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                            VOLUME: I
                            PAGES: 1-88
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    IN THE UNITED STATES DISTRICT COURT
   FOR THE WESTERN DISTRICT OF WISCONSIN
                  Case No. 11-CV-724-bbc
*******
KATHLEEN McHUGH and DEANNA
SCHEIDER, Individually and
on behalf of all persons
similarly situated,
         Plaintiffs
     VS.
MADISON-KIPP CORPORATION,
CONTINENTAL CASUALTY COMPANY, )
COLUMBIA CASUALTY COMPANY,
UNITED STATES FIRE INSURANCE )
COMPANY and ABC INSURANCE
COMPANIES 1-50,
         Defendants
*****
     DEPOSITION OF DAVID OZONOFF, M.D., a
witness called on behalf of the Defendant,
Madison-Kipp Corporation, pursuant to the
Federal Rules of Civil Procedure, before
Kelly G. Patterson, a Notary Public in and
for the Commonwealth of Massachusetts, at
The Charles Hotel, 1 Bennett Street,
Cambridge, Massachusetts, on Thursday,
February 7, 2013, commencing at 10:04 a.m.
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 1
    APPEARANCES:
 2
     THE COLLINS LAW FIRM
      (by Edward J. Manzke, Esquire)
     1770 Park Street, Suite 200
 3
     Naperville, Illinois 60563
     Tel. (630) 527-1595
 4
     ejmanzke@collinslaw.com
     for the Plaintiffs;
 5
 6
    MICHAEL BEST & FRIEDRICH, LLP
 7
      (by John A. Busch, Esquire)
     100 East Wisconsin Avenue, Suite 3300
 8
     Milwaukee, Wisconsin 53202
     Tel. (414) 271-6560
      jabusch@michaelbest.com
 9
     for Madison-Kipp Corporation;
10
11
    TROUTMAN SANDERS LLP
      (by Rebecca L. Ross, Esquire)
12
      55 West Monroe Street, Suite 3000
     Chicago, Illinois 60603
     Tel. (312) 759-1921
13
     becky.ross@troutmansanders.com
14
     for Continental Casualty Company and
     Columbia Casualty Company;
15
    MEISSNER TIERNEY FISHER & NICHOLS, S.C.
16
      (by Jennifer A.B. Kreil, Esquire)
     111 East Kilbourn Avenue, 19th Floor
17
     Milwaukee, Wisconsin 53202
18
     Tel. (414) 273-1300
      ibk@mtfn.com
19
     for United States Fire Insurance Company;
20
    NISTLER LAW OFFICE, S.C.
2.1
      (by Jacques C. Condon, Esquire)
     3235 North 124th Street
22
     Brookfield, Wisconsin 53005
     Tel. (262) 373-1420
23
     JCondon@NistlerLaw.com
     for Lumbermen's and American Motorists
24
      Insurance Company.
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4 PROCEEDINGS 1 2 (Curriculum vitae marked Exhibit 3 No. 1 for Identification.) 4 (Report of David Ozonoff, MD 5 marked Exhibit No. 2 for Identification.) DAVID OZONOFF, M.D., a witness 6 7 called for examination by counsel for the 8 Defendant, Madison-Kipp Corporation, having 9 been satisfactorily identified by the 10 production of her/his driver's license, 11 being first sworn by the Notary Public, was 12 examined and testified as follows: 13 DIRECT EXAMINATION 14 (By Mr. Busch) 15 Please state your name. Q. 16 Α. David Ozonoff. 17 Q. Have you been retained as an expert in this 18 matter, the McHugh matter? 19 I haven't been retained, but I have been Α. 20 asked to offer an opinion which I have 21 done. 22 When were you asked to render an opinion? 0. 23 Α. I think it was probably sometime in mid or 24 I don't remember late last spring.

5 1 exactly. 2 Who approached you? Q. 3 Α. Mr. Manzke. 4 Had you ever worked with Mr. Manzke in the Q. 5 past? 6 Yes. Α. 7 Q. In what regard? 8 Essentially, I was a witness in some cases Α. 9 that he had prior. 10 Do you recall the cases for which you were Q. 11 a witness? 12 Well, one of them was called the Lockformer Α. 13 case. I'm not exactly --14 Can you spell it? Q. 15 -- Lisle, Illinois, and then there was one Α. 16 in Indiana. I don't remember the name of 17 the case. 18 Do you recall any other cases? Q. 19 I don't, but if there was another case it's Α. 20 probably only one, but I'm not sure if 21 there was or not. 22 Was there a pollutant or a contaminant in Q. 23 the Lisle case upon which you rendered an 24 opinion?

6 1 Yes. Α. 2 What was that? Q. 3 That involved chlorinated ethylene like PCE Α. and TCE. 4 5 Was there a fate or transport mechanism in Q. 6 that case? By that I mean, was it a water 7 case, a vapor case, a ground case, or do 8 you recall? 9 I actually don't remember. Α. 10 Do you recall when that case was, when you Q. 11 were hired? 12 Five years. Four years. I'm not really Α. 13 sure. 14 In the Indiana case, was there a defendant Q. 15 in the Indiana case? 16 Α. Yes. 17 Who was that, do you recall? Q. 18 No, I don't. Α. 19 Do you recall the contamination or the Q. 20 toxic issue? 21 Yeah, I think everything I've done for 22 Mr. Manzke has been chlorinated ethylene. 23 Q. Do you recall whether there was any

particular method of transport of the

- chlorinated ethylenes in the Indiana case?

 By that I mean vapor, water, or --
- A. I actually don't remember. You know, I think it was so improperly managed so it wound up on the ground, wound up in the ground water. You know, whether the pathway to human exposure was through ground water or vapor intrusion, I don't remember that.
- Q. Let me show you what's been marked as
 Ozonoff Exhibit 1, which was proffered to
 us as your CV, or curriculum vitae. Take a
 moment and look at that, and my question
 is, is that your most recent CV?
- A. I think there is, you know, some minor changes from this.
- Q. As you sit here today, do you recall what those are?
 - A. Well, my term on the EPA Science Advisory

 Board has ended, so I think that's probably

 on here. Yes. I don't know if this says I

 was on the Faculty Senate or not but I am

 on the Faculty Senate again, and I'm on the

 Faculty Council for the University. I

8 1 think that's probably -- those are the 2 I think there's another changes. 3 publication. 4 The university of which you speak is Boston Q. 5 University? 6 Yes. Α. 7 Let me show you what's been marked as Q. 8 Ozonoff Exhibit No. 2. That's been 9 proffered to us as your report in this 10 If you take a look at it and matter. 11 confirm that that's what it is? 12 Α. Yes, I can confirm that. 13 Now, as of the date of this report, did the Q. 14 report contain all the, which is 15 November 29, 2012. As of the date of this 16 report, does the report contain all of the 17 opinions that you have in regard to this 18 matter? 19 Yes. Α. 20 Q. Since the date of this report, the 29th of 21 November 2012, have you formulated any 22 other opinions? 23 Α. No. 24 Have you been asked to formulate any other Q.

