that any
12 particular case of AML is, in fact, benzene induced. And
13 so I don't know how I can respond to the question if the
14 context of the question is with regard to one or more
15 individually identified human beings with the disease AML.

16 Q. All right. Well, let's try it this way, then.
17 Have you ever reviewed the epidemiologic literature of
18 benzene exposed -- strike that. Have you ever reviewed
19 the epidemiologic literature pertinent to benzene exposure
20 and the development of AML to determine the frequency of a
21 precedent myelodysplasia?

22 MR. AMUNDSON: Objection; vague.

23 THE WITNESS: I need to be very sure I understand
24 the question. Have I ever looked at the literature which
25 is or attempts to be descriptive of the frequency with

1 which the diagnosis of AML is known or thought to have
2 been preceded by a set of findings that could be described
3 as one of the MDS's? And I assume that we are excluding
4 here from the category of cases of AML those conditions
5 which arise as a blast crisis in a known case of MDS.

6 BY MR. METZGER:

7 Q. No. That's not my question at all. Off the
8 record.

9 (Discussion held off the record.)

10 (Lunch recess was at 12:17 p.m.)

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1 MONDAY, FEBRUARY 9, 2004, LONG BEACH, CALIFORNIA

2 AFTERNOON SESSION

3 1:13 P.M.

4 * * *

5

6 (The question read as follows:

7 "Q. All right. Well, let's try it this way,
8 then. Have you ever reviewed the epidemiologic
9 literature of benzene exposed -- strike that. Have
10 you ever reviewed the epidemiologic literature
11 pertinent to benzene exposure and the development of
12 AML to determine the frequency of a precedent
13 myelodysplasia?")

14 THE WITNESS: No.

15

16 CONTINUED EXAMINATION +

17 BY MR. METZGER:

18 Q. Do you have an opinion as to the frequency of
19 precedent myelodysplasia?

20 A. No.

21 Q. Do you know what caused Mr. Workman's MDS?

22 A. No.

23 Q. I think this question is probably the same, but
24 I'm going to ask it just to be thorough. Do you have an
25 opinion as to what caused Mr. Workman's MDS?

1 A. Well, truthfully, I don't see a distinction
2 between them, but the answer is no.

3 Q. For purposes of this next question I'd like you
4 to assume that Mr. Workman had AML. The question is:
5 Assuming that to be true, do you have an opinion as to
6 what caused Mr. Workman's AML?

7 MR. AMUNDSON: Okay. The question is vague and
8 ambiguous, compound, and an incomplete hypothetical.
9 You're asking him to opine as to the cause of a
10 hypothetical medical condition.

11 MR. METZGER: No, not at all. AML is not a
12 hypothetical medical condition. I'm asking him my
13 question. Your objections are noted for the record.

14 MR. AMUNDSON: Okay.

15 THE WITNESS: I'll need clarification of an
16 issue. When you say I shall assume that he had AML, I
17 want to be clear that I understand whether or not you mean
18 that he had a blast crisis in MDS that is often referred
19 to as AML, or, in fact, he developed the condition which
20 would be the same as the usual de novo AML, which usually
21 is not preceded by a diagnosable -- no. Which is usually
22 not preceded by a diagnosed MDS.

23 MR. AMUNDSON: Why don't you reread the question.

24 BY MR. METZGER:

25 Q. Let me approach it this way. You've made

1 reference to certain references in Mr. Workman's medical
2 records of a diagnosis of AML. I'm asking you to assume
3 that those references are true and that Mr. Workman did,
4 in fact, have AML.

5 A. Okay.

6 Q. Do you have an opinion as to the cause of his
7 AML?

8 MR. AMUNDSON: How can he answer that, Counsel?
9 You've given him a hypothetical situation.

10 MR. METZGER: Just make your objections for the
11 record.

12 +MR. AMUNDSON: Well, if you don't want to
13 clarify it, then I'll object that it's an incomplete
14 hypothetical, it's vague and ambiguous, and instruct him
15 not to answer.

16 MR. METZGER: You're not going to instruct him
17 not to answer.

18 MR. AMUNDSON: I just did.

19 MR. METZGER: I'll seek to exclude his testimony.

20 Q. Will you answer the question, Dr. Cole?

21 MR. AMUNDSON: If you think you understand what
22 he's asking, go ahead and respond to it.

23 THE WITNESS: I cannot respond to this question
24 yet because it has a premise, at least as I understood the
25 question -- it does go by a little fast -- that the record

1 or the records refer to this man as having a documented
2 case of AML.

3 Now, I may have misunderstood the question, but I
4 think that my recollection is that that is what you said.
5 I have never seen in the medical records a documentation
6 of the existence of AML in the case of Mr. Workman.

7 BY MR. METZGER:

8 Q. All right. Then let me approach it this way.
9 I'd like you to assume that Mr. Workman had AML, whether
10 that was a de novo or a progression of a preexisting
11 myelodysplastic syndrome. And I would like to know
12 whether you have an opinion as to the cause of the AML.

13 MR. AMUNDSON: Objection; vague and ambiguous,
14 compound, unintelligible, and an incomplete hypothetical,
15 calls for the witness to speculate.

16 THE WITNESS: The answer is no.

17 BY MR. METZGER:

18 Q. Okay. Among the general topics that you've told
19 me you had opinions on was whether Mr. Workman's condition
20 was caused by exposure to benzene. What is your opinion
21 on that topic?

