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COMMONWEALTH OF PENNSYLVANIA

1136

RECEIVED

PUBLIC UTILITY COMMISSION

NOV 26 1991

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 Letter of Notification of :  
Philadelphia Electric Company :  
 relative to reconstructing and :  
 rebuilding of the existing 138 kV :  
 line to operate as a Woodbourne- :  
 Heaton 230 kV line in Montgomery and :  
 Bucks Counties. :  
 :  
 Further hearing. :  
 :  
 -----x

SECRETARY'S OFFICE  
 Public Utility Commission  
 Docket No.  
 A-110550F055

Pages 1136 through 1207 Hearing Room No. ORIGINAL  
 State Office Building  
 Broad and Spring Garden Streets  
 Philadelphia, Pennsylvania

Friday, November 22, 1991

Met, pursuant to adjournment, at 10:00 a.m.

BEFORE:

HERBERT SMOLEN, Administrative Law Judge

SECRETED  
 DEC 04 1991

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C O N T E N T S

<u>WITNESSES</u>	<u>DIRECT</u>	<u>CROSS</u>	<u>REDIRECT</u>	<u>RECROSS</u>
Edward Paul Gelmann				
By Mr. Watson	1139	---	---	---
By Ms. McCloskey		1141	---	---
By Mr. Sugarman		1163	---	---

E X H I B I T S

<u>NUMBER</u>	<u>FOR IDENTIFICATION</u>	<u>IN EVIDENCE</u>
<u>Philadelphia Electric Company</u>		
Rebuttal Statement No. 3 (Gelman)	1139	---

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## P R O C E E D I N G S

1  
2 ADMINISTRATIVE LAW JUDGE HERBERT SMOLEN: This is a  
3 further hearing in the matter of the letter of  
4 notification in connection with the Philadelphia Electric  
5 Company's Woodbourne-Heaton line.

6 This morning's session has been reserved for the  
7 presentation of the rebuttal testimony of Dr. Edward  
8 Gelmann.

9 Are we ready to proceed?

10 MR. BONNEY: Yes, Your Honor. We have one  
11 preliminary. During the cross-examination of Dr. Bockman  
12 the other day, the Office of Consumer Advocate made an on  
13 the record data request for full citations to four  
14 scientific studies that were discussed in Dr. Bockman's  
15 testimony. We just wanted to indicate that we have  
16 provided the response to the on the record data request.

17 JUDGE SMOLEN: That is noted for the record.

18 Now, let's proceed.

19 MR. WATSON: Your Honor, we call Dr. Edward  
20 Gelmann.

21 Whereupon,

22 EDWARD PAUL GELMANN

23 having been duly sworn, testified as follows:

24 JUDGE SMOLEN: Please have a seat. Keep your voice  
25 up. State your full name and business address for the

1 record.

2 THE WITNESS: Edward Paul Gelmann, 3800 Reservoir  
3 Road, N.W., Washington, DC 20007.

4 JUDGE SMOLEN: Mr. Watson.

5 DIRECT EXAMINATION

6 BY MR. WATSON:

7 Q. Dr. Gelmann, do you have before you a copy of a  
8 document entitled Rebuttal Testimony of Dr. Edward  
9 Gelmann, Philadelphia Electric Company Rebuttal Statement  
10 No. 3?

11 A. I do.

12 (Whereupon, the document was marked  
13 as PECO Rebuttal Statement No. 3  
for identification.)

14 BY MR. WATSON:

15 Q. Do you have any corrections or additions to  
16 that document?

17 A. On page 15, line 14, a transcription error, the  
18 second "that" should be a "to", t-o.

19 JUDGE SMOLEN: What page was that again?

20 THE WITNESS: Page 15, line 14. Indicate "that  
21 exposure to power".

22 JUDGE SMOLEN: Yes, I found that one myself.

23 Go ahead.

24 BY MR. WATSON:

25 Q. Let me ask you to take a look at page 16, line

1 27. Could you read that sentence and see if there is  
2 anything missing from that?

3 (Witness perusing document.)

4 JUDGE SMOLEN: Are you saying "have" to "has"?

5 MR. WATSON: Yes. I think that, or "the effects  
6 have", one or the other. It seems like there is  
7 something left out.

8 JUDGE SMOLEN: I think the verb modifies her work.  
9 Is that what it meant to be?

10 THE WITNESS: "Either the effects of the  
11 phenomenon" have not been shown to occur in humans.

12 JUDGE SMOLEN: Do you want to follow it up?

13 BY MR. WATSON:

14 Q. Which one would you like?

15 A. "The effect" is fine.

16 Q. "The effect"?

17 A. Yes.

18 Q. The effects have not been shown to occur in  
19 humans?

20 A. In humans.

21 Q. So after "and" add "the effects", is that  
22 correct, Dr. Gelmann?

23 A. Yes.

24 Q. Any other additions or corrections?

25 A. No, sir.

1 Q. Dr. Gelmann, if I were to ask you while you are  
2 on the stand under oath here today each of the questions  
3 set forth in this document entitled Rebuttal Testimony of  
4 Dr. Edward Gelmann, would your answers be as set forth  
5 therein?

6 A. They would.

7 MR. WATSON: Your Honor, I pass the witness.

8 JUDGE SMOLEN: Ms. Burket.

9 MS. BURKET: No questions, Your Honor.

10 JUDGE SMOLEN: Mr. Dillon.

11 MR. DILLON: No questions, Your Honor.

12 JUDGE SMOLEN: Ms. McCloskey.

13 MS. McCLOSKEY: Thank you, Your Honor.

14 CROSS-EXAMINATION

15 BY MS. McCLOSKEY:

16 Q. Good morning, Dr. Gelmann. My name is Tanya  
17 McCloskey and I represent the Office of Consumer  
18 Advocate.

19 Dr. Gelmann, am I correct that you are testifying  
20 here today as a specialist in medical oncology and  
21 molecular and cellular biology?

22 A. Yes.

23 Q. And you are not testifying as an expert in  
24 electromagnetic fields or toxicology or epidemiology,  
25 correct?

1           A. Well, my testimony pertains to the studies that  
2 had to do with electromagnetic fields and cells and  
3 organisms. But I am not a physicist, I am not a  
4 toxicologist or an epidemiologist.

5           Q. Thank you.

6           Would you agree that the purpose of your testimony  
7 was limited to a review of the literature as to molecular  
8 and cellular biology as it relates to the question of  
9 electromagnetic fields and cancer?

10          A. The literature I have read is much broader than  
11 that, but my opinions are restricted to oncology and  
12 molecular and cellular biology.

13          Q. Now, if you could refer to page 22 of your  
14 testimony, lines 24 through 27, you refer at the end of  
15 line 27 to other adverse health effects. Do you see that  
16 reference?

17          A. Yes.

18          Q. And what are you referring to when you state  
19 other adverse health effects?

20          A. Any health effects that I would know of as a  
21 medical oncologist, also being an internist, that is,  
22 having specialty boards in internal medicine.

23          Q. And can you give us some example of those that  
24 you are considering as other health effects within your  
25 specialty?

1 A. I'm not sure what you are asking.

2 Q. What I am looking for is a specific type of  
3 health effect that you're discussing there. For example,  
4 you discuss cancer as one health effect. Is there  
5 anything else you are referring to when you use the term  
6 other adverse health effects?

7 A. Potentially any health effects that would fall  
8 under the general topic of internal medicine. There is  
9 particular interest in the lay press about cancer  
10 causation and there have been some epidemiologic papers  
11 that have addressed cancer, so cancer is the main topic.

12 I am not referring specifically to any other  
13 diseases that I could state. However, my reading of the  
14 literature did not cause me to conclude about any other  
15 possible diseases associated with electromagnetic fields  
16 from power transmission or distribution.

17 Q. As limited, as you stated before, going to the  
18 question of internal medicine and your medical specialty?

19 A. Yes.

20 Q. Dr. Gelmann, would you agree that the question  
21 of the health effects of electromagnetic fields is one  
22 that is interdisciplinary in nature, involving such as  
23 areas as biology, epidemiology, physics and other  
24 specialty areas?

25 A. It depends on what health effects you are

1 talking about.

2 Q. What I am referring to there is the full range  
3 of health effects that may be researched.

4 A. Well, the research that is done depends on the  
5 discipline for which it is pursued.

6 Q. Perhaps we are talking by each other. Let me  
7 try rephrasing it this way: would you agree that the  
8 question of cancer and its relationship to  
9 electromagnetic fields is only one of the areas that is  
10 being explored?

11 A. Well, there have been research papers in other  
12 areas of cancer, yes.

13 Q. And I believe you stated earlier that when you  
14 are looking at electromagnetic fields you are also  
15 looking in the area of physics?

16 A. I'm not sure I said that. I'm not sure what  
17 you mean.

18 Q. Earlier you stated that when one researches an  
19 area of electromagnetic fields you are not just looking  
20 at, for example, a molecule or a cell but you are looking  
21 at an interaction with the electromagnetic field, which  
22 is in the area of physics?

23 A. We must be talking around the question. If one  
24 asks what the effect of an electromagnetic field on a  
25 cell is, then one uses or one may use the electromagnetic

1 field as a stimulus. But one does not have to be a  
2 physicist to do that. But one is certainly using some  
3 sort of a physical entity as opposed to a chemical  
4 entity.

5 Q. Looking at pages four and five of your  
6 testimony, you refer and list numerous articles that you  
7 have published in scientific journals. Have you ever  
8 published any articles about electromagnetic fields and  
9 cancer?

10 A. No.

11 Q. And have you conducted any research into the  
12 question of electromagnetic fields and cancer beyond the  
13 literature review you have conducted for this testimony?