9 1 opinions? 2 Α. No. 3 As you sit here today, do you know how much Q. 4 time you spent in the work leading up to 5 this report? 6 You mean work done for this case? Α. 7 Yes, I mean this case. I don't mean your Q. 8 whole career. 9 Yes, a lot of work went into this report Α. 10 that was not related to this case. 11 How much work related to this case? 0. 12 I probably spent eight to ten hours, Α. 13 something like that. 14 Can you tell me, specifically during that Q. 15 eight to ten hours, what you did relating 16 to this case that's contained in this 17 report? 18 Well, a lot of my opinions have been Α. 19 previously written down and what I did was 20 I looked at the data involving the class 21 residences and the site that were provided 22 to me by counsel, and I looked at, you 23 know, some relative associated material, 24 like the website of the Wisconsin DNR, and

then I used the information to make the appropriate changes in what I had already prepared, essentially established knowledge about this.

- Q. Directing your attention to Page 1 of the report.
- A. Okay.

Q. At the bottom, there's a statement, and I'll just read it and then I'm going to ask you about it.

The statement is, "Reports indicate that a substantial contamination by chlorinated ethylene solvents of soil, groundwater and soil vapor occurred at the Madison-Kipp Corporation (MKC) facility located at 201 Waubesa Street, beginning decades ago and continuing until at least 1989, resulting from improper management and disposal of chlorinated ethylene solvents."

The sentence indicates that reports indicate, in particular, the fact that there was improper management and disposal of chlorinated ethylene solvents. Have you

done any independent work to ascertain the type of management and disposal of chlorinated ethylene solvents that Madison-Kipp engaged in?

- A. No, I haven't, but the fact that, you know, the groundwater and soils are contaminated with these materials indicates that they weren't disposed of properly. Exactly the details of the improper disposal, I don't know.
- Q. On the next page, Page 2, there's the statement that, and I'll just pick up at the semicolon on Page 1, "This contamination found its way into the groundwater, soil, soil vapor and indoor air at homes in the vicinity of the MKC facility and that this contamination has resulted in exposures through inhalation of chlorinated ethylene solvents (primarily PCE) to residents of these homes."

Is your opinion limited to the inhalation of chlorinated ethylene solvents in the MKC area?

A. Well, yes.

- Q. Do you know how many homes --
- 2 A. Let me just say.
 - Q. Go ahead.

- A. I hesitated for a moment because, in fact, when these solvents are in the air the principal root of exposure is through inhalation, but you can actually ingest it, so things like PCE are very lipid soluble so they can get into things like butter and olive oil that are in the house and you can ingest it that way. I'd expect that to be relatively minor in this case, but I tend to think of everything. I think this is primarily inhalation.
 - Q. Understood. Do you know how many homes of the 34 or so homes that are part of the Class have actually had reported exposures through inhalation of chlorinated ethylene solvents?
 - A. Well, I've seen the data. I can't give you a number right now. I've seen maps, for example, which have the homes in which there were detects located. I think it was probably most of them.

- Q. Are you aware that some of the homes have non-detect?
- A. Yes.

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Q. Is it your opinion that the homes that have non-detect are not exposed or --

MR. BUSCH: Strike that.

- Q. The homes that have registered non-detect do not have an unacceptable risk of cancer?
- A. So I'm not sure I understand your question.
- Q. Maybe I'll get to it another way. I
 believe it's your opinion, on Page 2, that
 the exposure to PCE in the residential
 environment presents an unacceptable risk
 of cancer; is that correct?
- 15 A. Yes.
 - Q. In the homes that have no detection of PCE, is it your opinion that they do have an acceptable risk of cancer?
 - A. If you were in an area where there's demonstrable contamination and yet there's no detectable level, I'm not ready to conclude that there's no exposure.
- Q. Are you -- do you conclude that there is exposure?

- A. I think it's likely that there is exposure.
- Q. What's the basis of that?
- A. Or at least potential for exposure.
- Q. On Page 2, the next sentence reads, "Data provided to me indicate that the concentrations of the chlorinated ethylene organic solvents in the indoor air to which residents have been, are currently, and in the future could be exposed present an imminent and substantial long term health danger." Is that your opinion?
- A. Yes.

- Q. Is there any it references the fact that the concentrations of the chlorinated ethylene organic found in the homes of the residents. What concentrations of chlorinated ethylene organic do you believe must be reached before an imminent and substantial long term health danger is presented?
 - A. It's my opinion that once you're able to measure it then it's already an unacceptable risk. The reason for that is that, in terms of the biological potential

that you have, it's plausibly reasonable, and it's certainly unacceptable, because there's no benefit to it; it only carries risk with it.

MS. ROSS: I'm sorry, I didn't hear the last of that sentence.

THE WITNESS: It only carries risk with it.

- Q. Going to the box in the opinion on Page 2.

 Is there any significance in your reportage as to the bolding and the placement of this language in a box?
- A. Not beyond the obvious one, which is it was meant to set it off so that it would be easy to see.
- Q. Okay. This really is at the core -- the boxed in areas tend to be the core of your opinions; is that fair to say?
- A. Well, I don't know what you mean by core of my opinions. I'm a scientist so I have lots of opinions on things. I think what's in the box was what I thought was pertinent about my opinions for this case, to some extent. If all I needed was what was in

the box, I wouldn't have had to have written anything else, so I'm not sure how to answer that.

- Q. Once again, the first sentence reads, "It is my opinion, within a reasonable degree of medical certainty, that exposures to PCE in the residential environment present a public health risk to the Class Area residents."
 - If I were to interpret what you said previously, that's because it's your opinion that once it's detectable, it's already unacceptable?
- A. Well, because, for this particular chemical, detectable amounts actually represent a substantial biological potential.
- 18 Q. And that's PCE?
- 19 A. Yes. It's not my opinion that once 20 anything is detectable.
- 21 Q. It's PCE?

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- 22 A. Right. I'm talking about PCE.
- Q. The next sentence says, "This risk is related to exposures to PCE and its

degradation products via inhalation through indoor air and ambient air."

Can you list for me the degradation products that you reference there?

A. Well, what happens with PCE, if you think of the chemical structure of PCE, it's two carbons connected with these double bonds, and then, like, four ears hanging off are these four chlorines. That's the tetrachloroethylene that's in its name.

What happens in the environment is that in anaerobic conditions, that is conditions without oxygen, microbes in the environment start stripping off those chlorines one by one. When you remove the first one, you're left with trichloroethylene. When you remove the second one, you're left with one of the isomers of dichloroethylene. And when you remove three of them, you only have one of the chlorines left, all the others have been replaced by hydrogen, and you have vinyl chloride. And then if you remove that one, you've gone all the way down to

- ethylene, which is a hydrocarbon. So the 2 degradation products are the anaerobic dechlorinated compounds that are produced from stripping off those chlorines.
 - Are there initials to describe Q. trichloroethylene?
 - Yeah, TCE. Α.

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- Are there initials to describe vinyl Q. chloride?
- 10 A lot of people call it VC. And 11 dichloroethylene is often abbreviated VDC 12 or DCE. VDC because it's vinylidene 13 chloride is sort of a generic name for it, 14 but it's chemical name is dichloroethylene 15 and you have to say which of the isomers.
- 16 Q. What do the initials VOC, if anything, 17 describe?
- 18 Volatile organic chemical. Α.
- 19 Are these that we just spoke VOCs? 0.
- 20 Α. They are.
- 21 Q. In the box it also indicates that it's your 22 opinion "within a reasonable degree of 23 medical certainty that the
- 24 weight-of-the-evidence favors the

proposition that exposure to PCE in the residential environment of Class Area members presents an increased and unacceptable risk of cancer to those exposed under the usual circumstances of living and working in a contaminated environment such as in Madison, Wisconsin."