22 A. The answer is that there is no evidence that it
23 was caused by benzene.

24 Q. On what do you base that conclusion?

25 A. The first is that benzene exposure is not a

1 generally accepted cause of MDS. The second is that from
2 what I would understand of his exposures to benzene in
3 this particular case, they would not be adequate to
4 increase his risk of any particular condition. Those are
5 the two reasons or the two bases for my statement.

6 Q. Do you have an opinion as to Mr. Workman's
7 cumulative received dose of benzene?

8 A. Yes, I do.

9 Q. What was it?

10 A. My opinion is that his exposure to benzene would
11 be minimal and far below that which is generally
12 considered to be the first doubling dose for the condition
13 known as AML.

14 Q. Are you able to quantify Mr. Workman's received
15 dose of benzene?

16 A. I can semi-quantify it.

17 Q. Well, give me your best quantification that you
18 can.

19 A. Okay. Minimal.

20 Q. I would say that's qualitative rather than
21 quantitative. Are you able to give me a quantitative
22 range or estimate of his cumulative benzene exposure?

23 A. It is far below that which is the first doubling
24 dose for AML.

25 Q. And what do you consider to be the first doubling

1 dose for AML?

2 A. 100 ppm years.

3 Q. Are you aware of any study which has validated
4 part per million years as the most sensitive indicator of
5 benzene-induced leukemia?

6 A. May I ask what you mean by the statement "the
7 most sensitive indicator of"?

8 Q. Sure. I'll answer your question with an
9 explanation, and we'll do it this way. There are
10 different metrics which can be used to assess benzene
11 exposure. True?

12 A. Yes.

13 Q. And one of those is a cumulative part per million
14 year dose. Correct?

15 A. Yes.

16 Q. One can also assess benzene exposure by duration.
17 True?

18 A. Yes.

19 Q. And one can also assess benzene exposure by peak
20 exposures. True?

21 A. Yes, yes.

22 Q. These are all different metrics of benzene
23 exposure.

24 A. Yes.

25 Q. Are you aware of any study which has validated

1 part per million years as the most accurate or sensitive
2 metric for assessing causality of benzene-induced
3 leukemia?

4 A. But that's my question. It's not whether or not
5 there are other metrics and what they may be, but what you
6 mean by the most sensitive or accurate.

7 Q. Okay. What I mean is the one that most -- that
8 best conforms to epidemiology.

9 MR. AMUNDSON: Does that make sense to you?

10 THE WITNESS: I think it will in another minute
11 or two.

12 MR. AMUNDSON: Okay.

13 BY MR. METZGER:

14 Q. Let me approach it this way, Dr. Cole. Maybe
15 this will help. Are you aware of any epidemiologic
16 studies which have evaluated odds ratios for risk of
17 leukemia in benzene-exposed workers in which the studies
18 compare different metrics?

19 A. There are several such studies as that.

20 Q. And can you name any?

21 A. I can't name any offhand. But I don't know how
22 one would know from any small number of such studies that
23 the metric -- how to decide which metric is the most
24 accurate predictor of risk enhancement. There has been a
25 tendency to use the metric that gives the highest odds

1 ratios or SMRs. I think this is fundamentally fallacious.
2 Generally speaking -- no. I want to retract the words
3 "generally speaking."

4 It is considered to be true for all known human
5 carcinogens that some measure of cumulative exposure,
6 whether it's ppm years or some other way of combining
7 intensity and duration, is the most accurate descriptor of
8 risk with the exception -- no. I'll end my response by
9 saying this: With one exception and one possible
10 qualification.

11 Q. What are you referring to?

12 A. By the exception?

13 Q. Yes.

14 A. I am referring to the circumstance that the risk
15 of leukemia subsequent to radiation, ionizing radiation
16 exposure, in some circumstances may reach a plateau and
17 not be a strict monotonic function of exposure. That's
18 one. And the qualification is that -- I hope I said
19 possible qualification. I should have said possible
20 qualification -- many epidemiologists and toxicologists
21 are of the point of view that it may turn out to be true
22 that there is an intensity below which the exposure should
23 not contribute to the measure of ppm years for -- well,
24 possibly in general.

25 Q. Are you aware of any study that has proved a

1 threshold for benzene?

2 A. First I have to ask if you could use some other
3 word other than "proved" because I hold proof at least in
4 scientific circles to be a virtually unattainable
5 standard.

6 Q. Well, using the word "proved" as you've just
7 indicated, are you aware of any study that has proved that
8 there is a threshold exposure to benzene below which
9 leukemia cannot occur?

10 MR. AMUNDSON: Objection; the question is vague
11 and ambiguous, argumentative as worded, as in the use of
12 the word "proved." That's it.

13 THE WITNESS: I should answer?

14 BY MR. METZGER:

15 Q. Please.

16 MR. AMUNDSON: If you can.

17 THE WITNESS: I understood the question to end
18 with the expression "below which benzene does not occur."
19 I'm sorry. "Below which leukemia does not occur."

20 BY MR. METZGER:

21 Q. Cannot occur.

22 A. Cannot occur. May I ask if the question could be
23 qualified to below which an increase in the risk of
24 "benzene" cannot occur?

25 Q. No. I'd like you to answer the question that I

1 asked you.