14 A. No.

15 Q. Now, if you could refer back again to pages 21  
16 through 23 of your testimony, and I believe starting on  
17 page 21 you refer to three terms, cause, promote or  
18 contribute to the development of cancer.

19 MR. WATSON: Could we have a line, Counsel?

20 MS. McCLOSKEY: I'm sorry. It was page 22, line  
21 26.

22 BY MS. McCLOSKEY:

23 Q. And first, could you please explain what you  
24 mean by cause?

25 A. What I mean by cause?

1 Q. Yes.

2 A. Cause means cause. That magnetic fields give  
3 -- can give an animal or human cancer.

4 Q. And would that be the same as initiation of  
5 cancer?

6 A. It could mean the same as initiation and it  
7 could also be broader.

8 Q. And in this sentence, though, you are referring  
9 to it in the broader sense, is that correct?

10 A. Yes.

11 Q. When you refer to promote, could you please  
12 tell us what you mean by promote?

13 A. Cancer promotion.

14 Q. What does cancer promotion mean? I am just  
15 trying to get a little background.

16 A. A cancer promoter is an agent that stimulates  
17 cell proliferation.

18 Q. And when you use the term contribute to the  
19 development of cancer, what do you mean by contribute to  
20 the development of cancer?

21 A. Aid in cancer formation in any other way that  
22 you may imagine.

23 Q. Are you familiar with the term progression of  
24 cancer?

25 A. Yes.

1 Q. Is progression different than cause, promote or  
2 contribute, or does it fit into one of those areas?

3 A. It would fit into contribute.

4 Q. Now, turning back to page 21 of your testimony,  
5 line 17, you refer to a substantial body of well done  
6 research. About how many studies are you referring to  
7 when you make that statement?

8 A. Predominantly the ones cited in the testimony.

9 Q. Do you think that more research should be done  
10 in the area of electromagnetic fields and the question of  
11 cancer in your area of cellular and molecular biology?

12 A. That depends what questions you want to ask by  
13 doing the research.

14 Q. What if we wanted to look into the question of  
15 causation of cancer? Do you believe that additional  
16 research should be done in your area concerning the  
17 causation of cancer and E/MF?

18 A. No.

19 Q. What if we want to look at the question of  
20 promotion of cancer and its relationship to E/MFs? In  
21 your area do you think that additional research would be  
22 necessary?

23 A. No.

24 Q. And what if we wanted to look into the area of  
25 contributing to the development of cancer? In your area

1 of specialty do you believe that additional research is  
2 necessary into that question?

3 A. No.

4 Q. Dr. Gelmann, I believe you have answered no to  
5 all three areas. How does the research community in  
6 science determine that enough studies have been conducted  
7 so that additional studies would not be necessary?

8 MR. WATSON: Objection, Your Honor. It's beyond  
9 the scope of his testimony. He is not here as an expert  
10 on the research community of science or the philosophy of  
11 science as it is sometimes called. He is testifying as  
12 an expert in the fields he has indicated.

13 MS. McCLOSKEY: I believe he is testifying as an  
14 expert in the area of cancer research and with that  
15 should have some familiarity with the research community.  
16 I am trying to get behind his --

17 JUDGE SMOLEN: Overruled. You may answer.

18 A. Well, I can't speak for the research community.  
19 I can give you my opinion, and my opinion is that  
20 sometimes research is instigated by important scientific  
21 issues, sometimes by political concerns and sometimes in  
22 response to public concerns that may or may not have any  
23 basis in scientific fact.

24 A good example of that, of the latter, is the  
25 research that was done with the anti-cancer agent

1 Laetrile, which had a tremendous amount of interest among  
2 cancer patients. There were books written about it.  
3 There was a huge amount of hope and money on the part of  
4 the suffering public expended in seeking this agent, in  
5 many cases in an illegal way. And finally the National  
6 Cancer Institute was basically forced by the situation to  
7 sponsor a very rigorous trial which after a lot of public  
8 funds were expended and a lot of cancer patients were  
9 enrolled showed that the stuff was absolutely worthless.  
10 But at least it put the question to rest.

11 BY MS. McCLOSKEY:

12 Q. You have given us one example. Let me ask you  
13 how you have determined as to these areas, what analysis  
14 have you pursued in these three areas to reach your  
15 conclusion that no additional research is necessary?

16 A. The analysis is based on my reading of the  
17 experiments described in the literature.

18 Q. And let's say you read one study that said that  
19 there was some effect shown and you read one study that  
20 said there was no effect shown. How, then, do you  
21 distinguish between those two studies to reach your  
22 conclusion?

23 MR. WATSON: Objection, Your Honor. It assumes  
24 facts not in evidence here. There is obviously more than  
25 one study on each side and there is no way for him to

1 wash out his mind and pretend that there is only one on  
2 each side.

3 MS. McCLOSKEY: Your Honor, it was a hypothetical.

4 JUDGE SMOLEN: It was a hypothetical.

5 MR. WATSON: I don't think it is a hypothetical  
6 that is relevant to the circumstances presented here.

7 JUDGE SMOLEN: My problem with the question, and I  
8 am not ruling on your objection, is that when you say  
9 effects, do you mean adverse effects or do you mean any  
10 effect as part of the question?

11 MS. McCLOSKEY: I can change it to mean adverse  
12 effects. What I am trying to do is get behind what  
13 analysis he is performing.

14 JUDGE SMOLEN: You are withdrawing the former  
15 question. Let's hear the next question.

16 BY MS. McCLOSKEY:

17 Q. Let's assume a hypothetical, Dr. Gelmann, and  
18 state that you have read one study that shows an adverse  
19 effect and you have read another study that shows no  
20 effect. How do you determine between those two studies  
21 or how do you make a distinction between those two  
22 studies and reach a conclusion?

23 MR. WATSON: The same objection.

24 JUDGE SMOLEN: I will overrule it. You can answer.

25 A. It depends on what you mean by effects. You

1 read the scientific data and based on my education,  
2 training and experience I am available to evaluate the  
3 scientific data and I come to conclusions. That is why  
4 we have scientific publications, so that the actual data  
5 is presented before scientific peers and can be assessed.

6 BY MS. McCLOSKEY:

7 Q. Let me ask it a different way, Dr. Gelmann.  
8 When you are evaluating the scientific literature do you  
9 look to see if studies have been replicated?

10 A. Yes.

11 Q. And do you look to see if studies have resulted  
12 in consistent results, or do you look for consistency?

13 A. Yes.

14 Q. And in the area of electromagnetic fields and  
15 cellular and molecular research, how many of these  
16 studies have been replicated?

17 A. I can't answer that sitting here, and I'm not  
18 sure what you mean exactly by replicated. If you want to  
19 discuss specific studies then I am glad to look at them  
20 with you.

21 Q. I'm not trying to get into specific studies. I  
22 am trying to get into you have reviewed many studies that  
23 are presented in your selected references and when you  
24 reviewed those studies did you look to see if there had  
25 been some replication or some consistency in the results

1 that were produced by those studies?

2 A. As part of my reading of the literature, having  
3 read many papers to this point, I am always registering  
4 as I read the next paper what is consistent or in some  
5 way replicative of previous work. But I can't possibly  
6 tell you sitting here, give you a number of how many.  
7 That is beyond what I have in front of me.

8 Q. Now, Dr. Gelmann, again back on page 21, line  
9 17, you refer to the substantial body of well done  
10 research. And in that substantial body of research do  
11 you know how many of those studies were designed to test  
12 whether or not exposure to E/MFs is carcinogenic?

13 A. I cannot tell you which of the studies actually  
14 uses the word carcinogenicity in the body of the text,  
15 and I can't go into the mind of the researchers. But the  
16 studies that I cite as far as I am concerned are relevant  
17 to the question of cancer. And I will assume somewhat  
18 that that crossed the minds of the people who wrote those  
19 papers.

20 Q. Now, I believe I understand from your testimony  
21 that cancer causation is always related to a direct  
22 interaction with DNA. Is that accurate?

23 A. Cancer causation involves a permanent change in  
24 DNA.

25 Q. And do cancer promotion and contribution to the

1 development of cancer always involve a direct interaction  
2 with the DNA?

3 A. No, that is not what I said.

4 Q. Right. I am asking you about the next two  
5 areas.

6 A. I understand that, but that is not what I said.  
7 Cancer promotion is usually caused by an agent which  
8 facilitates cell proliferation.

9 Q. And what about contribution to the development  
10 of cancer?

11 A. That's not a rigorous scientific term. Can you  
12 be more specific.

13 Q. How about the progression of cancer?

14 A. The progression of cancer is something which  
15 can happen all by itself.

16 Q. Now, looking at page ten of your testimony at  
17 line 32, and on to the top of page 11, where you refer to  
18 the Whitson's study on the relationship between DNA  
19 repair and E/MF exposure, do you see that reference?

20 A. Yes.

21 Q. Just to get a starting point, could you attempt  
22 to put in as layman's terms as possible the importance of  
23 the DNA repair mechanism and its relationship to cancer?

24 A. Changes in DNA are necessary for the cause of  
25 cancer. The cell -- a normal cell has the capacity

1 through a number of different enzymatic mechanisms to  
2 repair DNA. There are in-born genetic diseases involving  
3 errors of DNA repair that show the severe impact of  
4 having an abnormal DNA repair on a person. And the study  
5 by Whitson looked at one kind of DNA repair, that is, to  
6 damage caused by ultraviolet radiation and whether power  
7 frequency fields could alter the cell's ability to repair  
8 that damage.