And the unacceptable risk once again here is the anytime PCE is detected, correct?

- A. Well, if there's enough PCE to detect it with the usual analytic methods then the biological potential to produce harm and no benefit at all makes it unacceptable.
- Q. So PCE at any level once detected presents an unacceptable risk of cancer in your opinion?
- A. Well, that's not what I said. I said once detected then it's present at a level which presents unacceptable harm. You had those two things reversed. I'm not saying at any level whatsoever.
- Q. Once detected it presents an unacceptable risk?
- A. Yeah. But if your instruments can detect

it, then the arithmetic really has worked against you because there's quite a lot of it around once it's detected. Even though the units of detection are sometimes expressed in a way that make it sound small, like a part per billion, in biological terms, actually, that's a very large exposure because in terms of the number of molecules, which are the number of potential interactions with a cell that could produce a cancer is very, very large at that point.

- Q. Is your report limited to risk of cancer or is it broader than risks of cancer?
- A. Well, my -- I think this report is largely related to cancer. There are risks that are non-cancer risks, some of which are produced by literature that I've contributed to.
- Q. This opinion is primarily about cancer?
- A. Yeah, this is primarily about cancer, but if you want to know what my opinion is, actually, since this was written I'm much more concerned, not much more concerned,

but I am concerned about non-cancer risks, and I think that when I gave my opinions it was sort of implicit there that there are public health risks in general not completely restricted to cancer.

- Q. Have you done any analysis of non-cancer risk since your report?
- A. Well, we published about several papers and I can't remember when the last one came out because it takes awhile for these things to go through the publication.
- Q. Have you done any work in this case in regard to assessing non-cancer risks since the promulgation of your report?
- 15 A. No.

- Q. At Page 3, you reference two government-sponsored studies which you are currently the principal investigator or co-principal investigator. Can you name what those are for me, please?
 - A. Let me see which ones those are when I wrote this. I don't remember which ones they were but I'll tell you the two that exist now.

Q. Okay.

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Α. One of them is an EPA grant for which I'm a co-investigator, not a principal investigator. It's in the EPA STAR program. STAR is an acronym that stands for science to achieve results, and it's a program that EPA -- it's a grant program that EPA established, at least the part that we're involved in, to deal with issues of cumulative risk, and so the principal investigator of that, Professor Madeleine Scammell, was my last graduate student, and I'm actually very pleased to say that she's my boss now on this grant, since I'm a co-investigator on her grant, and it makes me very proud to say that.

But I also have another grant which she is on, so I'm her boss on that one, and the other grant is an NIH grant, and it's something that I've had for 17 years. It's at the Superfund Research Center, and it's a multi-project grant funded currently at the level of about 2.1 million dollars a year. I'm the program director of it.

There are maybe six or seven project leaders of which at least five of them are senior faculty members leading their own projects with me as the overall program director. There are five projects, one of which is a PCE project, and three -- five core facilities.

Q. Those two studies, two programs that you're involved in, the NIH and the STAR program, what, if any --

MR. BUSCH: Strike that.

- Q. Of the two programs in which you are involved, the NIH grant and the STAR program, do any of them relate to PCE or its degradation products?
- 16 A. Yes.

- Q. Which ones and how?
 - A. Well, the Superfund Research Center has an entire project devoted to PCE, and that's been going on since probably the late 1980s, and it's an environmental exposure to PCE and almost only through drinking water, and we publish many papers for them.
 - Q. Are some of those the ones that are listed

- on Page 6 in the Footnote 3?
- A. I'm not sure if this has all of them.
 - Q. At least some of them are?
 - A. Yeah, probably most of them. There may be one that's not on there because it came out after this. I'm not sure.
 - Q. In regard to the NIH grant, is there any specific study that's being done with regard to PCE in which you were involved?
- 10 A. This is the NIH grant.
- 11 Q. How about the STAR?
- 12 A. The STAR grant is a methodology grant.
- 13 It's more theoretical and it has
- applications to PCE but it's about
- cumulative risk to all sorts of things in
- the environment.
- 17 Q. In this matter, have you been asked to
- render any opinions in regard to PAHs?
- 19 A. No.

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- Q. Have you been asked to render any opinions in regard to PCBs?
- A. No. I know that there are PAHs and PCBs there and I have opinions about them.
 - Q. You didn't report them in your report, did

you?

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- A. No -- well, I wasn't aware of any exposure pathway to the residents here so I didn't actually address that.
- Q. You have not been asked to render any opinions with regard to PAH or PCB, correct?
- A. No, I haven't, but of course whether I will give opinions about it, I'm not completely in control of because you may ask me for my opinion.
- Q. You haven't been asked by plaintiffs in this case to render opinions on PAH or PCBs?
- 15 A. No. I could possibly be asked by you, I suppose.
- 17 Q. Directing your attention --
- A. While we're stopped for a second. I like to stop once an hour because I have bone spurs in my neck.
- 21 Q. You control whatever you want.
- 22 A. I know. We're a long way from that.
- Q. You control the whole thing, sir.
- A. Okay, then let's go home.

Q. Which I'm sure is a rarity in your life.

On Page 7, you have a discussion that continues about the weight-of-the-evidence methodology; do you see that?

A. Yes.

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- Q. Did you employ weight-of-the-evidence methodology in arriving at your opinions in this case?
- A. Yes.
- Q. Did you use any other methodology?
- 11 Well, you know, weight-of-the-evidence Α. 12 methodology is sort of a term of art for a 13 lot of different things, which includes 14 making judgments about the evidence and 15 which pieces to weigh, how much importance 16 you give them in your decisions, and I'm 17 not speaking quantitatively there, but 18 qualitatively, so I used lots of other 19 methodologies in pursuing the 20 weight-of-the-evidence.
 - Q. Those are the ones that you discuss at some length in this report?
 - A. Well, I discuss quite a bit the nature of scientific method and scientific judgments

and then I employ them.

- Q. Directing your attention to Page 17. One of the issues that this report addresses is the question "Can chlorinated ethylene solvents cause cancer in human beings?" Do you see that?
- A. Yes.

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- Q. In opining on that, did you use the weight-of-the-evidence methodology?
- 10 A. Well, that's not -- yes, I think the short
 11 answer to that is yes. Its got a more
 12 complicated long answer.
 - Q. Did you use the weight-of-the-evidence methodology in arriving at any opinion other than the one that "Can chlorinated ethylene solvents cause cancer in human beings?"
- 18 A. In this report you mean?
- 19 Q. Yes, I'm sorry, in this report.
- A. Well, I think the answer here is -- I was going to say the answer is yes but now I don't remember what the question was.
- MR. BUSCH: Can you read back that question?

(Previous question is read back by the Court Reporter.)

- Q. I'll restate it. Did you use the
 weight-of-the-evidence methodology in
 arriving at an opinion other than "Can
 chlorinated ethylene solvents cause cancer
 in human beings?"
- A. Well, I do use weight-of-the-evidence methodology for arriving at my scientific opinion. To the extent that I have given scientific opinions in this report, that's what I did.
- Q. Okay.

(Discussion off the record.)