2 A. No.

3 Q. That is your answer?

4 A. Yes. My answer is no.

5 Q. Now, isn't it true that what is known of the
6 mechanisms by which benzene causes leukemia is
7 inconsistent with the parts per million year metric?

8 MR. AMUNDSON: Objection; vague and ambiguous.

9 THE WITNESS: My answer is I don't know.

10 BY MR. METZGER:

11 Q. Okay. Have you researched the literature to
12 ascertain the dose of benzene that is necessary to cause
13 the 5q- syndrome?

14 A. No.

15 Q. Have you researched the literature to ascertain
16 the dose of benzene which is necessary to cause the
17 monosomy 7?

18 A. No.

19 Q. Are you familiar with any articles by Aksoy
20 regarding benzene and myelodysplastic syndrome?

21 A. Yes.

22 Q. Which ones?

23 A. Well, I don't remember which ones now, but I am
24 aware that there was one.

25 Q. And what was the finding of that study?

1 A. I don't recall.

2 Q. Are you familiar with any studies by Aul, A-u-l,
3 regarding benzene and myelodysplastic syndrome?

4 A. Yes.

5 Q. Which one?

6 A. I don't recall.

7 Q. What was the finding of that study?

8 A. I don't recall.

9 Q. Are you familiar with any studies by Baslo,
10 B-a-s-l-o, regarding benzene and myelodysplastic syndrome?

11 A. No, I am not.

12 Q. Are you familiar with any studies by Brandt,
13 B-r-a-n-d-t, regarding benzene exposure and
14 myelodysplastic syndrome?

15 A. No, I am not.

16 Q. Are you familiar with a study by Chen entitled
17 "Benzene-Induced Myelodysplastic Syndrome"?

18 A. Yes.

19 Q. And what is the finding of that study?

20 A. I believe there was a representation on the part
21 of the authors that there was an increased risk of
22 myelodysplastic syndrome among people who they categorized
23 as occupationally exposed to benzene.

24 Q. And do you recall whether that finding was
25 significant?

1 MR. AMUNDSON: Objection; vague as to the term
2 "significant."

3 THE WITNESS: I was about to ask you: Did you
4 mean statistically significant?

5 BY MR. METZGER:

6 Q. Yes. I make it a practice of whenever using the
7 word significant to refer to statistical significance, and
8 when I'm not talking about that, I use the word
9 "important."

10 A. Okay. I'm glad to know that. So I can assume
11 that to be true?

12 Q. Yes. I'm talking about statistically
13 significant.

14 A. I don't recall whether it was statistically
15 significant or not.

16 Q. Are you familiar with a study by Ciccone,
17 C-i-c-c-o-n-e, regarding benzene exposure and
18 myelodysplastic syndrome?

19 A. No.

20 Q. Are you familiar with a study by Cowles,
21 C-o-w-l-e-s, regarding benzene exposure and
22 myelodysplastic syndrome?

23 A. Yeah. I think that's -- I think her name is
24 pronounced Cowles. And that is Sally Cowles or S. Cowles.
25 Correct?

1 Q. Yes.

2 A. Yes, I am.

3 Q. And what was the finding of that study?

4 A. I don't recall the finding.

5 Q. Are you familiar with a study by Crane regarding

6 benzene exposure and myelodysplastic syndrome?

7 A. No.

8 Q. Are you familiar with a study by Cuneo on that

9 topic?

10 A. Could you spell that.

11 Q. C-u-n-e-o.

12 A. Yes.

13 Q. What was the finding of that study?

14 A. I don't recall the finding.

15 Q. Are you familiar with a study by Farquhar,

16 F-a-r-q-u-h-a-r, regarding benzene and MDS?

17 A. No.

18 Q. Are you familiar with a study by Farrow regarding

19 benzene and MDS?

20 A. No.

21 Q. Are you familiar with study by Fengtong on that

22 topic?

23 A. No.

24 Q. Are you familiar with a study by Goguel,

25 G-o-g-u-e-l, on that topic?

1 A. Could you read me the title of that paper.

2 Q. Sure. "Les Leucemies Benzeniques de la Region
3 Parisienne entre 1950 et 1965."

4 A. Yeah. I am familiar with that paper. My
5 recollection is that it is a study of the epidemiology of
6 leukemia. I don't recall anything in it about MDS.

7 Q. Incidentally, was MDS previously referred to as
8 preleukemia?

9 A. It's possible that some people made that
10 equivalence of terminology. It was never a standard or
11 accepted terminology.

12 Q. Well, before myelodysplastic syndrome was called
13 that, what was it called by clinicians?

14 A. Dismyelopoiesis.

15 Q. And have you read any articles where it had been
16 previously referred to as preleukemia?

17 A. I have certainly read literature on the condition
18 known as preleukemia. I have never seen an epidemiologic
19 study of it. Whether or not some individual cases of what
20 was to come to be known as MDS were labeled preleukemia or
21 not, I don't know. But I do not understand the word
22 preleukemia as it was ordinarily used to be the equivalent
23 of what we now call any of the MDS's.

24 Q. Are you familiar with a study by Goldberg
25 regarding benzene and MDS?

1 A. Again, could I ask for the title of that?

2 Q. "Survey of Exposure to Genotoxic Agents in
3 Primary Myelodysplastic Syndrome."

4 A. No.

5 Q. Are you familiar with a study by the Groupe
6 Francais regarding benzene and myelodysplastic syndrome?

7 A. No.

8 Q. Are you familiar with a study by Hasselbalch
9 regarding benzene and MDS?

10 A. No.

11 Q. You mentioned this study by Hayes. Which study
12 by Hayes are you referring to?

13 A. I'm not sure how to identify it to you. I'd have
14 to see it. As far as I know, there was only one in which
15 myelodysplastic syndrome or similar language was
16 incorporated into the title.