9 Q. And just to be sure we have come full circle,  
10 would it be correct to state that if the damage to the  
11 DNA is not repaired that is when the problem begins, the  
12 change then becomes more permanent? And I don't want to  
13 put words in your word, but I just want to come full  
14 circle.

15 A. Not repairing the DNA results in a mutation.  
16 Whether the mutation is important or not is the next  
17 question.

18 Q. But one of the mutations, then, for example,  
19 could be important in the causation of cancer? Is that  
20 the link?

21 A. Yes.

22 Q. Now, I believe you stated that the Whitson  
23 study looked at one DNA repair process. How many DNA  
24 repair processes do we now know about?

25 A. Many. I can't be more accurate than that.

1 Q. And you note at the top of page 11 the Frazier  
2 study and you use the term another DNA repair process.  
3 Did they look at a different process than in the Whitson  
4 study?

5 A. Yes.

6 Q. Do you know how many repair processes the  
7 Frazier study looked at?

8 A. I don't know how many and I'm not sure that  
9 Frazier knew how many. However, he looked at damage  
10 caused by X-rays, by gamma radiation, which another term  
11 would be ionizing radiation, and looked at the ability of  
12 cells to repair that damage and recover from exposure to  
13 ionizing radiation and found no effect of power frequency  
14 E/MF on the ability of a cell to recover from exposure to  
15 ionizing radiation. Ionizing radiation can cause several  
16 different kinds of breakage in DNA and the experiments  
17 basically encompassed all of those kinds of repairs just  
18 by virtue of the way the experiments were set up.

19 Q. Now, on page 19 of your testimony, at lines 26  
20 and 27, you state that there is no data connecting  
21 melatonin levels to breast cancer in humans. First of  
22 all, what type of research would one rely upon to connect  
23 melatonin levels to breast cancer in humans?

24 A. I don't know. It depends on what research is  
25 done.

1 Q. Let me try and focus the question a little  
2 more. Would that be something along the lines of an  
3 epidemiologic study or something in the area of cellular  
4 and molecular biology?

5 A. Well, studies in a number of scientific  
6 disciplines could be done to ask the question.

7 Q. Now, are you aware of data or research that  
8 connects melatonin levels to breast cancer in animals?

9 A. Which studies are you talking about  
10 specifically?

11 Q. Specifically, are you familiar with a study by  
12 Tamarkin that is entitled Melatonin Inhibition and  
13 Pinealectomy Enhancement of  
14 7,12-Dimethylbenz(a)anthracene-induced Mammary Tumors in  
15 Rats?

16 A. Yes.

17 Q. And would you agree with me that this study  
18 showed that the removal of the pineal glands in the rats  
19 leads to an increase in chemically induced breast cancer  
20 in rats?

21 A. I have read that paper. There were some  
22 effects shown. To discuss specifics I would need you to  
23 give me a copy. I have not read it, reviewed it,  
24 recently. But there were some effects after pinealectomy  
25 and also as a result of administering the melatonin to

1 the rats, as I recall.

2 Q. I unfortunately have not been able to obtain a  
3 copy of that. I have another article that discusses  
4 that. Maybe I can show you that and you can tell me  
5 whether you are comfortable after reviewing that and if  
6 that refreshes your memory.

7 A. No, if we are going to discuss Larry Tamarkin's  
8 paper I would prefer to have his paper in front of me.

9 Q. Why don't I show you the article that I have  
10 and after reviewing it you can let me know.

11 A. Okay.

12 Q. I don't want to discuss it in depth.

13 (Document handed to witness.)

14 MR. WATSON: This is a Stevens paper published in  
15 the American Journal of Epidemiology?

16 MS. McCLOSKEY: Yes.

17 MR. WATSON: This is Steves' discussion of  
18 Tamarkin's paper.

19 MS. McCLOSKEY: Tamarkin's paper. And I am asking  
20 after reviewing that discussion whether that refreshes  
21 his memory sufficiently on the Tamarkin study.

22 MR. WATSON: I thought he said if he was going to  
23 discuss Tamarkin's paper he would want to see Tamarkin's  
24 paper.

25 JUDGE SMOLEN: That is what he said.

1 MS. McCLOSKEY: I'm just asking him if this gives  
2 sufficient detail other than the Tamarkin study.

3 THE WITNESS: I am familiar with this review and it  
4 does not give me sufficient information on the Tamarkin  
5 study.

6 MS. McCLOSKEY: Fine.

7 BY MS. McCLOSKEY:

8 Q. Dr. Gelmann, do you know what the function of  
9 melatonin is in the human body?

10 A. I do not.

11 Q. Now, on page 20 of your testimony you refer to  
12 a study by Ross concerning cell proliferation, and you  
13 state that Ross reported a decreased proliferation of  
14 cells in vitro, is that correct?

15 A. Yes.

16 Q. I have the Ross study with me. Why don't I  
17 just hand that to you.

18 (Witness perusing document.)

19 Q. Is this the Ross study that you were referring  
20 to?

21 A. Yes.

22 Q. Now, when you state that Ross reported  
23 decreased proliferation of cells in vitro, were you  
24 referring to a specific table of data when you reached  
25 that conclusion?

1           A. For example, some of his data is shown in  
2 Figure 1.

3           Q. If we could stay with Figure 1, isn't it also  
4 true that Ross reported increased proliferation of cells  
5 when the signal amplitude began to exceed 0.6 millitesla?

6           A. Yes.

7           Q. And I'm not sure if you could make this  
8 conversion. We have been discussing about milligauss and  
9 gauss in this proceeding. Can you make the conversion  
10 between a millitesla and a milligauss?

11          A. Yes. One millitesla is ten gauss. So you are  
12 talking about 10,000 milligauss. Or 0.6 millitesla would  
13 be 6,000 milligauss.

14          Q. And at page 20, line 12 through line 15 --

15          A. I'm sorry. We are back --

16          Q. I'm sorry. We are back on your testimony.

17          A. Okay.

18          Q. As it relates to the Ross study.

19          A. Page 20 -- could you say it again?

20          Q. Yes. Lines 12 through 15.

21          A. Yes.

22          Q. And you are stating that the Ross study  
23 consistently reports that no such effect occurs as a  
24 result of exposure to power frequency fields. First of  
25 all, could you define what you mean by the power

1 frequency fields?

2 A. Fields that come from 60 hertz transmission  
3 lines.

4 Q. You were not limiting that to a specific gauss  
5 level or milligauss level in this review, is that  
6 correct?

7 A. Correct.

8 Q. And after reviewing Figure 1 does that change  
9 your conclusion that the studies consistently report no  
10 such effect occurs as a result of exposure to power  
11 frequency fields?

12 A. No.

13 Q. And why not?

14 A. The proliferation effects as described in the  
15 Ross paper are nearly insignificant. In terms of  
16 amplitudes, magnetic field intensities, et cetera,  
17 different authors report effects that change at different  
18 levels. There has never been any consistent dose  
19 response with respect to the level of the fields. And  
20 therefore, putting this in context with the other  
21 research that I have reviewed, I think it has no bearing  
22 at all on the cancer question.

23 Q. Now, if we could go back to pages nine and ten  
24 of your testimony where you discuss the work of  
25 Dr. Liboff?

1 A. Yes.

2 Q. And just for summary purposes, would you agree  
3 that Dr. Liboff's work measured the uptake of tritiated  
4 thymidime?

5 A. The assay he used in the paper that I discuss  
6 in the testimony was tritiated thymidime uptake in cell,  
7 yes.

8 Q. I'm sorry. I was referring to the study you  
9 discussed.

10 A. Yes.

11 Q. And it was your conclusion in your testimony  
12 that this had no relationship to DNA synthesis, is that  
13 correct?

14 A. Yes.

15 Q. Is an increase in thymidime uptake by the cell  
16 consistent with an increase in DNA synthesis?

17 A. If there was an increase in DNA synthesis there  
18 would be an increase in tritiated thymidime labeling.  
19 But an increase in tritiated thymidime uptake does not  
20 mean there was an increase in DNA synthesis.

21 Q. You have used the terms labeling and uptake as  
22 obviously two different things. Could you explain the  
23 difference?

24 A. For the last answer it would be synonymous.

25 Q. If the word uptake and labelling is synonymous

1 in your last answer, would it be possible that Liboff's  
2 observations may be related to DNA synthesis?

3 A. Well, a lot of things are possible. But the  
4 experiments as performed and reported unfortunately don't  
5 tell us anything about DNA synthesis.

6 Q. And could this issue be resolved by repeating  
7 the experiment using a more direct assay for DNA  
8 synthesis than what was used in the Liboff study?

9 A. You are asking me could the experiment having  
10 done better or could a better designed experiment address  
11 the answer more specifically?

12 Q. Yes.

13 A. Yes.

14 Q. And do you think it is important to do so, to  
15 use a better designed experiment to address that question  
16 of DNA synthesis?

17 A. I think it has been done.

18 Q. And where has that been done?

19 A. I think other papers have looked at the effect  
20 of fields on cell growth and it has been included in  
21 other work and there have not been any clear  
22 demonstration that you can increase cell growth with  
23 power frequency fields.

24 MS. McCLOSKEY: I have no further questions, Your  
25 Honor.

1 JUDGE SMOLEN: Mr. Sugarman.

2 CROSS-EXAMINATION

3 BY MR. SUGARMAN:

4 Q. Do you agree with the U.S. Environmental  
5 Protection Agency, Air and Waste Management Division,  
6 report of August 9, 1991, that a wide variety of studies  
7 indicate an influence of E/MF on pathways that regulate  
8 cell growth and proliferation?

9 A. You are talking about the EPA draft report?

10 Q. No. I am talking about a technical paper  
11 entitled Status Report of the Research in Electromagnetic  
12 Fields prepared by the U.S. EPA, August 9, 1991.