- Q. At Page 21, you make the statement that
 "Toxicology is an experimental science,
 while epidemiology is an observational
 science." Does that observation play any
 role in your opinion?
- A. Just for the record, there's also a footnote there that suggests that there are possible exceptions with respect to epidemiology.
- Q. Okay.

- A. Does this play a role? I'm not sure what you mean by "play a role."
- Q. Well, do you view your opinion -- you view your opinion in this matter as an epidemiological opinion as opposed to a toxilogical or both or neither?
- 7 It's a scientific opinion. Α. 8 epidemiologist but I do use toxicology --9 there is a branch of epidemiology that 10 could be called experimental, so that's 11 part of my professional expertise, but most 12 of the evidence that we're talking about is 13 not in epidemiology, it's from the 14 observational portion of epidemiology, and 15 I am primarily an observational 16 epidemiologist.
- 17 Q. The methodology that you use in
 18 observational epidemiology is described, at
 19 least in part, in your report, correct?
 - A. Yes, in part.

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- Q. Is there any part of observation or of
 epidemiology that's important for your
 report that's not contained in your report?
 - A. No, I don't think that's important for my

- report. There's quite a lot that's not here. I'm writing a book now on the subject. But I don't think it affects any of the opinions here.
- Q. At Page 41 -- excuse me, Page 40 of your report, you reference at Paragraph D,

 "Relationship with time," and in

 Paragraph E, "Dose-response relationship".

 Do either of those, "Relationship with time" and "Dose-response relationship" bear on your opinion in this case and if so how?
- A. Well, my opinion here is not a specific causation opinion, it's a general causation, and it's not it's about the ability of these chemicals to do certain kinds of health effects, so these bear upon the interpretation of epidemiological studies, as described here, and I don't know what to say beyond that.
- Q. It certainly comes into play but your opinion is not reliant upon any particular dose-response or relationship with time; is that fair to say?
- A. Yes, except in so far as those things are

related to the interpretation of the studies that are considered in this report.

Q. Okay. Directing your attention to Page 48.

There's a statement, "It is my opinion,

within a reasonable degree of medical

certainty, that exposure to PCE in the

residential environment presents a public

health risk to the Class Area. This risk

is related to exposures to PCE and its

degradation products."

How, if at all, does that opinion differ from the opinion set forth on Page 2?

- A. I think it's saying it's the same general idea in different language.
- Q. At Page 68, in the box, there's a statement that "At the very least, it is clear there is independent, informed, scientific opinion that accepts the proposition that TCE and PCE are probable human carcinogens."

You italicized the word "probable"; do you see that?

A. Yes.

- Q. In your opinion, is there a difference between the use of the word "probable" and "likely"?
- A. No. At least that's not my understanding there's a difference in EPA's language, and I think in ordinary parlance there isn't either.
- Q. Much of your work at Boston University and through grants has been relating to exposure to PCE in drinking water, correct?
- A. Didn't you just say how much.

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- 12 Q. No, I just made a statement. Is it correct
 13 that much of your work over the past
 14 several years at Boston University and
 15 otherwise has been in regard to exposure to
 16 PCE in drinking water?
 - A. Yes, probably the last 25 years.
- Q. Is the primary means of ingestion in those studies the actual consumption of water that has PCE in it, as opposed to vapor that may come from the water?
- 22 A. It's hard to say. Of course a lot of
 23 estimates are that when you have all of the
 24 organics in drinking water that about half

- of the exposure may be through inhalation, but that varies from setting to setting.
 - Q. Have you done any --
 - A. And there's dermal exposure, too.
 - Q. Have you done any studies isolated on PCE and its degradation bi-products -- that's a bad term.

MR. BUSCH: I'll strike it.

- Q. Have you done any studies on PCE, DCE or TCE limited solely to vapor being the means of ingestion, inhalation?
- 12 A. No.

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- Q. At Page 137, in the last paragraph, you reference some testimony from Michael

 Schmoller and some information from John Hausbeck referencing mitigation systems.

 Do you see that?
- 18 A. Yes.
- Q. Are you aware of the types of mitigation systems that are being offered to certain residents in the Class Area?
 - A. From what I recall from descriptions that this is -- I can't remember exactly what the exact term is, sub-slab ventilation or

exhaustion or something like that.

- Q. Have you had any or have you studied at any point in time the efficacy of such sub-slab mitigation systems?
- A. No. I say that our Superfund Center, not me personally, but the center and the program I direct, does do vapor intrusion work.
- Q. Your opinion in -- you have not been asked to render nor are you rendering on opinion on the efficacy of sub-slab mitigation systems as a means of addressing vapor intrusion, are you?
- 14 A. No.

- Q. Directing your attention to Page 138.

 There's a -- the first phrase in the first sentence says that "current uncertainties do not allow precise estimation of cancer risk from exposure to PCE and potentially TCE and VC in the residential environment at levels seen in the Class Area." Do you see that?
- 23 A. Yes.
 - Q. Can you list for me the current

uncertainties of which you refer?

- A. Just about everything that goes into making these kinds of estimates. The biological mechanistic bases of the models, the parameters used in the models. The uncertainty in the inputs into the models and the fact that the models produce expected values and many of them don't produce distributions of possible risks.
- Q. Excuse my ignorance, but can you be more -can you elaborate a little bit more on what
 you mean by "failure to produce
 distributions"?
- A. So they tend to produce expected values or average values, in layman's terms. So if you have two people, one who is five feet tall and one person who is six feet tall, their average is five-foot six, but nobody in that sample is five-foot six feet tall, so the distribution is five feet and six feet. The average is five-foot six.
- Q. What, if anything, do you believe could be done to eliminate the uncertainties that you believe to be current in that Class

Area?

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- A. Eliminate exposure.
- Q. The exposure which we talk about are the detected exposures, correct?
- A. Well, I'm saying eliminating exposure.
- 6 Q. At any level?
 - A. Yes. That would eliminate the uncertainties, if that's the question.
- 9 The last sentence indicates that the, Q. 10 or states that it's reasonable and 11 supportable "for residents of the Class 12 Area to believe that the measured levels of 13 PCE, TCE and VC contamination of their 14 groundwater, soil, soil vapor and indoor 15 air presents them with an excess risk of 16 cancer not balanced by any benefit and 17 could be considered unacceptable by a 18 reasonable person."

In the context of this report, what do you mean by "excess risk of cancer"?

- A. Cancer that's attributed to the exposure to PCE.
- Q. At any level above that which would be there in its absence?

A. No, not necessarily.

- Q. What makes it excess?
- A. Well, first of all, if you can measure it, then there's plenty of it around, because our instruments are not that sensitive that we can get down to levels that don't have, I would say, biological potential of public health significance.
- Q. So once again, the fact that it's measured makes it in excess?
- A. No, the fact that the level at which it's measured makes it an excess. If we had instruments that were maybe a thousand times more sensitive, you might be able to get down to a level at which people would say -- I don't know.
- Q. But based upon the fact that with the current level of instrumentation that it can be detected, that in and of itself represents an excess risk?
- A. Yes, I think that's primarily a question of arithmetic, and I think in this report, I went through that arithmetic, and essentially it's because molecules are very

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small and a microgram of PCE has got an awful lot of molecules. Each of those molecules has got some biological potential to cause some harm, but if there were a handful of them, maybe a million of them or ten million or a hundred million, but we're talking about one with fifteen zeros after We're talking about very, very, very large numbers of potential and biological interactions, and that's purely a function of the fact that what chemists refer to as Avogadro's number. It's the number of molecules in one gram molecular weight of a chemical, and it's a huge number. 6.023 times ten to the 23rd. That's one with 23 zeros after it.