17 Q. Okay. Are you familiar with a study by Hotz,
18 H-o-t-z, regarding benzene and MDS?

19 A. No.

20 Q. Are you familiar with a study by Ido, I-d-o,
21 regarding the epidemiology of MDS?

22 A. No.

23 Q. Are you familiar with a study by Iurlo,
24 I-u-r-l-o, regarding MDS?

25 A. No.

1 Q. Are you familiar with a study by Jacobs
2 concerning MDS?

3 A. No.

4 Q. Are you familiar with a study by Kim concerning
5 benzene and MDS?

6 A. No.

7 Q. Are you familiar with an article by Levine
8 entitled "Secondary Myelodysplastic Syndromes and
9 Leukaemias"?

10 A. No.

11 Q. Are you familiar with a study by Levine entitled
12 "Leukemias and Myelodysplastic Syndromes Secondary to
13 Drug, Radiation, and Environmental Exposures"?

14 A. No.

15 Q. Are you familiar with a study by Mecucci
16 regarding benzene and MDS?

17 A. No.

18 Q. Are you familiar with a study by Mele, M-e-l-e,
19 regarding leukemia and preleukemia?

20 A. Yes.

21 Q. What is that study?

22 A. I have read the paper, but I don't recall it.

23 Q. Are you familiar with a study by Mironova
24 regarding --

25 A. The answer is no.

1 Q. Are you familiar with a study by Nagata?
2 A. No.
3 Q. Are you familiar with a study by Nisse,
4 N-i-s-s-e?
5 A. No.
6 Q. Are you familiar with a study by Preudhomme
7 regarding MDS?
8 A. No.
9 Q. Are you familiar with a study by Rigolin
10 regarding MDS?
11 A. No.
12 Q. Are you familiar with a study by Rodella
13 regarding MDS?
14 A. No.
15 Q. Are you familiar with a study by Rozman
16 concerning MDS?
17 A. The first name?
18 Q. R-o-z-m-a-n.
19 A. No.
20 Q. Are you familiar with a study by Ruiz, R-u-i-z,
21 regarding benzene and MDS?
22 A. No.
23 Q. Are you familiar with a study by Sandler
24 regarding MDS?
25 A. Yes.

1 Q. Which study?

2 A. I think I have it here.

3 MR. AMUNDSON: You've identified it already.

4 MR. METZGER: Oh, we have.

5 Q. And are you familiar with any other studies by

6 Sandler concerning MDS?

7 A. No.

8 Q. Are you familiar with a study by Sanz, S-a-n-z,

9 regarding MDS?

10 A. No.

11 Q. Are you familiar with a study by Shen, S-h-e-n,

12 regarding MDS?

13 A. No.

14 Q. Are you familiar with a study by Smith regarding

15 benzene exposure and risk of MDS?

16 A. No.

17 Q. Are you familiar with a study by Snyder entitled

18 "Benzene and Leukemia" which concerns MDS?

19 A. No.

20 Q. Are you familiar with a study by Stafford

21 concerning MDS?

22 A. No.

23 Q. Are you familiar with a study by Stillman

24 concerning MDS?

25 A. No.

1 Q. Are you familiar with a study by Tasaka regarding
2 MDS?

3 A. Could you read the title of that.

4 Q. Sure. "Translocation 3;21, et cetera, Found in a
5 Patient with Myelodysplastic Syndrome and Long-term
6 Exposure to Organic Solvents."

7 A. Yeah. I have read that paper.

8 Q. And what is the finding of that paper?

9 A. I think it's essentially captured in the title.
10 It's an individual case report. It's not an epidemiologic
11 study.

12 Q. Are you familiar with a study by Travis
13 concerning benzene and MDS?

14 A. Again could you read the title.

15 Q. "Hematopoietic Malignancies and Related Disorders
16 Among Benzene-exposed Workers in China."

17 A. Yes.

18 Q. What is the finding of that study?

19 A. I don't recall.

20 Q. Are you familiar with a study by Tsai, T-s-a-i,
21 regarding benzene and MDS?

22 A. May I ask if the author's first initial is S?

23 Q. Yes, it is.

24 A. Yes, I am familiar with that paper.

25 Q. What's the finding of that study?

1 A. I don't recall.

2 Q. Are you familiar with a study by Van den Berghe
3 regarding MDS?

4 A. No.

5 Q. Are you familiar with a study by Vineis regarding
6 MDS?

7 A. Could you read me the title of that paper. I
8 think it might be here.

9 MR. AMUNDSON: I think we identified that.

10 THE WITNESS: No. That was the paper that I
11 brought. Maybe read me the title.

12 BY MR. METZGER:

13 Q. Yes, I will. "Cytogenetics and Occupational
14 Exposure to Solvents: A Pilot Study on Leukemias and
15 Myelodysplastic Disorders."

16 A. I am familiar with that paper.

17 Q. What is the finding?

18 A. I don't recall the finding.

19 Q. Are you familiar with a study by Vineis entitled
20 "Solvent Exposure and Myelodysplastic Syndrome"?

21 A. Well, now that you read them in immediate
22 juxtaposition, I can't say which one I am familiar with.
23 I am not familiar with two.