13 MR. WATSON: Your Honor, I object to the question  
14 unless he shows him the document.

15 JUDGE SMOLEN: Why don't you show the witness the  
16 document.

17 MR. SUGARMAN: Sure.

18 BY MR. SUGARMAN:

19 Q. Directing your attention to the first full  
20 paragraph on page five --

21 MR. WATSON: Why don't you let me see it first.

22 (Document handed to Mr. Watson.)

23 MR. WATSON: This is a document prepared by a  
24 Michael Buccigrossi?

25 MR. SUGARMAN: Prepared by the agency. It is

1 issued by the agency. I object to your characterization.

2 MR. WATSON: I think it says at the bottom of page  
3 12 prepared by Michael Buccigrossi.

4 MR. SUGARMAN: It is issued by the agency.

5 MR. WATSON: I think it is actually issued by  
6 Region II.

7 MR. SUGARMAN: That is the agency.

8 MR. WATSON: That is one of the regions of the  
9 agency.

10 JUDGE SMOLEN: What is the question, now?

11 BY MR. SUGARMAN:

12 Q. Do you agree or disagree with the statement in  
13 this published report, "A wide variety of studies on the  
14 other hand, indicate an influence of E/MF on pathways  
15 that regulate cell growth and proliferation"?

16 A. I have never seen this document. I don't know  
17 specifically what studies it is referring to. I would be  
18 glad to take time to read it and then talk to you about  
19 it. But to take a sentence out of it and to ask me  
20 whether it is true or not is just something that I  
21 can't --

22 Q. You don't have an opinion about -- you don't  
23 have to read the report to have an opinion on the  
24 subject. The statement is, "A wide variety of studies on  
25 the other hand, indicate an influence of E/MF on pathways

1 that regulate cell growth and proliferation." Do you  
2 have any opinion on that subject?

3 MR. WATSON: Objection. Asked and answered.

4 MR. SUGARMAN: He didn't answer it.

5 JUDGE SMOLEN: Well, it's answered in his direct  
6 testimony but it is cross-examination. So I will let him  
7 answer the question.

8 A. What pathways are you discussing?

9 BY MR. SUGARMAN:

10 Q. Any pathways.

11 A. The statement that you are reading to me is  
12 hopelessly broad and I can't answer it in any effective  
13 way.

14 Q. Are you familiar with the work on protein  
15 kinase activity in response to E/MF by Byus, et al. in  
16 1984?

17 A. Yes.

18 Q. And the statement is made in here that "protein  
19 kinase activity has been altered in response to E/MF."  
20 Do you agree with that?

21 A. I am familiar with the paper by Byus. I don't  
22 necessarily agree with that conclusion about it but I am  
23 familiar with the paper.

24 Q. The statement is made, "magnetic fields have  
25 also been demonstrated to react a membrane bound

1 receptors in bone tissue of mice which stimulated  
2 collagen synthesis," citing Luben, 1982. Are you  
3 familiar with Luben's work?

4 A. I may have read it but I don't recall it right  
5 now.

6 Q. A statement is made, "another indication of  
7 cell proliferative activity, RNA transcription, was shown  
8 to increase with exposure to E/MF," citing Goodman and  
9 Henderson. Do you agree with that statement?

10 A. I am familiar with the work of Goodman and  
11 Henderson. Many of the papers that you're rapidly citing  
12 do not use just -- first of all, don't use just power  
13 frequency fields but some of them use pulse fields.

14 Secondly, the work of Goodman and Henderson about  
15 RNA transcription in some papers uses sinusoidal fields  
16 and in some cases uses pulse field.

17 So if we are going to talk about the data and what  
18 it means we will have to get more specific.

19 Q. The statement is made, "further supporting  
20 evidence shows human cancer sells were found to produce  
21 more colonies after exposure to magnetic fields or  
22 electric and magnetic fields combined but not electric  
23 fields alone," citing Philips and Winters, 1987, and  
24 stating, "this increased proliferative response was  
25 correlated with increased numbers of transferant

1 receptors, also indicative of growth potential." Are you  
2 familiar with Philips and Winters?

3 A. Yes.

4 Q. Do you agree with that characterization of  
5 their work?

6 A. No, for several reasons.

7 Q. The next statement is made, "E/MF is also  
8 suspected of having the potential to enhances the effect  
9 of other promoters. TPA, a known cancer promoter, was  
10 more effective in causing tumors when mice were also  
11 exposed to a magnetic field," citing Stuchly, et al., in  
12 1989. Are you familiar with Stuchly?

13 A. Yes.

14 Q. Do you agree with that characterization of his  
15 work?

16 A. It is Maria. It's her work, Stuchly. And, no,  
17 as a matter of fact that is an absolutely incorrect  
18 reading of the data. There were no statistical  
19 significant differences in that work.

20 Q. So you, then, disagree -- well, do you disagree  
21 with the conclusion on page six, "individually and  
22 collectively the results of these studies support the  
23 hypothesis that E/MF influences cell growth patterns and  
24 as such may have the potential to promotion cancer  
25 cells"?

1           A. I have not read that document. Using the  
2 statement that you just read and the studies that you  
3 have just reeled off, I completely disagree. It is an  
4 erroneous conclusion and it reflects a complete  
5 misunderstanding of the biology of cancer and of cell  
6 biology.

7           Q. Now, you were asked by Ms. McCloskey about  
8 whether you considered any of the epidemiological studies  
9 in formulating your opinion as expressed in your direct  
10 testimony. Is it true that you have not considered any  
11 of the epidemiological studies?

12          A. I don't know what you mean by considered. I  
13 have read them.

14          Q. Well, do you agree that the epidemiological  
15 studies by Wortheimer, Savitz, and -- well, I will  
16 withdraw the question and ask a couple of preparatory  
17 questions.

18           Are you familiar with Matanoski's report or study?

19          MR. WATSON: Your Honor, I am going to object to  
20 this right now so that there is no question of waiver  
21 beyond this. This witness has testified in a specific  
22 field. He has identified that field, he said he is not  
23 an epidemiologist. He didn't testify on it. And it is  
24 totally inappropriate cross-examination to begin to  
25 attempt to cross-examine him on the field that Dr. Cole

1 testified in just the other day. It is beyond the scope  
2 of his direct examination.

3 JUDGE SMOLEN: Mr. Sugarman.

4 MR. SUGARMAN: Well, he has testified that he read  
5 the epidemiological studies. I am asking if he  
6 considered them. By consider I mean did they have any  
7 bearing on your opinion one way or the other.

8 JUDGE SMOLEN: I will allow that question.

9 MR. SUGARMAN: I am withdrawing that question. I  
10 want to know, because it is obvious if he didn't consider  
11 them --

12 JUDGE SMOLEN: Just ask the question.

13 MR. SUGARMAN: Fine.

14 BY MR. SUGARMAN:

15 Q. Did they have any bearing on your opinion one  
16 way or the other?

17 A. In terms of what I have written in my report?

18 Q. And what you have answered Ms. McCloskey's  
19 questions.

20 A. No.

21 Q. Do you consider it scientifically appropriate  
22 to exclude epidemiology?

23 MR. WATSON: Objection, Your Honor. Argumentative.

24 JUDGE SMOLEN: I am going to sustain that. He is  
25 offered as an expert in a particular field and that is

1 what he has testified to.

2 BY MR. SUGARMAN:

3 Q. Do you consider it be accepted procedure in  
4 your field, oncology, to exclude consideration of  
5 epidemiology in forming conclusions about your work?

6 MR. WATSON: Objection, Your Honor. The same  
7 objection. It's argumentative. He has testified to a  
8 particular field.

9 MR. SUGARMAN: It's not argumentative.

10 JUDGE SMOLEN: Sustained. Next question.

11 MR. SUGARMAN: It's not argumentative.

12 JUDGE SMOLEN: He has testified -- the objection  
13 was sustained.

14 MR. SUGARMAN: But I have a right to find out  
15 whether he considers it appropriate to -- whether it is  
16 his understanding of his field that epidemiology should  
17 or should not be considered.

18 JUDGE SMOLEN: That's a different question.

19 MR. SUGARMAN: Okay.

20 BY MR. SUGARMAN:

21 Q. Do you consider it appropriate -- do you have  
22 an opinion about whether your field considers it  
23 appropriate to exclude epidemiology in forming opinions  
24 in your field?

25 A. Yes, I have an opinion.

1 Q. What is your opinion?

2 A. My opinion is that in scientific knowledge one  
3 must take all information and assess it for its worth.

4 Q. Well, did you assess the epidemiology for its  
5 worth?

6 A. I was not asked to assess the epidemiology, and  
7 it's beyond the scope of my expertise.

8 Q. So, then, if in all scientific fields you must  
9 assess the related information, as you have just said,  
10 and yet you can't do it, how could you fulfill your  
11 scientific or how did you fulfill your scientific  
12 obligation?

13 MR. WATSON: Objection, Your Honor. Argumentative.

14 JUDGE SMOLEN: Sustained.

15 BY MR. SUGARMAN:

16 Q. You said it has to be done. You said you  
17 couldn't do it. How do you square those two?

18 MR. WATSON: Objection, Your Honor. Argumentative.  
19 It's the same question.

20 MR. SUGARMAN: This is cross-examination.

21 BY MR. SUGARMAN:

22 Q. How could you reach a conclusion -- I am trying  
23 to be non-argumentative. This is not a rhetorical  
24 question. How do you go about reaching a conclusion if  
25 you can't do something that is necessary to do?