So if you have even a fraction of this, say one billionth of a mole gram molecular weight, then you still have one with 15 zeros after it or 14 zeros after it. It's an incredibly large number. The fact that a part per billion doesn't sound very big, that's just a function of the unit that's being used, and if you use

39 1 units of molecules, then that number 2 suddenly is a very, very large exposure. 3 MR. BUSCH: This would be a good 4 time to break. We're an hour into it. 5 THE WITNESS: Sure. That's 6 perfect actually. 7 MR. BUSCH: Okay. 8 (Recess.) 9 Doctor, do you know what regional screening Q. 10 levels are from the EPA? 11 You mean what the levels are? Α. 12 No, just generally the concept of regional Q. 13 screening levels? 14 Α. Yeah. 15 What do you understand a regional screening Q. 16 level to be? 17 They are -- my understanding is that Α. 18 they're sort of -- well, it depends a 19 little bit on what the relationship of EPA 20 to the state is as to whether the state has 21 prelacy or not, but they're some kind of 22 guidance or direction to people who are 23 trying to deal with environmental problems 24 as to when they should take certain

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- Q. Do you agree that exposures below regional screening levels can be considered not to present toxicological concerns?
- A. Well, since regional screening levels differ from region to region, that can hardly be true.
- Q. Assuming that all regions agree as to an appropriate screening level, do you agree with the proposition that exposures below screening levels can be considered to not present a toxilogical concern?
- 13 A. No. EPA doesn't believe that and neither do I.
 - Q. Did you consider at all in your opinion the site specific dose and duration of exposure?
 - A. I'm not sure what you mean by that.
- 19 Q. Did you consider site specific information 20 in that part of your opinion that addresses 21 dose-response?
- 22 A. I actually don't understand the question.
- Q. In your opinion, you do take into consideration dose, correct?

- A. You mean specific doses?
- 2 Q. Yes. Or do you not?
 - A. I take -- well, first of all, there is no risk if you're not exposed.
 - Q. Okay.

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- A. And what I -- I took dose into account to the extent that we've already discussed, which is that if you can see it, then we're talking about a biological potential here that concerns me as a public health scientist, so to that extent the answer is yes, I took it into account in that sense.
 - Q. Did you take into consideration or into account the frequency and duration of exposure?
- 16 A. Yes, I think so.
- 17 Q. How?
- A. That when you're living in a house, the
 frequency is daily and the duration is the
 amount of time that you spend in that
 environment, so when I talk about risk to
 people living under ordinary circumstances,
 or whatever the exact language was, I was
 referring to frequency and duration.

Q. By the way, of the eight to ten hours that you spent in compiling the report, how much of it did you spend in reviewing the site specific data, do you know?

- A. Well, you know, for example, not for example, but I review that because I wanted to take what I had written about PCE and make it appropriate to the setting, so I needed to see what the setting was.
- Q. But if the total amount of time spent was eight to ten hours, how much of it was in reviewing the data?
- A. Probably at least half of it. I can't give you an exact. I wasn't doing one thing all at once. I would go back and forth.
- Q. Would you agree with the definition, the following definition, that risk assessment is the characterization of the potential adverse health effects of human exposures to environmental hazards?
- A. Well, I don't think I object to it. I
 think one could probably come up with
 different definitions of risk assessment.
 I think that probably describes a lot of

what's done.

- Q. Did you engage in risk assessment in formulating your opinions as set forth in the report?
- A. So when you -- you're saying risk assessment now, you're specifically referring to this definition?
- Q. Let's go back. Do you use the term "risk assessment" in your practice?
- 10 A. Yes.

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- Q. Would you define "risk assessment" for me as you use it.
- A. Well, when I've done risk assessments and
 when I hear other people talking about it,
 they usually are talking about some kind of
 point or interval estimate using one or
 another kind of a model, so quantitative
 estimate, and a risk is a probability.
 - Q. Did you engage or did you undergo -MR. BUSCH: Strike that.
- Q. Did you perform a risk assessment in rendering your opinion as set forth in Exhibit 2?
 - A. No, I didn't perform a quantitative risk

assessment, that is to say a point or interval estimate of average risk.

Just to add to that. I did perform an assessment of risk. I assessed the risk, but if you want to put — if you want to put the word assessment after risk then you're referring to a particular kind of operation, but I think my report is really an assessment of risk.

- Q. On a qualitative as opposed to quantitative basis?
- A. It's not purely qualitative. When you talk about quantitative basis, in the context of risk assessment, you're talking about a point or interval assessment of a probability.
- Q. You did not do that in this case?
- A. I did not do that, no, but I did other quantitative things. For example, there's a fairly complete review of quantitative aspects of the literature up through 2003 or so.
- Q. Did you use at all in your opinion or reference at all or take into consideration

45 at all the EPA's screening level of 9.4 1 2 micrograms per cubic meter for PCE? 3 That refers to what? Α. 4 The EPA screening level. Q. 5 For what? Α. 6 PCE. Q. 7 Well, are you talking about soil, soil gas, Α. 8 sub-slab, indoor air? 9 Excuse me, vapor. Indoor air. Q. 10 Indoor air? Α. 11 0. Yes. 12 Screening level of what? Say it again. Α. 13 Q. 9.4 micrograms per cubic meter. 14 Well, the Massachusetts screening level is Α. 15 .21 parts per billion, so a part per 16 billion is about seven micrograms per cubic 17 meter so talking about 1.4. 18 Q. 9.4? 19 1.4 parts per billion screening level, I Α. 20 believe, is what it is in Massachusetts, 21 micrograms. 22 Whatever the screening level is that the Q. 23 EPA adopts, it was not specifically used in 24 your report or referenced in your report

that I saw; is that correct?

A. No, it wasn't.

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- Q. Do you agree that indoor air typically contains volatile organic chemicals, including PCE, from consumer products, building materials, and outdoor air?
- A. Yes, it often does.
 - Q. Is indoor air concentration resulting from these sources commonly called background?
- 10 A. Yes, I think commonly but probably
 11 inappropriately called background.
- Q. Do you know, for example, some of the sources from which background PCE may emanate?
- 15 A. Yes.
 - Q. Give me some examples, if you would.
- A. Well, PCE is used in dry cleaning. It's found in some kind of products like drain
- cleaners, you know, other household things.
- I don't know what they all might be. Most
- of the dry cleaning exposure is gone by the
- time you get the clothes home but it
- contributes to urban background.
 - Q. That is the more concentrated the

- population the more background PCE, as a general proposition?
- A. May or may not be. It depends upon local conditions. So many dry cleaners are now moving away from PCE because of its toxicity so my dry cleaner no longer uses PCE.
 - Q. Is PCE a banned substance from any use in the United States?
- 10 A. Well, it will be -- in California I think
 11 it's going to be banned for dry cleaning
 12 use. If not already, in a year or two, but
 13 it's not yet banned but likely will be in
 14 the not too distant future.
 - Q. Do you know if it's banned in Wisconsin for use in dry cleaning?
- 17 A. I don't know.