24 Q. Are you familiar with any studies by West
25 regarding benzene and MDS?

1 A. No.

2 Q. Are you familiar with any studies by -- I think
3 you did mention a study by Y-i-n.

4 A. Yin, Y-i-n.

5 Q. You did?

6 A. Yes. I just want to be clear that it's Yin, not
7 Yen. I think you said Yen.

8 Q. It is Yin, Y-i-n.

9 A. Okay.

10 Q. How many studies by Yin are you familiar with
11 regarding benzene and MDS?

12 A. Just one.

13 Q. And what is the finding of that study?

14 A. It represents that among the group of people
15 described as benzene workers that there was an excess of
16 MDS.

17 Q. Was the excess statistically significant?

18 A. I don't recall.

19 Q. Do you recall how many cases of MDS there were in
20 this study?

21 A. No.

22 Q. Are you familiar with a study by Zapata Gayon?

23 A. No.

24 Q. Are you familiar with any studies that have found
25 a statistically significant excess of AML in truck

1 drivers?

2 A. I believe there is -- well, I don't know if it's
3 statistically significant, so I guess I'll have to answer
4 the question in the negative.

5 Q. Are you aware of any studies finding a
6 statistically significant excess of AML in workers exposed
7 to diesel exhaust?

8 A. No.

9 Q. Are you familiar with any studies finding a
10 statistically significant excess of MDS in truck drivers?

11 A. No.

12 Q. Are you familiar with any studies finding a
13 statistically significant excess of MDS in diesel exhaust
14 exposed workers?

15 A. No.

16 Q. Are you familiar with any studies finding a
17 statistically significant excess of MDS among smokers?

18 A. No.

19 Q. Incidentally, we've been referring to the term
20 "statistically significant," and we haven't actually
21 defined that. But in answering my questions have you
22 assumed that I've meant statistically significant to a 95
23 percent confidence interval?

24 A. I have understood that.

25 Q. Yes. And would you agree that that is the

1 convention for statistical significance, although it is --
2 well, that much?

3 A. Mr. Metzger, I'm not sure if you're going
4 someplace with this or not, so I just want to say that it
5 is not statistically significant to the 95 percent
6 confidence interval. It is statistically significant at
7 the 5 percent level of probability. I just don't know if
8 we're going down that road or not.

9 Q. Yes. I misstated it. Being a lawyer rather than
10 an epidemiologist, I tend to lapse.

11 A. It really doesn't matter if that's the end of the
12 story. It does matter if we go down to what's the meaning
13 of confidence interval relative to the meaning of P
14 values.

15 Q. That's a venture I don't want to embark on right
16 now.

17 A. Neither do I.

18 Q. Okay. Are you aware of any mechanistic studies
19 regarding benzene which indicate that as benzene -- as the
20 concentration of benzene increases to about 25 parts per
21 million, there is an increasingly lesser production of
22 toxic metabolites?

23 A. I am not aware of any such study. Well, let me
24 ask for some clarification. When you say a "lesser
25 production," do you mean in a relative or an absolute

1 sense?

2 Q. I mean a relative sense.

3 A. Okay.

4 Q. And absolute as a matter of fact.

5 A. Yeah. My question back was not well phrased. I
6 should have asked whether or not it is relative, and
7 you've answered that.

8 My answer is that there is one paper that
9 addresses this issue. I don't recall exactly where the
10 cut point was. I think it might have been 20, but perhaps
11 it was two alternatives, one being 20, one being 40. So
12 yes, there is at least one paper that deals with the human
13 situation and the relationship between low level exposures
14 and possible proximate leukemogens or toxins, if you wish.

15 Q. Do you know what the toxic metabolites of benzene
16 are?

17 A. I think there are several, but I assume that the
18 one we're interested in is the proximate leukemogen.

19 Q. Well, let me just ask you: What are the toxic
20 metabolites of benzene?

21 A. I don't know.

22 Q. Do you have an opinion as to whether benzene
23 causes AML by inducing genetic damage?

24 MR. AMUNDSON: I'll object. That question is
25 vague as to the term "genetic damage." Go ahead.

1 THE WITNESS: Yes. I need to ask for a question
2 of clarification. In your question do you mean benzene
3 per say, or do you mean benzene or one of its metabolites?

4 BY MR. METZGER:

5 Q. Benzene and/or one or more of its metabolites.

6 A. The answer to your question, then, is yes.

7 Q. I forgot what my question was.

8 A. I think it related to whether or not benzene or
9 one of its metabolites causes AML by genetic damage.

10 Q. Okay. Are you able to identify for me the genes
11 which benzene damages in causing AML?

12 A. No.

13 Q. Are you able to identify for me the chromosomes
14 which benzene damages in causing AML?

15 A. No.

16 Q. You also had an opinion regarding the possible
17 role of smoking in Mr. Workman's condition. And what is
18 your opinion on that topic?

19 A. I believe that there would be no way to
20 substantiate the point of view that Mr. Workman's
21 cigarette smoking would have produced his case of MDS.

22 Q. Okay. I think I've gone through the general
23 topics that you provided me earlier. As we've gone
24 through this process, have any additional topics or
25 opinions occurred to you that you have not yet related to

1 me?