1 MR. WATSON: Objection, Your Honor. Argumentative.  
2 Counsel knows well that this expert has testified in a  
3 particular field. If you want an opinion in the field of  
4 epidemiology you go to an expert in epidemiology and get  
5 it.

6 MR. SUGARMAN: I don't want an opinion in the field  
7 of epidemiology. I don't want that. What I want is I  
8 want a response --

9 JUDGE SMOLEN: Temperance, Mr. Sugarman.  
10 Temperance, please. Go ahead.

11 MR. SUGARMAN: I'm sorry.

12 JUDGE SMOLEN: What was your response to the  
13 objection?

14 MR. SUGARMAN: My response to the objection is that  
15 the witness has said that it is necessary to consider all  
16 relevant information. I asked him if he assessed the  
17 relevancy of the epidemiology. He said he couldn't do  
18 it. So my question is how can he form an opinion without  
19 doing it. If it's necessary to do it -- it seems to me  
20 it's a syllogism. So I am asking him to explain.

21 JUDGE SMOLEN: You can respond.

22 MR. WATSON: Your Honor, I think this started with  
23 the same series of questions. It's an argumentative  
24 question. It is based upon a fundamental  
25 misunderstanding or position that Mr. Sugarman has that

1 somehow this expert must go into the field of  
2 epidemiology necessarily to reach a conclusion, even if  
3 he has enough in the field of molecular genetics alone to  
4 reach an answer.

5 MR. SUGARMAN: Well, if that is his answer let him  
6 give that answer. But that is Mr. Watson's testimony.

7 JUDGE SMOLEN: One voice at a time.

8 MR. WATSON: And this expert has said that as to  
9 offering an expert opinion in the field of epidemiology,  
10 he has left that to somebody with expertise in that  
11 field.

12 MR. SUGARMAN: That is not what he said.

13 MR. WATSON: He has made that, I think, clear, and  
14 all of this is basically argument.

15 MR. SUGARMAN: That's not what he said --

16 JUDGE SMOLEN: Let's ask the question again.

17 Please, let's keep composure here. Ask the question  
18 again.

19 MR. SUGARMAN: I mean, I would just like to  
20 response -- I will ask the question, Your Honor, but I  
21 would like to respond to what I would consider the  
22 epithet that something is argumentative whenever  
23 Mr. Watson doesn't like it. I don't understand the term  
24 if it means everything that is cross-examination. I  
25 mean, I don't believe Mr. Watson either believes what he

1 says or knows what he is saying when he characterizes  
2 questions as argumentative simply because they take  
3 previous testimony and ask the witness if he can  
4 reconcile it.

5 JUDGE SMOLEN: Ask your question.

6 BY MR. SUGARMAN:

7 Q. You testified that it is necessary to consider  
8 all relevant data.

9 JUDGE SMOLEN: Wait. Was that your testimony?  
10 Finish that piece of the question.

11 MR. SUGARMAN: Can we read it back?

12 JUDGE SMOLEN: Yes.

13 MR. SUGARMAN: Well, it was before the last  
14 question. First I asked is it necessary to consider  
15 epidemiology and he said yes, it's necessary --

16 JUDGE SMOLEN: To do what? Necessary to consider  
17 epidemiology in what.

18 MR. SUGARMAN: In forming a scientific opinion.

19 JUDGE SMOLEN: About what?

20 MR. SUGARMAN: In his field.

21 JUDGE SMOLEN: And what was your answer? Put that  
22 on the record again. What was your answer to that?

23 THE WITNESS: Can I get the question one more time?  
24 I'm sorry.

25 JUDGE SMOLEN: Just repeat that question.

1 BY MR. SUGARMAN:

2 Q. I asked in your field of oncology is it  
3 necessary to consider epidemiological evidence in forming  
4 opinions --

5 A. Yes.

6 Q. -- as a matter of scientific practice.

7 JUDGE SMOLEN: He answered yes. Go ahead.

8 BY MR. SUGARMAN:

9 Q. And I asked you, then, did you consider the  
10 epidemiology, and you said no, it was outside the scope  
11 of your assignment and outside the scope of your  
12 expertise, isn't that right?

13 MR. WATSON: Your Honor, I object to that. That is  
14 a misstatement of the record. He did not say -- he said  
15 he did not consider it with respect to the issues  
16 presented in --

17 JUDGE SMOLEN: Well, let's ask the question and get  
18 the witness to answer. If it's repetitive, it's  
19 repetitive. Ask the second half of your question again.

20 MR. SUGARMAN: The only trouble is if I get a  
21 different answer I want to go back to the earlier one  
22 because Counsel is trying to coach the witness through  
23 his arguments.

24 BY MR. SUGARMAN:

25 Q. I will ask it fresh as if I didn't ask it

1 before. Did you consider the epidemiology and did it  
2 affect your conclusion one way or the other? I think you  
3 answered no, right?

4 A. I read the epidemiology; it did not affect my  
5 conclusion.

6 Q. Right. Then I asked you did you consider it.  
7 I will ask you that again. Did you consider it?

8 A. I considered it as I would any epidemiologic  
9 study that I would read as a medical oncologist.

10 Q. Do you remember answering me five minutes ago  
11 that it was outside the scope of your assignment and you  
12 don't have the expertise? Do you remember giving me that  
13 answer five minutes ago?

14 A. Yes.

15 Q. Okay. Now, if it was outside of your  
16 assignment and outside of your expertise, how did you  
17 consider it?

18 A. It is not abnormal or unreasonable to read  
19 something and be informed about it. For the purposes of  
20 the area that I was asked to review and comment on based  
21 on my expertise I focused on molecular and cellular  
22 biology and the tumor cell genetics of the question.  
23 There is nothing about the epidemiology that is  
24 considered in the report. I am aware of the  
25 epidemiology.

1 For example, if I were reading a paper about the  
2 epidemiology of some other cancer, I would read it and  
3 weigh it as I could. If I were very concerned about it  
4 and if I wanted to know more I would go to an  
5 epidemiologist and say what do you think about the data  
6 in this study and I would utilize that expertise.

7 Q. Did you ask Counsel whether you could consult  
8 with an epidemiologist as part of your scope of work?

9 A. The epidemiology was outside the scope of the  
10 work. I wasn't asked to consider it as part of the  
11 studies that I have cited. I am focusing here on the  
12 cell biology and the genetics of the question.

13 Q. Didn't you just say a little while ago that it  
14 is necessary to consider the science -- I'm sorry -- that  
15 it is necessary to consider the epidemiology in doing the  
16 science?

17 MR. WATSON: Objection, Your Honor.

18 JUDGE SMOLEN: I am going to sustain it. It has  
19 been asked and answered. We are going over the same  
20 thing. Go to the next question.

21 BY MR. SUGARMAN:

22 Q. About the epidemiology, if it were true that  
23 the epidemiology showed a trend of association between  
24 childhood leukemia and exposure to E/MF, would that  
25 affect your conclusions?

1 MR. WATSON: Objection, Your Honor. It assumes  
2 facts not in evidence. There is no evidence to support a  
3 proposition that the epidemiology would at all support  
4 that view. In fact, the only evidence by a qualified  
5 epidemiologist is directly to the contrary.

6 MR. SUGARMAN: Rosenbaum testified on the basis of  
7 literature review. Liboff testified on the basis of his  
8 participation that the epidemiology shows that  
9 association. The witness Cole conceded there is an  
10 association. He challenged the epidemiology, but he  
11 agreed there are epidemiological studies that purport to  
12 show an association between E/MF and childhood leukemia.  
13 So there is plenty of evidence in the record.

14 MR. WATSON: Your Honor, just to respond to that,  
15 neither Rosenbaum nor Liboff have any qualifications in  
16 the field of epidemiology whatsoever, and Dr. Cole did  
17 not testify as Mr. Sugarman described it.

18 MR. SUGARMAN: He certainly did.

19 JUDGE SMOLEN: One says yea and the other says nay.  
20 Let's get the question again.

21 BY MR. SUGARMAN:

22 Q. If there were studies that showed --

23 JUDGE SMOLEN: Is this a hypothetical question?

24 MR. SUGARMAN: Yes. But it is based on evidence in  
25 the record. I don't expect the witness have read the

1 record --

2 JUDGE SMOLEN: Let's hear it.

3 MR. SUGARMAN: -- since part of it was made  
4 yesterday.

5 BY MR. SUGARMAN:

6 Q. If there were studies that showed an  
7 association or purported to show an association between  
8 exposure to E/MF and childhood leukemia would that change  
9 your opinion?

10 JUDGE SMOLEN: About what?

11 BY MR. SUGARMAN:

12 Q. About your conclusion that exposure to E/MF has  
13 no effect on cancer.

14 JUDGE SMOLEN: You may answer.

15 A. It would not change my opinion based on the  
16 studies I reviewed and my scientific expertise.

17 BY MR. SUGARMAN:

18 Q. Now, if there were epidemiological studies  
19 which purported to show an effect of exposure to E/MF on  
20 cancer incidence in occupational fields of adults, would  
21 that change your testimony?

22 A. No.

23 Q. Now, did you study potential effects of E/MF on  
24 the pineal gland?

25 A. Can you be more specific?

1 Q. Did you study whether E/MF affects the  
2 production of the pineal gland?

3 A. I have read some papers in that regard, but  
4 E/MF effects on pineal gland secretion is outside of my  
5 expertise.

6 Q. So when you gave your opinion in your direct  
7 testimony at page 22, lines 24 to 27, that, quote, the  
8 extensive research on molecular and cellular biology and  
9 cancer provides no scientific basis to conclude that  
10 power frequency electric and/or magnetic fields cause,  
11 promote or contribute to the development of cancer or  
12 other adverse health effects, end of quote, you did not  
13 include in that possible cancer impacts of effects on the  
14 pineal gland, is that right?