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- 18 Q. Is it banned in Massachusetts for use in dry cleaning?
- A. Not yet. Actually, I think Los Angeles
 county is the only place where such a ban
 has actually been put into effect or about
 to be put into effect, but Los Angeles
 county is bigger than most countries in the

48 1 world so. 2 Do you know if the use of PCE is banned in Q. 3 various cleaners and cleaning substances? 4 Α. Not that I'm aware of. 5 Is it banned at all in any application to Q. 6 your knowledge? 7 Well, I think we just talked about dry Α. 8 cleaning. 9 In Los Angeles but how about nationwide? Q. 10 Α. Not yet. 11 Are you aware that a study was done by the Q. 12 United States Environmental Protection 13 Agency in regard to background indoor air 14 concentrations of volatile organic 15 compounds? 16 Α. Yes. 17 It was promulgated sometime in 2011? Q. Well, there have been numerous studies. 18 Α. 19 Q. Are you aware of one that was promulgated 20 in 2011? 21 I don't know what you mean by 22 "promulgated". 23 Published. Q. 24 Α. No.

- Q. To your knowledge, is there an estimated level of PCE nationwide that's deemed to be background?
- A. You mean an ambient outdoor air or indoor air?
- Q. Indoor air.

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- A. Well, I think my general impression that the 50th percentile in a distribution for indoor air concentration is somewhere around half a part per million billion volume.
- Q. What is the significance, from your perspective of being in the 50th percentile?
- A. It has no particular significance other than it's one of the places in the distribution that's frequently used as a marker. It's the median.
- 19 Q. Does -- when it is -- when it's expressed 20 in terms of the 50th percentile, what is 21 meant by that from a lay perspective?
- 22 A. It's the median.
- Q. So the median of indoor air background of PCE is what again?

- A. I haven't looked at this for a bit, but my recollection is somewhere around a half part per billion as a volume measurement, so that means that 50 percent of households will have that or less.
- Q. Is a half part per billion a measurable level of PCE?
- 8 A. Yes.

- Q. Another way to put it is, that's a detectable level of PCE?
- 11 A. Yes.
- Q. Does that mean that, on average, I know you don't like to -- I won't say that. That the level -- does that mean that 50 percent of the houses have one half part per billion or that all houses have, on average, a half a part per billion?
 - A. It's not an average, it's a median, and that's an extremely important difference.
 - Q. In the context of this, the median is the mid-point number, correct? It means that half of the detections -- excuse me, that the highest, the mid-point between the highest and the lowest detection, is that

the median in this context?

- A. Well, no, it includes all the non-detects, so supposing that you had 100 measurements and 49 of them were non-detects and the 50th was a half part per billion, then that would be the median. In other words, you take all the measurements and you line them up in order and you go halfway down the line, so it doesn't take into account the distribution at all.
- Q. Are you aware of any studies that take into account the distribution of PCE?
- A. Yeah, the problem -- there's a different kind of problem there because the non-detects are not zero. Some of them may be zero but a lot of them aren't, so in order to figure out what the non-detects are, you have to make an assumption about what the underlying distribution of the data that it might be.

So there's different ways to do it.

One of them is you can take all the

non-detects and call them zero. I think

what EPA frequently does is they fit a

lognormal distribution to it. That's a bell-shaped curve which has been transformed logarithmically, so it's now skewed, and they fit that and assume that a lot of the non-detects are — there's stuff there, but it goes according to the lognormal distribution. That's not a bad way to do it but it can produce certain kinds of bias when you do it, and you don't really know what the measurements are below your level of detection, so that's kind of a long-winded way of saying we don't know.

- Q. I appreciate that. I believe you said that one of the more prevalent uses of PCE, at least here to for, has been in the dry cleaning industry?
- A. Yes, that and degreasing are probably the two principal uses.
- Q. Assume for the moment that my laundry, the laundry that I use to do my shirts, for example, uses PCE, and assume that I wear five laundered shirts a week and every two weeks I take them to the laundry and I pick them up and put them in my car and I

drive ten shirts that are laundered in PCE or have some PCE component in them from the dry cleaning. Am I, as you understand it, am I exposed during my car ride to a detectable level of PCE?

A. The data that I've seen, and I haven't looked at it for awhile -- well, first of all, my advice to you would be to find another dry cleaner because a lot of them are moving away from PCE not because so much the risk to consumers, although consumers don't like it when they find out, but the risk to the workers.

So the answer to your question is that the data that I've seen in the past, when people weren't quite as careful with PCE, was that if you had dry cleaning, let's, say not your shirt but your jacket, your suit jacket, and you took it home on a very hot day wrapped up in plastic from the dry cleaners, that in a certain percentage of them there might be some measurable PCE in your car from that, but mainly not.

That's not -- my impression, that's not a

- significant exposure. I don't think there's probably anything to speak of from shirts.
- Q. But it's mostly those items that are truly dry cleaned, like suits and woven fabrics?
- A. Yeah, and of those, only under special circumstances would there be a brief exposure under not well-defined circumstances, like really hot days and only from some dry cleaners. Dry cleaners differ. So you might bring it home from one place and there might be no exposure from another place, and now that they're using the transfer method, there's not as much exposure that way.
- Q. Have you taken any position at all publicly in regard to the desirability of banning PCE from all use in the United States?
- A. It's my opinion it should be banned from all use. Have I ever taken a public position on it? I can't remember. If anybody asked me about it, that's what I would say. I think I and a lot of people consider it an unreasonably dangerous

- product in the sense that you don't need it.
- Q. Are you familiar with the U.S. EPA's vapor intrusion screening level calculator that was published in March of 2012?
- A. Well, I mean, I have looked into what EPA is doing on vapor intrusion a little bit, so I don't know that they have actually publicly put anything out there. There was a leaked graph vapor intrusion that inside EPA had, but I don't think that's up on their website. I think it has either been withdrawn or so the answer is, I know that there is something, but I don't think it's really out there.
- Q. Whether it's out there or not, you did not use an EPA vapor intrusion screening level calculator in coming up with your opinions, correct?
- A. I did not.

- Q. Did you read the expert report of Barbara
 Beck?
- 23 A. I only took a briefest glance through it.
 - Q. You've not been asked to rebut any of her

opinions, have you?

- A. No. Well, to be honest with you, from what I quickly saw from what she said, she seems to agree with me, but I can't say that from a detailed reading of it. I expect that, you know, what she was asked to do is criticize me, and I was not surprised to see, but her bottom line seems to be the same as my bottom line; this is a likely cause of cancer in human beings, or it's likely to cause cancer in human beings.
- Q. You have not been asking to rebut any of her specific opinions?
- 14 A. No.

MR. BUSCH: I want to take
five minutes. I may be able to eliminate
some of this stuff.

(Recess.)