2 A. No. But I understand that it's possible that I
3 may be asked to read some other depositions, whether of
4 plaintiffs or defense witnesses, and so it is possible
5 that other opinions will come to mind.

6 There is one area that you did not ask me about,
7 which is important to my point of view. And there is
8 another area that you asked me about but from a
9 perspective that did not allow me to state a fundamental
10 part of the basis for my point of view.

11 Q. Please tell me.

12 A. Both?

13 Q. Both, sure. I'm here to find out your opinions,
14 so whichever you'd like to tell me.

15 A. The first one is that the probability of
16 conversion during the remaining lifetime of a case of MDS
17 to blast crisis and/or possibly AML if you wish to call it
18 that is known, and it differs among the different types of
19 MDS. And it is the lowest and is, in fact, rather low for
20 refractory anemia with ringed sideroblasts. So I would
21 just state that to be a fact and -- I don't know how to do
22 this. Do I show how I documented, or do I wait to see if
23 you ask me about that?

24 MR. AMUNDSON: Let him ask that.

25 MR. METZGER: I will.

1 Q. But before you do that, you have another matter
2 you want to tell me about. So why don't you tell me about
3 that.

4 A. The occupational exposure to diesel exhaust and
5 which usually also entails occupational exposure to diesel
6 fuel is something that has been studied in large number of
7 rather large studies with interest in a full array of end
8 points by which I mean all causes of death or at least all
9 major causes. These studies have had a focus on cancer of
10 the lung and to a lesser extent -- well, to a much lesser
11 extent cancer of the bladder. I know of none of these
12 major studies of the adverse effects of high level
13 exposures to diesel exhaust that show an excess of any
14 form of leukemia.

15 I will mention, incidentally, that they don't
16 show MDS either, but they don't actually present data on
17 that condition. Therefore, a large part of the
18 justification for my point of view that Mr. Workman's
19 exposure to diesel exhaust and/or fuel did not produce his
20 MDS is based not on a reverse view of what is known about
21 the causes of MDS but, rather, on the forward view of what
22 is known about the adverse health effects of diesel
23 exhaust. So those are my two points.

24 Q. Of the studies that you're referring to regarding
25 diesel exhaust, how many of those studies had sufficient

1 statistical power to detect a significant increase in AML?

2 A. Mr. Metzger, this question will need a little
3 work before I can answer it. May I try to suggest how we
4 could improve the question?

5 Q. Well, let me just ask you if you can tell me the
6 number of those studies --

7 A. No, I can't.

8 MR. AMUNDSON: Let him finish the whole question.
9 I think you interrupted him.

10 THE WITNESS: Sorry.

11 BY MR. METZGER:

12 Q. Let me ask another question, and we'll explore a
13 little further. How many of the studies, epidemiologic
14 studies that you are referring to regarding diesel exhaust
15 have sufficient statistical power to detect a significant
16 increase of MDS?

17 A. Again, I cannot answer the question as phrased.

18 Q. Okay. How would you like me to change the
19 question? Not that I'm going to, but I'll entertain the
20 suggestion.

21 A. I think the question would be converted into
22 something I could answer if you change two parts of it.
23 One was that instead of talking about statistical power,
24 you asked the more general question about the precision of
25 the study. I consider and I think most epidemiologists

1 consider power to be what is termed a front end or
2 planning issue in study design, and its counterpart is
3 precision of the width of the confidence interval at the
4 analytical or interpretative phase.

5 The second thing is you asked to detect a
6 significant, and I understand you to mean statistically
7 significant, excess if there were in truth to be one. It
8 is not possible to answer that question without stating
9 what level of precision -- I'm sorry -- what level of
10 significance would have been requested.

11 There is another point, and that is that when you
12 ask a question of this sort -- I'll reverse that. When I
13 would try to answer a question of this sort, and when it
14 is phrased in the context of an array of studies, the
15 question of the precision and statistical significance of
16 any one individual study becomes a relatively minor issue.
17 The issue becomes whether or not an array of studies which
18 are in at least some sense comparable in terms of the
19 cause-effect relationship that they seek are displaying
20 similar or dissimilar results.

21 Q. Have you conducted any meta-analyses of these
22 studies?

23 A. No.

24 Q. What is the frequency that you believe exists for
25 the conversion of RARS to AML?

1 A. With the understanding that I will accept your
2 language of conversion of AML to be the equivalent to my
3 language of blast crisis, the answer is 5 percent.

4 Q. Now, of that 5 percent, what percentage of those
5 are --

6 A. I'm sorry. I need to ask if we could go back to
7 that question. Could I have it read back because I might
8 have missed an important part. I might have assumed
9 something in my answer that you didn't actually say.

10 Q. Why don't you just clarify.

11 A. I was speaking about the conversion to blast
12 crisis or AML, if you will, wish, in that form of MDS that
13 Mr. Workman had.

14 Q. Which is RARS?

15 A. Yes. Is that what you asked?

16 Q. I said specifically RARS.

17 A. I lost that.

18 Q. That's fine. Now, my next question is:
19 Regarding that approximately 5 percent that you
20 mentioned -- first of all, what is the source of your
21 information for that? Is that Wintrobe?

22 A. Yes.

23 Q. And of that approximately 5 percent, what
24 percentage of those were cases of patients who had been
25 administered alkylating agents?

1 A. I don't know.

2 Q. Are you aware of the frequency or percentage of
3 conversion of RARS to blast crisis or AML for patients who
4 have RARS as a result of alkylating agent therapy?