15 A. Yes, I did.

16 Q. Well, how could you do that if you didn't study  
17 -- please tell us, and this is a non-argumentative -- can  
18 you please tell you how you could consider them in  
19 forming that conclusion if you didn't have the expertise  
20 to consider whether E/MFs affect pineal gland production?

21 MR. WATSON: Objection, Your Honor. It is  
22 argumentative. He asserts in there that the witness  
23 didn't have the expertise.

24 MR. SUGARMAN: All I did was quote his own  
25 testimony.

1 JUDGE SMOLEN: I am going to overrule the  
2 objection. You can answer that question.

3 A. You are talking about two different things.  
4 The effects on the pineal gland in animals is an  
5 endocrinologic question. The issue of whether the  
6 primary pineal hormone, which is melatonin, which has  
7 something to do with cancer in humans is an oncologic  
8 question, and the answer is that it has nothing to do  
9 with cancer in humans.

10 BY MR. SUGARMAN:

11 Q. So you are not testifying as to whether the  
12 effect on the pineal gland could have some other health  
13 effects other than cancer?

14 A. I didn't say that.

15 Q. So, well, if you don't have the --

16 JUDGE SMOLEN: I'm not sure the witness finished  
17 his answer.

18 THE WITNESS: I'm done.

19 JUDGE SMOLEN: Okay. Sorry.

20 BY MR. SUGARMAN:

21 Q. What is your basis or did you -- let me ask the  
22 question a different way. Do you wish to express an  
23 opinion about whether effects on melatonin production by  
24 the pineal gland have any health effects or may have any  
25 health effects?

1 A. No.

2 Q. Do you wish to express an opinion on whether  
3 they have any adverse health effects? That is, on  
4 whether reduction of production in the pineal gland has  
5 any adverse health effects. Do you wish to express an  
6 opinion on that?

7 A. Not outside my area of expertise or what I have  
8 written in my report.

9 Q. Well, you have written in your report, quote,  
10 other adverse health effects, which could cover  
11 everything or nothing. And it does not say any other.  
12 So it may say some other or whatever. So I am asking you  
13 specifically with respect to the potential adverse health  
14 effects of decreases in melatonin production by the  
15 pineal gland, are you wishing to express an opinion as to  
16 whether that might have any adverse health effects?

17 MR. WATSON: Objection, Your Honor. Asked and  
18 answered.

19 MR. SUGARMAN: Oh, no.

20 JUDGE SMOLEN: Wait. Give him a chance.

21 MR. WATSON: I would like to finish my objection  
22 before Mr. Sugarman reacts.

23 I think the witness clarified when Ms. McCloskey  
24 asked him the specific phrase "other adverse health  
25 effects" that he was referring to adverse health effects

1 within the field of molecular and cellular biology and  
2 cancer only. He made that clear. He has already  
3 answered this.

4 JUDGE SMOLEN: What I am going to do is I am going  
5 to overrule the objection. If the witnesses wants to  
6 answer it that way then he will answer. He is an  
7 sophisticated witness. He is an expert. He can answer  
8 that question, what he meant, really, by adverse health  
9 effects. If it's surplusage, it's surplusage.

10 THE WITNESS: Can I get the question read back,  
11 please?

12 (Whereupon, the reporter read from the record as  
13 requested.)

14 A. No.

15 BY MR. SUGARMAN:

16 Q. Thank you,

17 Now, are you familiar with the work of Grahm, et  
18 al., dealing with a slowing of human heart rate during  
19 exposure to magnetic fields?

20 MR. WATSON: Objection, Your Honor. Beyond the  
21 scope of his testimony. He didn't testify as a  
22 cardiologist.

23 BY MR. SUGARMAN:

24 Q. It says here the extensive research on  
25 molecular and cellular biology and cancer provides no

1 scientific basis to conclude that power frequency  
2 electric and/or magnetic fields cause, promote or  
3 contribute to the development of cancer or other adverse  
4 health effects. What does that phrase other adverse  
5 health effects mean? You talk cellular and molecular  
6 biology as if that limits the phrase other adverse health  
7 effects. Fine. Now I need to know is the slowing of the  
8 human heart a molecular or cellular biology question.

9 JUDGE SMOLEN: Well, that is okay.

10 MR. WATSON: If he asks that question that would be  
11 different.

12 BY MR. SUGARMAN:

13 Q. You indicated, and Mr. Watson --

14 JUDGE SMOLEN: Well, wait a minute. Let's get an  
15 answer to that.

16 BY MR. SUGARMAN:

17 Q. Is the slowing of the human heart a molecular  
18 or cellular biology question?

19 A. It may be and it may not be.

20 Q. And as you gave your testimony, other adverse  
21 health effects, were you intending to give an opinion and  
22 are you now intending to give an opinion as to whether  
23 E/MF slows human heart rates?

24 A. You are asking me if E/MFs slow human heart  
25 rates?

1 Q. I am asking you if you are intending to give an  
2 opinion on that subject.

3 A. No.

4 Q. Are you familiar with the work of Creim, et  
5 al., 1989, concluding that behavioral effects have also  
6 been investigated and that rats made more errors in  
7 performance in a radial arm maze when exposed to a DC  
8 magnetic field combined with a 60 hertz magnetic field?

9 A. No.

10 MR. WATSON: Your Honor --

11 JUDGE SMOLEN: Well, he answered. He's not  
12 familiar with the study.

13 BY MR. SUGARMAN:

14 Q. Now, did you include in your opinion anything  
15 having to do with effects of E/MF on reproduction?

16 MR. WATSON: Your Honor, objection. This is --

17 BY MR. SUGARMAN:

18 Q. What does the phrase other adverse health  
19 effects mean?

20 MR. WATSON: He has testified several times that by  
21 other adverse health effects, and he started with the  
22 cross-examination by the OCA, that it refers to, and he  
23 said it at least twice, molecular and cellular biology in  
24 cancer.

25 MR. SUGARMAN: Okay. Let me ask the other

1 question.

2 JUDGE SMOLEN: Sustained.

3 BY MR. SUGARMAN:

4 Q. Are reproduction effects within the term  
5 molecular and cellular biology?

6 MR. WATSON: Your Honor, that is not -- I object to  
7 that. That is not the same question.

8 MR. SUGARMAN: It's a different question.

9 JUDGE SMOLEN: I sustained your objection before.  
10 Now it's a new question.

11 MR. SUGARMAN: I am asking the question.

12 A. They could be.

13 BY MR. SUGARMAN:

14 Q. Were you intending and are you now intending to  
15 express an opinion as to the effect of E/MF on  
16 reproduction?

17 A. I have not done an exhaustive literature search  
18 on the issue of reproduction. Some of the papers that I  
19 have read included information about reproduction.

20 Q. My question is were you and are you now  
21 intending to express an opinion about reproduction as  
22 affected by E/MF?

23 A. You want to know if I am saying that E/MF has  
24 an effect on human reproduction?

25 JUDGE SMOLEN: No. That wasn't the question. Do

1 you intend to express an opinion on that?

2 THE WITNESS: No.

3 BY MR. SUGARMAN:

4 Q. Thank you.

5 Now, on the question of cyclotron resonance you  
6 expressed an opinion with respect to some of Dr. Liboff's  
7 work, specifically at pages nine and ten of your  
8 testimony and 19 and 20 of your testimony. My question  
9 is this: if I understand -- and this is a very  
10 complicated area for me at least so this question may  
11 seem too simplistic to you -- if I understand your  
12 testimony you are not challenging that Dr. Liboff might  
13 have demonstrated that the ion cyclotron resonance effect  
14 occurs. Your question is does it have any effect on  
15 human health. Is that what I -- and you believe that he  
16 has not demonstrated that it has any effect on human  
17 health. Am I correctly interpreting your testimony?

18 A. You lost me. I don't think I said anything  
19 about ion cyclotron resonance. I'm not sure I used the  
20 term in the report. Where are you referring to?

21 Q. I am referring to Liboff's work on DNA  
22 synthesis and on cell proliferation.

23 A. That is the paper referred to on page nine of  
24 my report, Liboff et al., in Science, 1984?

25 Q. Right. And Liboff's paper, 1991, cited at

1 pages 19 and 20 of your report. Liboff is referred to in  
2 lines 34 to 35 as Liboff's recent abstract using lymphoma  
3 cells.

4 JUDGE SMOLEN: What is the question again so that  
5 the witness knows what he is being asked?

6 BY MR. SUGARMAN:

7 Q. Is it your intention in either section of your  
8 testimony that I am referring to to challenge the  
9 statement or conclusion or to question the work as to  
10 whether electromagnetic fields cause ion movement across  
11 cell membranes?

12 MR. WATSON: Objection, Your Honor. That is a  
13 compound question, I believe.

14 JUDGE SMOLEN: Is a compound question. Break it  
15 up.

16 MR. SUGARMAN: All right. I'll break it down.

17 JUDGE SMOLEN: Sustained, then.

18 BY MR. SUGARMAN:

19 Q. Dr. Liboff -- you are familiar with  
20 Dr. Liboff's work as reflected in the references on pages  
21 nine and 19. Are you questioning whether there is a  
22 mechanism of action on E/MF by way of cyclotron  
23 resonance?

24 MR. WATSON: Objection, Your Honor. That is the  
25 same question, I believe. He has referred to page nine

1 and then page 19. They are two different sets of  
2 studies. On page nine it is an '84 study --

3 JUDGE SMOLEN: Sustained.

4 MR. SUGARMAN: Let's try it separately.

5 BY MR. SUGARMAN:

6 Q. In your critique or discussion of Liboff at  
7 pages 19 and 20, do you challenge his conclusions with  
8 respect to the mechanism of cyclotron resonance in  
9 causing movement of ions when exposed to E/MF?