- Q. Doctor, in your opinion, are there any members of the Class who are not exposed to an unacceptable risk of cancer?
- A. Well, just looking at the environmental setting here, the environment that's substantially contaminated and the

57 1 groundwater and the soil and in the air, 2 and I think you have to be worried about --3 it's reasonable to consider that there's a 4 risk of harm to anybody who lives bordering 5 on this facility. This is pretty close 6 quarters. 7 Have you been to the site? Q. 8 No, I haven't. Α. 9 Have you interviewed any of the homeowners? Q. 10 Α. No. 11 Other than discussions with your 0. 12 attorney -- excuse me, with the attorney 13 for the Class and with your review of the 14 information provided to you, have you 15 talked with anyone else? 16 Α. You mean specifically about this case? 17 About this case. Q. 18 I have colleagues. I ask them about stuff. Α. 19 0. Not about this case? 20 Α. No. 21 MR. BUSCH: I'll pass the baton to 22 the others. 23 CROSS-EXAMINATION 24 (By Mr. Jacques Condon)

- Q. Doctor Ozonoff, my name is Jacques Condon.

 I just have a few follow-up questions. Can you pull out Exhibit No. 1, which is your

 CV. I noticed in here -- you described yourself as an epidemiologist, correct?
- 6 A. Yes.

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- Q. For awhile you were in the staff at the Department of Neurology at the Boston VA Medical Center?
- 10 A. Yes.
- Q. What's the difference between neurology and epidemiology?
- 13 A. They're completely different disciplines.
- Q. What are they? Can you explain the difference?
- A. Neurology is the clinical discipline about
 diseases of the nervous system, and
 epidemiology is a methodology for
 understanding determinants of distribution
 of a disease in a population.
 - Q. When you were at Cornell, was your emphasis in neurology, epidemiology?
- A. Are you asking me why I was in the Department of Neurology?

Q. Yes.

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There's a very simple answer to that Α. question, which is that I was the co-director, along with a colleague who was a neuropsychologist, of the Boston Environmental Hazard Center, which was the principle Gulf War research center for the Department of Veterans' Affairs. were located at the Veterans' Hospital and I was given an appointment on the staff of the hospital, which meant that I could see patients if I was so inclined, which I wasn't, because I'm not a diagnosing or treating physician at this point, although I'm licensed to do that. It was essentially just an administrative slot for me as the director of this center in a clinical facility, and the reason it was in the Department of Neurology was because my colleague is a neuropsychologist. actually succeeded me in the department at Boston University.

Q. So it was more a circumstance of being part of the VA that you're listed under the

Department of Neurology?

Α. Yes.

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- The report that has been marked as Q. Exhibit 2, you said in your earlier testimony that came from either prior versions or it came from other sources; is that right?
- It is in part, which is this is a report Α. 9 that I sort of developed over a period of time because this is what I do is chlorinated ethylenes, and I wanted a way 12 to explain this, not only to explain 13 chlorinated ethylenes, but to explain the 14 whole process of how we understand these 15 A lot of people have read this, so things. it's not like you're the only one to have 17 read it, but it's also useful in 18 circumstances like this and so each of the 19 circumstances like this that I've used it 20 with have had specific parameters to them, 21 and so I make the changes that are 22 appropriate to that.
 - Q. There's a lot of background material in terms of methodology,

weight-of-the-evidence, other things in
this report, right?

- A. Yes. Actually, one reason is because it has become important when offering opinions these days to explain exactly how you arrived at your opinion, and I think that I took a lot of care to explain that and that's applicable to lots of different cases, not just this one.
- Q. The opinions in some of the background material that's in your report, have you published that separately?
- A. No. Well, I'm an academic, so I write

 papers and I'm sure that these ideas appear

 in other forms in different ways or they

 were first part of papers and appear here.

 I'm writing a book now on mathematical

 foundations of epidemiology and obviously,

 this is part of that.
- Q. When you sat down to prepare this report, were you taking it from one or two sources, did it come from different papers; how did you come up with what we have as a 140 page --

- A. You mean the origin of this?
- 2 Q. Right.

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- 3 A. My head.
- 4 Q. Over time.
- A. Yeah. It's original with me. I wrote it
 myself. I didn't -- it's not copied or
 taken from another source, except where
 cited. I cited everything.
 - Q. In the eight to ten hours you spent looking at things in this case, did you write 140-page report in that time?
 - A. No, I think, as I described, I essentially spent the time in this case finding those things which were necessary in order to make this relevant.
 - Q. You took what you thought was necessary to make it relevant, you inserted those into this report; is that right?
- A. I adapted this report so that it addressed things that are relevant to this case.
- 21 Q. What source did you adapt it from?
- A. You know, as I described, the data on residents and site specific --
 - Q. No, I mean -- sorry. We're not

- communicating very well. You say you adapted it. Does that mean you took down -- you had a report already, you took things out --
- A. Yes, I had a report already that had lots of stuff in it, and, in fact, there are things that I've written in the past that talk about autoimmune disorders and birth defects, which could very well have been in this one.
- 11 Q. The report that you had already, what case was that?
- 13 A. Its been used in a number of cases.

 14 There's a case out in Burbank. I can't

 15 remember what the caption was.
 - Q. You talked about the Indiana case and another case. Were those reports, would they look similar to what I see in Exhibit 2?
 - A. Yes, they would.

- Q. Same information with the exception of information that would be case specific, right?
 - A. Probably pretty much so. I can't remember

exactly.

- Q. I believe those cases were five or six years ago, or what was the timeframe of those?
- A. Something like that. There's more

 up-to-date citations in this one, but it's

 not systematic. I do, obviously, keep

 track of the literature because this is

 what I do for a living, PCE epidemiology,

 and there are lots of citations in papers

 that I've co-authored on that have come out

 in this period. I don't know if they're

 all cited in here or not.
- Q. If you go back to Exhibit No. 1 and look at Page 8. Look at the very top. There's something you published in the New England Journal of Medicine. This goes back awhile, 32 years ago. "Artificial Sweeteners and Bladder Cancer." Did you come to a conclusion in that article?
- A. Yeah. This was actually a response to an article written by Morrison in New England Journal of Medicine, a case control study. The artificial sweetener involved was

saccharin, which was actually banned under the Delaney Clause. It was a comment on Morrison's study, and he and I ran into each other, unfortunately he passed away a number of years ago, but he and I ran into each other and I said, "I wrote that because what you said was going to be misunderstood." He said, "It's not my job to teach people."

- Q. What was your conclusion?
- 11 | A. Well --

- Q. Thirty-two years ago, what was your conclusion?
 - A. I'm guessing that you've read it more recently than I have since I read it 32 years ago when I wrote it. I can't remember exactly what the issue was anymore, to be perfectly honest.
 - Q. Do you recall whether you were -- either the article you were commenting on or your comment was negative towards saccharin?
 - A. Yeah, I thought that the saccharin ban under the Delaney Clause was reasonable.
 - Q. If you go to Page 11. Let me know when

you're there.

A. Yeah.

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- O. You're there?
- 4 A. Uh-huh.
 - Q. Look at No. 64. It talks about "Cancer in the Vicinity of a Department of Defense Superfund site in Massachusetts," and this was something that was apparently published in a Toxicology and Industrial Health publication. Do you see that?
- 11 A. Yeah.
- 12 Q. Do you recall if you reached a conclusion
 13 in what this particular article or whatever
 14 this was?
 - A. Well, reach a conclusion. We reported an association.
- 17 Q. Association of what?
- A. A statistically significant association
 between breast cancer and, I think it might
 have been lung, and these mortar training
 positions on Otis Air Force Base on Cape
 Cod.
 - Q. You said there was a statistically significant correlation?