5 MR. AMUNDSON: Objection; compound, vague and
6 ambiguous, overbroad. If you understand it, go ahead and
7 answer it.

8 THE WITNESS: My answer is no.

9 BY MR. METZGER:

10 Q. Okay. Let me just review. I think we're done.

11 MR. AMUNDSON: Give me two minutes.

12 BY MR. METZGER:

13 Q. What I'd like to do is we're all going to take a
14 short break for various reasons. And during that break I
15 would like you to think, Dr. Cole, if you've told me now
16 all of the opinions that you have in this case, and we'll
17 come back in a short minute and find out if there's
18 anything left.

19 A. So when we resume, you would like me to tell you
20 whether or not I feel that I have been asked about all of
21 my opinions?

22 Q. Whether you've related them to me, whether I've
23 asked you about them or not.

24 A. Well, if you've asked them, I've related them.

25 (A recess was taken.)

1 BY MR. METZGER:

2 Q. Dr. Cole, I have asked you before the break if
3 you would think during the break if there are any other
4 opinions that you have that you have not yet related to
5 me.

6 A. No.

7 Q. Okay. I have a few more questions for you, and
8 we'll be done shortly. You have told me that -- I'm not
9 sure I'm stating this accurately, but that it's your
10 opinion that benzene does not cause MDS. First of all, am
11 I right on that, that that's your opinion?

12 A. No. It would have depended on the context how I
13 would have phrased it. But since you ask it pointedly,
14 then I have to say no.

15 Q. Well, what is your opinion on that topic?

16 A. My opinion is that the available scientific
17 evidence does not support a causal relationship between
18 benzene and MDS, but I used the expression for shorthand
19 that you originally said.

20 Q. Fine. Now, when you are referring to the
21 available evidence, what types of evidence or data are you
22 referring to?

23 A. Now, when you say when I refer to the available
24 evidence, you mean as in the statement that I made just
25 prior to this question when I said the available

1 scientific evidence?

2 Q. Yes.

3 A. I refer to the totality of it but, of course,
4 with my primary emphasis being on the epidemiology.

5 Q. Okay. So at least part of it is epidemiology?

6 A. No. It's not part of it. It's the overwhelming
7 majority of it, and it is in the tradition of the original
8 rules of the IARC for putting the substance into
9 Category 1.

10 Q. Other than epidemiology, what types of studies
11 are you relying on for your opinion on that point?

12 A. And when you say "on that point," you mean --

13 Q. Benzene causing --

14 A. -- benzene as a cause of MDS?

15 Q. Yes.

16 A. I also rely upon my understanding of the animal
17 studies.

18 Q. Anything else?

19 A. No.

20 Q. Okay. What animal studies are there that you
21 have reviewed regarding benzene and MDS?

22 A. I haven't reviewed any.

23 Q. What animal studies regarding benzene and MDS
24 have you considered in forming your opinion?

25 A. It is not that I have considered any particular

1 study or studies, but it is my understanding that the
2 available animal evidence does not establish benzene as a
3 cause of MDS in animals.

4 Q. And what is your -- upon what are you basing
5 that?

6 A. Well, I can't give you a citation. That's just
7 my recollection at this time.

8 Q. Have you ever attempted to ascertain what animal
9 studies there are that report morphological features which
10 are those of MDS?

11 A. No, I haven't attempted to do that.

12 Q. Now, you mentioned IARC. Has IARC ever published
13 a monograph regarding benzene and MDS?

14 A. You mean a monograph that was specifically given
15 over to that question?

16 Q. Well, let me rephrase. IARC publishes monographs
17 regarding the carcinogenicity of chemicals. True?

18 A. Yes.

19 Q. And the last IARC assessment of benzene was what?
20 1989? Is that right?

21 A. I really don't recall. For the purpose of this
22 case, I didn't check benzene. I checked diesel exhaust.

23 Q. In any of the IARC evaluations of the
24 carcinogenicity of chemicals, does IARC look at MDS as a
25 carcinogenic end point?

1 A. No. I would assume not. Let me just be clear.
2 You asked me whether or not I considered benzene to be a
3 cause of MDS, and I said that if I were to use the
4 criteria that were the old criteria of IARC, basically
5 what I was trying to say there was that the available
6 evidence in human beings is sufficient, then the answer
7 would be no.

8 Q. And what specific criteria are you referring to?

9 A. Let me be sure I understand the question. You
10 mean what are the IARC's criteria?

11 Q. I can read the IARC criteria, and I have. You've
12 already told me that IARC does not evaluate MDS as a
13 carcinogenic end point. So my question is: What criteria
14 are you employing for your assessment that you've
15 provided?

16 A. I understand that now. I think there was a
17 little confusion generated by perhaps the way I phrased my
18 answer. As you know, apparently, IARC begins with two
19 subclassifications of the evidence, one for humans and one
20 for animals. And then from those two subclassifications,
21 it attempts to derive an overall categorization. What I
22 was trying to say when I referred to IARC was that that's
23 what I would do for benzene and MDS. That is I would ask
24 whether or not the human evidence is sufficient, limited,
25 or inadequate, and I would ask whether the animal evidence

1 is sufficient, limited, or adequate. And I would require
2 before I could make a causal inference that the human
3 evidence is sufficient. And as I would have understood it
4 in this case, it wasn't for benzene but for diesel
5 exhaust. However, I say the same thing for benzene.