10 MR. WATSON: Your Honor, I don't see a challenge to  
11 conclusions here or any reference to what he is talking  
12 about. I think the question is vague.

13 JUDGE SMOLEN: It's a good question. It may not be  
14 -- it's not objectionable. So I will overrule the  
15 objection.

16 You may answer if you understand the question?

17 A. Are you asking my conclusions about cyclotron  
18 resonance?

19 BY MR. SUGARMAN:

20 Q. Right.

21 A. I think you are better off to ask a physicist.

22 Q. You testified that the melatonin research does  
23 not provide a pathway by which exposure to E/MF could  
24 cause cancer in humans. That's page 19, lines 23 and 24.  
25 You then testify there are no data connecting melatonin

1 levels to breast cancer causation in humans. Are you  
2 including in the term no data epidemiological data? Or  
3 do you mean experimental data?

4 A. To my knowledge as a medical oncologist there  
5 is no data connecting melatonin levels to breast cancer  
6 causation in humans.

7 Q. Have you made any inquiry as to whether there  
8 is any epidemiology on that?

9 A. I have read several papers in the field. I  
10 couldn't quote them to you at this time.

11 Q. Now, the question that you were answering asks  
12 whether there was a pathway, whether they provide a  
13 pathway. And your answer is no to that question. Have  
14 you found any pathway by which any agent causes cancer in  
15 humans?

16 A. You want to know if I know of some agent that  
17 causes cancer in humans and how?

18 Q. I am using the term pathway. Have you found  
19 any pathway by which any agent causes cancer?

20 A. We know of a number of agents that cause  
21 cancer.

22 Q. That is not the question, sir. The question  
23 relates to the term pathway. Have you found any pathway  
24 by which any agent causes cancer?

25 A. I personally know of several agents that

1 through biochemical pathways cause cancer.

2 Q. Through biochemical, but can you tell us what  
3 the biochemical pathway is and how it works?

4 A. Yes.

5 Q. Okay. Tell us one, please.

6 A. Breast cancer causation.

7 Q. Go ahead. Tell us the pathway and how it  
8 works.

9 A. Estrogen is required for breast cancer  
10 causation.

11 Q. Right.

12 A. Estrogen stimulates breast cell proliferation  
13 and is a tumor promoter in humans as well as in animal  
14 models.

15 Q. How does it work to do that? What is the  
16 pathway? How does estrogen do that?

17 A. Perhaps your definition of pathway is different  
18 from mine. What specifically do you mean by pathway?

19 Q. How does it work?

20 A. It works by being a tumor promoter.

21 Q. So when you say does the research provide a  
22 pathway by which exposure to E/MF, referring to Wilson's  
23 research and the others on melatonin, what is the  
24 difference between the answer you just gave about  
25 estrogen and the answer that they give about melatonin

1 levels in terms of being able to describe exactly how it  
2 happens? What is the difference?

3 A. Well, would you please tell me what you are  
4 talking about when you talk about Wilson and what is it  
5 that he said that you want me to comment on?

6 Q. They show that reduced levels of -- they show  
7 that E/MF reduces melatonin levels. Melatonin levels  
8 have been associated with cancer. What do you mean by  
9 saying that there is no pathway?

10 MR. WATSON: Objection, Your Honor. It assumes  
11 facts not in evidence. In fact, they haven't shown that.  
12 That is just Counsel's assertion.

13 BY MR. SUGARMAN:

14 Q. Do you agree that they purport to show that?

15 A. Do I agree that they purport to show what?

16 Q. That melatonin levels are lowered.

17 A. In a study done by Wilson? I would have to  
18 look at the study to tell you what they show.

19 Q. It says here, "Have you reviewed the research  
20 by Wilson and others on melatonin?"

21 "Yes, I have reviewed that research. For example,  
22 Lerchl, '91; Reiter, '90; Wilson, '81, '86 '88 and 99."

23 A. Yes.

24 Q. Do you agree that that shows that E/MF exposure  
25 lowers melatonin production?

1 A. No.

2 Q. Do you agree that it purports to show that?

3 A. Yes.

4 Q. Do you agree that melatonin reduction is a  
5 cancer promoter?

6 A. No.

7 Q. I understand now.

8 (Pause.)

9 Q. Are you familiar with the use of  
10 electromagnetic fields to promote bone healing?

11 MR. WATSON: Objection, Your Honor. Beyond the  
12 scope. He didn't testify in the area of --

13 JUDGE SMOLEN: Sustained. Go ahead.

14 BY MR. SUGARMAN:

15 Q. Are you testifying that E/MF does not affect --  
16 and this is not a rhetorical question or an argumentative  
17 question -- are you intending to testify that E/MF does  
18 not affect cell formation and proliferation?

19 A. My testimony was that power frequency fields do  
20 not affect cell proliferation.

21 Q. Now, do you know whether they affect cell  
22 proliferation in connection with healing of bones?

23 A. No, I don't.

24 Q. Are you familiar with the use of E/MF in the  
25 medical community to affect bone healing?

1           A. I am vaguely aware that there are devices which  
2 are used to try to affect bone healing.

3           Q. And have you made any inquiry to determine  
4 whether they have any success?

5           A. Yes. Actually it is a very interesting area.

6           Q. And what is the nature of the inquiry that you  
7 made?

8           MR. WATSON: Your Honor, I assume that these  
9 questions are related to his testimony about the impact  
10 on cells as distinguished from the separate field of bone  
11 healing.

12           JUDGE SMOLEN: Cell proliferation, I think.

13           MR. WATSON: If that is the case I have no  
14 objection as long as they are within that arena.

15           MR. SUGARMAN: Cell proliferation, yes.

16           BY MR. SUGARMAN:

17           Q. Go ahead.

18           A. The question had to with what my inquiry was?

19           Q. Yes.

20           A. I have read several papers in the area.

21           Q. Do you have any reason to doubt that  
22 electromagnetic fields are successful in promoting cell  
23 proliferation as a means of healing bones?

24           A. I do.

25           Q. What is your reason for doubting it?

1           A. Well, the rigorous test of a medical or  
2 therapeutic modality often is a randomized trial. And I  
3 looked hard for randomized trials in this area and only  
4 one was started and it got to the point where 14 patients  
5 were randomized, 14 patients with fractures had not  
6 healed for more than a year, which is a long time, were  
7 randomized, 14 to receive the application of a device  
8 which was not turned on. So these patients were put to  
9 bed with this device strapped on but it was not working.  
10 And half of the patients were given the same treatment,  
11 only the device was turned on. And an equal number of  
12 fractures healed in both of those groups. So in the one  
13 study that was initiated and the one preliminary report  
14 that I read there was in fact no difference between those  
15 two groups.

16           And because I was particularly curious about it,  
17 because it is sort of intriguing, I couldn't find any  
18 evidence that when this was tested rigorously that at  
19 least one device did any good whatsoever.

20           Q. Is the device approved for use by the FDA?

21           A. I don't know. I don't use them. I am not an  
22 orthopedist.

23           Q. Have you talked to any clinicians who are using  
24 it?

25           A. No, I have not.

1 Q. Do you consider clinical evidence to be a valid  
2 basis for forming conclusions about the scientific  
3 processes?

4 A. Clinical experiments have to be evaluated for  
5 their rigor just as any other scientific study does.

6 Q. I didn't mean clinical studies, I meant  
7 clinical results, clinical effects?

8 A. Clinical anecdotes are virtually worthless.

9 Q. What was the study that you referred to that  
10 indicated equivalent or comparable healing for control  
11 and treated patients or individuals?

12 A. I believe Bassett was one of the authors but I  
13 would have to get it out of my files for you. I don't  
14 recall it.

15 Q. Do you agree that the Adey laboratory has  
16 interpreted their data as having a relevance between E/MF  
17 and cancer?

18 A. Which data are you --

19 MR. WATSON: Objection, Your Honor. I don't see  
20 how this witness could possibly be an expert on how  
21 someone who is not here in the courtroom today interprets  
22 his own data. There is no way he could know that.

23 MR. SUGARMAN: I don't mean in their heads. I mean  
24 in the reports.

25 MR. WATSON: If he shows him a report and asks him,

1 that would be a different question.

2 JUDGE SMOLEN: I sustain the objection.

3 BY MR. SUGARMAN:

4 Q. Have you previously testified, specifically in  
5 a New Jersey Public Utility Commission hearing, that in  
6 response to a question, "In reviewing the published work  
7 by the Adey laboratory isn't it true that in  
8 interpreting their data they suggest it has relevance to  
9 cancer?"

10 "Answer. In the discussion sections of the papers  
11 they do make those speculations."

12 Do you recall that question and answer?

13 A. Yes.

14 MR. WATSON: Could we have the page and line on  
15 that?

16 JUDGE SMOLEN: He has answered the question but,  
17 please, supply the page and line number of that.

18 MR. SUGARMAN: The page?

19 JUDGE SMOLEN: Yes.

20 MR. SUGARMAN: Here it is, page 52 and 53 of his  
21 cross-examination in New Jersey.

22 BY MR. SUGARMAN:

23 Q. And the next question was, "I take it from your  
24 direct testimony it is fair to say that to the extent  
25 that the OTA, Office of Technology Assessment, report

1 agrees that their research may have a relationship with  
2 cancer, you disagree with the OTA report, right?"

3 "Answer. With some of the assessments of the  
4 biological significance of the results from Adey's lab,  
5 yes. I think it is not critically assessed."