6 Q. Well, you're referring to IARC Group 1. Correct?
7 When you say sufficient, sufficient evidence of
8 carcinogenicity in humans, you're referring to a Group 1
9 designation by IARC. True?

10 A. Yes. I think that is equivalent.

11 Q. And the standard that IARC employs for making
12 that determination is in part, in large part epidemiology.
13 True?

14 A. There are two answers to that question or two
15 parts to an answer. It used to be true that the judgment
16 with regard to the human evidence had to be in the
17 category sufficient for an agent to fall into the overall
18 judgment of Group 1 sufficient evidence. However, about
19 five or six years ago, they modified that criterion and
20 have relaxed it somewhat. I consider that inappropriate,
21 and I have not made that relaxation.

22 Q. Fine. Under the former unrelaxed standard, what
23 was the standard as you understand it for epidemiologic
24 evidence for IARC to deem a chemical carcinogenic to
25 humans?

1 A. In the final analysis, it became -- it could be
2 defined as a majority vote of the working group.

3 Q. Okay. And the working group evaluated the
4 evidence using certain criteria, however. True?

5 A. Well, those criteria -- the answer to your
6 question is yes, that's true.

7 Q. And are those the same criteria that you're using
8 for your assessment or not?

9 A. Well, now I have to develop the answer because --

10 Q. Please.

11 A. -- each individual member of the working group
12 was allowed to reach his or her own judgment in his or her
13 own way. And beyond that the criteria were rather general
14 and basically were of the type that there is a reasonably
15 consistent, often statistically significant, unlikely to
16 be confounded set of findings. And whether or not those
17 criteria were fulfilled was a judgment that each member of
18 the working group made, and I'm saying that that's a
19 judgment that I also attempt to make.

20 Q. And your personal judgment on this issue based
21 upon what you've reviewed is that there's insufficient
22 evidence for benzene and MDS?

23 A. And for diesel and MDS.

24 Q. Okay.

25 A. And for diesel and AML.

1 Q. Okay. Now, it's true at that IARC has never
2 evaluated benzene and MDS employing those criteria. True?

3 A. I can't affirm that that is true, but I would say
4 that it is probably true, and I don't know that they have.
5 I wouldn't think they would because MDS is not a cancer.

6 Q. Right. Okay. Now, lastly, we did get faxed or
7 e-mailed here a list of cases. I'd like you to take a
8 look at that.

9 A. You said last week?

10 Q. Lastly.

11 A. Lastly. Sorry.

12 Q. And my question is: Which of those cases there
13 concern benzene?

14 A. All right. I'll need a minute to go through
15 this.

16 Q. Sure.

17 A. (Witness peruses document.)

18 If we look at No. 9, this is actually a diesel
19 exposure and to some extent benzene. However, the outcome
20 is not an LHC.

21 13, there are actually two cases there. The
22 Martindale case is a benzene, and as I recall it was CLL,
23 and the Casas is benzene and pancreas cancer.

24 Q. What was the name of the attorney who took your
25 deposition in those cases?

1 A. In Casas and Martindale?

2 Q. Yes.

3 A. I don't recall his name.

4 Q. Was it Herschell Hobson, by chance?

5 A. No.

6 Q. Have you been deposed by Herschell Hobson?

7 A. On one occasion many, many years ago. This
8 person may have been in his firm, but I don't recall his
9 name. Let me just finish.

10 Q. Go ahead.

11 A. Well, 14, of course, is the same. 15 is a case
12 in which benzene is one of the issues. Is this your copy
13 or mine?

14 Q. Yes, it was. Thank you. Okay. We are done.
15 Same stip?

16 MR. AMUNDSON: Same stip.

17 MR. METZGER: Based upon the prior agreement that
18 I believe we reached regarding signing and execution and
19 changes, the stipulation will be that the court reporter
20 may forward the original transcript directly to Dr. Cole;
21 and that he may have until February 19th to make the
22 corrections that he wishes and sign the transcript and
23 notify me of the changes, notify all counsel of the
24 changes; he will then forward the original transcript to
25 me, and I will keep it; if the original is not signed or

1 is somehow lost, a certified copy may be used with full
2 force and effect.

3 (Discussion held off the record.)

4 MR. AMUNDSON: So we're amending the stipulation
5 that in the event Dr. Cole does not receive the
6 deposition, the original transcript until after the 19th,
7 or even if it's a day before the 19th, we will give him an
8 extra week from his receipt to sign it. And then we'll
9 make our best efforts to immediately notify all parties.

10 MR. METZGER: He's notifying us.

11 MR. AMUNDSON: He will notify us, or perhaps
12 he'll ask us to notify you. But you will get notification
13 either way --

14 MR. METZGER: Right.

15 MR. AMUNDSON: -- within a week of his receipt.

16 THE REPORTER: Do you want a copy, Counsel?

17 MR. AMUNDSON: Yes, please.

18 (Proceedings concluded at 2:23 p.m.)

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DECLARATION

I hereby declare I am the deponent in the within matter; that I have read the foregoing deposition and know the contents thereof, and I declare that the same is true of my knowledge except as to the matters which are therein stated upon my information or belief, and as to those matters, I believe it to be true.

I declare under the penalties of perjury of the State of California that the foregoing is true and correct.

Executed this day of ,
200 , at , California.

PHILIP COLE, M.D.