6 Do you still hold that opinion?

7 A. Yes.

8 Q. Now, the Adey studies were related to ODC  
9 levels, were they not?

10 A. There is one paper that I am familiar with on  
11 ornithine decarboxylase level. Let's just make sure we  
12 are talking about the same one. Which one are we talking  
13 about?

14 Q. That one. I am not going to try to pronounce  
15 it. Are you familiar with that, right?

16 A. I am familiar with the paper in Cancer Research  
17 which addresses ornithine decarboxylase levels, yes.

18 Q. And that is the research in which Adey and his  
19 group purported to associate the impacts of E/MF with  
20 cancer, is that not correct?

21 A. I'm not sure about that.

22 Q. I thought you said that -- "Isn't it true that  
23 in interpreting their data they suggest that it has a  
24 relevance to cancer?"

25 "Answer. In the discussion sections of the paper

1 they do make those speculations."

2 A. That particular paper in their discussion  
3 section says that the data presented in the paper -- that  
4 any connection between cancer causation and that data is  
5 entirely speculative. I believe it is phrased something  
6 like that.

7 Q. Now, is it true that in the New Jersey  
8 proceeding you were not asked to get into the initiation  
9 and promotion process?

10 A. I don't remember.

11 Q. I show you Mr. Watson's statement at pages 33  
12 and 34 of the record, where it says, "Here we have a  
13 medical oncologist testifying on the medical oncology and  
14 the cellular biology," referring to you. "Dr. Boutwell  
15 was testifying solely limited to the question of cancer  
16 initiation and promotion processes, which is a separate  
17 area. And he was not a medical oncologist. He was a  
18 regular oncologist. So while they may use the same  
19 study, we have the same situation here."

20 And he goes on to say, "We are not asking him" --  
21 he says, "You've have heard the same study mentioned by  
22 several different people and you will probably hear a  
23 number of studies mentioned by several different people,  
24 but we are confining them to the area of expertise that  
25 they are in and what that study tells them only within

1 that area. We are not asking him," that is referring to  
2 you, "to get into the initiation and promotion process. I  
3 think you can tell already we have not asked him that."

4 MR. WATSON: Your Honor, I object to this. He is  
5 entitled to cross-examination the witness about a prior  
6 inconsistent statement, but nothing more. There is no  
7 statement by the witness here.

8 JUDGE SMOLEN: I don't have a question yet.

9 BY MR. SUGARMAN:

10 Q. Do you remember in the New Jersey proceeding  
11 that you did not get into initiation and promotion?

12 MR. WATSON: Objection, Your Honor. Irrelevant.  
13 He is only entitled to ask him about a prior inconsistent  
14 statement from an another case.

15 JUDGE SMOLEN: I'm going to sustain that.

16 BY MR. SUGARMAN:

17 Q. Is it true that you are not an expert in  
18 initiation and promotion?

19 A. No.

20 Q. Are you an expert in initiation and promotion?  
21 Or should I break it down separately?

22 A. I have an expert understanding of the molecular  
23 events of the initiation and promotion of cancer.

24 Q. Did you indicate an unwillingness to testify in  
25 that area previously?

1 A. No.

2 MR. WATSON: Objection, Your Honor. Irrelevant.

3 JUDGE SMOLEN: He answered.

4 You mean in this proceeding?

5 MR. SUGARMAN: No. Previously in response to  
6 Mr. Watson's request.

7 JUDGE SMOLEN: In the other case?

8 MR. SUGARMAN: Yes.

9 JUDGE SMOLEN: Sustained.

10 BY MR. SUGARMAN:

11 Q. You answered Ms. McCloskey that you didn't see  
12 any indication of any need for more research in the  
13 field, is that correct? Of E/MFs and cancer.

14 A. My answer sounded something like that but it  
15 wasn't exactly that.

16 Q. Could you tell us how I misstated it and  
17 correct me?

18 A. I would have to rehear her questions.

19 MR. WATSON: Objection, Your Honor. This is  
20 repetitious.

21 JUDGE SMOLEN: It may be repetitious of what  
22 Ms. McCloskey asked, but we have a different attorney  
23 representing a different client and I think he is allowed  
24 some latitude. So I will overrule.

25 BY MR. SUGARMAN:

1 Q. Is it your opinion that there is no need for  
2 additional research relating to E/MF exposure in cancer  
3 within your field?

4 A. Yes.

5 Q. That's what I thought. Okay.

6 And do you recognize Reiss, R-e-i-s-s? I think you  
7 cite him in your direct.

8 A. Isn't it R-e-e-s-e?

9 Q. Okay.

10 A. Are we talking about the same Reese?

11 Q. Yes, possibly so. His work of 1988. Yes,  
12 Reese, Jostes and Frazier, Exposure of Mammalian Cells to  
13 60 hertz Magnetic or Electric Fields, cited at page 32 of  
14 your testimony.

15 A. Yes.

16 Q. Do you agree that he recommends additional  
17 research be done?

18 A. I don't recall what was said in that regard in  
19 that paper.

20 Q. In the New Jersey proceeding you were quoted a  
21 page of that report. Page 12 of your testimony, "The  
22 research represents part of a systematic study by his  
23 laboratory to determine mechanisms by which electrical or  
24 magnetic fields interacted with cells and cellular  
25 constituents." You agreed that that was a correct

1 statement.

2 MR. WATSON: Your Honor, I think he ought to show  
3 him the statement if he is going to read from another  
4 transcript. And there does not seem to be any prior  
5 inconsistent statement anyway.

6 JUDGE SMOLEN: It may be the groundwork for a prior  
7 inconsistent statement.

8 BY MR. SUGARMAN:

9 Q. Did you agree in New Jersey that Reese  
10 recommended additional research on the possibility that a  
11 low rate of DNA damage if linked to cell cycle would not  
12 have been detected after a one hour exposure?

13 MR. WATSON: Objection, Your Honor. There is prior  
14 inconsistent statement.

15 JUDGE SMOLEN: Of this witness?

16 MR. WATSON: Of this witness.

17 JUDGE SMOLEN: Sustained.

18 MR. SUGARMAN: I will withdraw the question. I  
19 don't have the Reese report. I think it is easier just  
20 to withdraw the question.

21 BY MR. SUGARMAN:

22 Q. You testified that not all genetic change  
23 results in cancer at page seven of your testimony, lines  
24 15 to 25. How would you go about determining whether any  
25 particular genetic change causes cancer?

1           A. There are numerous experimental techniques to  
2 ask those questions depending on what genetic change you  
3 are specifically referring to.

4           Q. What studies have been done to determine  
5 whether genetic changes lead to -- caused by E/MFs --  
6 lead to cancer?

7           A. Well, as I have outlined in my report, the  
8 first questions that have been asked, or questions that  
9 have been asked focus on whether E/MF can cause genetic  
10 change, and the answer to that is conclusively no. The  
11 second questions that have been asked are whether  
12 exposing small animals with short lifespans to E/MF  
13 results in the formation of cancers, and the answer to  
14 that question has also been no. So those are two kinds  
15 of experiments that can be done.

16          Q. That's it?

17          A. Those are the main E/MF studies that I have  
18 relied on, and that is a very short summary of what is  
19 referred to in here.

20          Q. Now, is a genetic change the only way that an  
21 agent can cause cancer?

22          A. The only way that cancer can be caused is by  
23 genetic change.

24          Q. Are changes in melatonin production genetic  
25 changes?

1 A. No.

2 Q. Are changes in cell proliferation genetic  
3 changes?

4 A. No.

5 Q. So your conclusion is that changes in melatonin  
6 and genetic changes -- changes in melatonin and cell  
7 proliferation couldn't cause cancer?

8 MR. WATSON: Asked and answered, Your Honor.

9 MR. SUGARMAN: Okay.

10 BY MR. SUGARMAN:

11 Q. How do you explain the positive findings of the  
12 epidemiological studies?

13 MR. WATSON: Objection, Your Honor. Beyond the  
14 scope.

15 JUDGE SMOLEN: Sustained.

16 MR. SUGARMAN: No further questions.

17 MR. WATSON: No questions, Your Honor.

18 JUDGE SMOLEN: The witness is excused. Thank you  
19 very much for appearing and testifying today.

20 (Witness excused.)

21 JUDGE SMOLEN: I didn't bring my calendar in, but  
22 there is a date for the exchange of the testimony of your  
23 other witnesses.

24 MR. WATSON: Yes, Your Honor. And I believe we  
25 also owe everybody here a report on Dr. Rosenberg. And I

1 think we will need some calendars in order to talk about  
2 that, the scheduling.

3 JUDGE SMOLEN: Let's take a break. We will go off  
4 the record now and discuss the scheduling.

5 (Recess.)

6 JUDGE SMOLEN: Back on record.

7 Our scheduling still stands, then. Rebuttal  
8 testimony is due from the electric company on December 5  
9 on its remaining rebuttal witnesses. If there is any  
10 problem with any one of the witnesses then we will have a  
11 telephone conference and iron that out at that time.

12 And the next set of hearings will be on December 16  
13 and 17 here commencing at 10:00 a.m.

14 Anything further?

15 (No audible response.)

16 JUDGE SMOLEN: There is no response. The hearing  
17 is adjourned. Thank you all very much for appearing  
18 today.

19 (Whereupon, at 11:58 a.m., the hearing was  
20 adjourned, to be reconvened at 10:00 a.m., on Monday,  
21 December 16, 1991, in Philadelphia, Pennsylvania.)  
22  
23  
24  
25

C E R T I F I C A T E

1  
2 I hereby certify, as the stenographic reporter,  
3 that the foregoing proceedings were taken  
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