- A. Association, yes.
- Q. Is that based on studies that you reviewed or what was that?
 - A. It was based on studies we did.
 - Q. When you say "we," was it you, part of a grant, what was it, if you can recall?
- 7 A. It was part of a grant and, you know, those are my co-authors listed with me.
 - Q. Was it a grant from governmental --
- 10 A. Yeah, it was either the Commonwealth of
 11 Massachusetts or NIH and I, on that date, I
 12 don't remember exactly who the funder was.
- 13 Q. Okay.

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- 14 A. Just to explain -- do you want me to
 15 explain what it was about or you don't
 16 care?
- Q. Let's move on. If you can go to your report, which is Exhibit 2, and in particular I want you to look at Page 6.
- 20 A. Okay.
- Q. In the large footnote number three, if you go eight lines down, there's a reference to "Cancer risk and residential proximity to cranberry bog cultivation in

Massachusetts." Do you see that?

A. Yes.

- Q. Are you familiar with this cancer risk in residential proximity to cranberry bog in Massachusetts?
- 6 A. Yes, I'm co-author of it.
- 7 Q. What was going on?
 - A. Cape Cod, which is where we've done a lot of work, and this was either funded by Massachusetts or NIH. I think it was --
 - Q. You said NAH?
 - A. NIH. I think it was the Commonwealth of
 Massachusetts at this point. There are two
 states, maybe three, actually, Wisconsin
 may be one of them, that produce
 cranberries so a cranberry bog is like a
 giant pool full of cranberries, and in
 order to grow them, they put pesticides on
 them, and often that's done through the
 water. It's call chemigation. At one
 point it was done by airplanes, aerial
 spraying of cranberry bogs.

Now people live right along there, their houses border on the cranberry bogs,

and so we used a drift model for that have been used by the pesticide people about how pesticides drift away when you're spraying them.

- Q. Just so I'm clear, when you say "drift model," is this an actual physical model or more a model from a scientific --
- A. I'm not sure what you mean by a physical model.
- Q. When you say "drift model," what is a drift model?
- 12 Α. It's a, in this case it was an equation 13 predicting how pesticides drifted when you 14 spray things, although we did something, 15 now that I'm telling you, we used 16 information on drift models, but we 17 actually used a buffer around the cranberry 18 I think it was 2500 meters, bogs. 19 something like that, so we compared the 20 cases of brain cancer within that buffer 21 and outside that buffer zone, and that's 22 where this association came from.
- Q. Was that also a grant?
 - A. Yeah.

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- Q. When you're part of this grant proposal and you're doing your research and you create a buffer around the zone, are you there literally taking samples or how does that work?
- A. Okay. It was not a grant proposal, it was a grant. A proposal is how you get the grant.
- Q. Thank you.

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- Α. We were funded to do research by the Commonwealth of Massachusetts on cancer on Cape Cod, different kinds of cancer. Ι think there were seven different kinds of cancer. One of them was brain cancer. one of the things we decided to look at was whether living near cranberry bogs, because they're sprayed aerially, was related to brain cancer, and there was actually a pretty strong association with brain cancer living close to the bog and the vicinity of the bog, so this was a study that we did using the state's cancer registry and interviewing people.
 - Q. What was your ultimate conclusion on the

spraying?

- A. There was a relative risk, something like four, four and a half, living close to the cranberry bog.
- Q. What do you mean by "relative risk"?
- A. In other words, the risk living close to the bog compared to living farther away from the bog.
- Q. When you say "relative risk," that's a percentage?
- A. I presume too much, I'm sorry. So if I
 were to ask what the relative risk of, say,
 being in this room versus not being in this
 room. I would take the risk, a measured
 risk of being in this room and compare it
 to the measured risk outside the room and
 take their ratio. That's the relative
 risk. So a relative risk of ten would mean
 that it was ten times riskier to be in this
 room than outside this room.

A relative risk of four for cranberry bogs means that it was four, four and a half times riskier to live within 2500 meters of the cranberry bogs than to live

- outside of that 2500 meters by part of the study group, which were the members who lived on Cape Cod?
- Q. In that particular study, you came up with the relative risk numbers?
- A. Yeah, it was estimated with something else called an odds ratio.
- Q. Odds ration, I saw that in your report.

 You refer to it as OR.
- 10 A. Yes.

- 11 Q. There is some odds ratios related to births
 12 and other things in some reports?
 - A. Yeah, so often you can't measure a relative risk directly because of the way your observations are collected, and if you use a study design called a case control design, you don't actually get the relative risk, you get something called an odds ratio, which is the odds of having the disease if you're exposed compared to the odds of having the disease if you're not exposed, but it turns out when the risk of getting the disease is relatively low, less than ten percent or less than one percent,

- then the odds ratio and the risk ratio are basically the same thing.
- Q. There are other references, and you just mentioned in the Cape Cod that it sounds like it was an extensive study in Cape Cod?
- A. We have been studying them probably for 20 years, maybe longer, and it's not the same datus. We keep collecting new data.
- Q. It's an ongoing study?

- 10 A. Yes, its been ongoing and now it has been extended into Rhode Island.
- 12 Q. Is it because of how the geography of Cape
 13 Cod or what's --
 - A. That's a good question. The original impetus for the study was that when people looked at the state's cancer registry they saw that the risk of cancer was about 25 percent higher if you lived on Cape Cod compared to the rest of the state by a particular kind of measure, and then the question was why. So we were asked by the state and funded by the state to try to come up with an answer to that question.

So we looked at a number of things.

One of them was cranberry bogs and one of them was one of the original suspicions, which was Otis Air Force Base might be the source of contamination. Relevant to this case, it turned out that there was another source of contamination on the Cape that people sort of knew about but didn't know what the extent of it was.

Q. What was that?

- A. PCE contamination of the water. Now, the really interesting part about this is where that PCE contamination came from. It turned out that it came from the lining of the water mains, which made this an extremely unique situation because it became like a gigantic natural experiment.
- Q. How long did it take to realize it's the lining of the water mains --
- A. It was going on for a full ten years before anybody realized it, and they discovered it by accident in Rhode Island when they did some routine water testing and they found PCE in the water and they couldn't figure out where it was coming from because this

case is typical of where PCE comes from,
water and the air, which is someone throws
it on the ground and it gets into the
groundwater, but they couldn't find any
source of PCE here, and it took many months
for the EPA to figure this out, and here's
what the story turned out to be.

- Q. Were you part of the team that figured it out or was it EPA acting alone?
- A. EPA and Commonwealth of Massachusetts and Rhode Island.
- Q. Okay. Keep going.

A. Sure. So here's what happened. That there is very soft sort of corrosive water in the northeast and the water mains had been coated with sort of tar, asphalt type substance to protect the water mains from corroding and so on, but with this soft corrosive water it was creating color and taste and odor problems. So in the late 1960's, two companies, Johns Manville Corporation and, I think, CertainTeed, who are makers of asbestos cement pipe said "Well, you know, if you're in one of these

areas, we'll give you a new kind of water pipe. We'll coat the inside of it with a plastic," and a plastic is something called Piccotex. It's a resin.

Q. Piccotex?

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Yes, P-i-c-c-o-t-e-x. It's on the outside Α. of milk cartons. So its been tested to be safe for contact with water and stuff like So the question is how do you get this on the inside of the pipe. Well, what they decided to do is dissolve it in PCE and then paint the inside of the pipe with it and under the assumption that the PCE would evaporate and they would have a lined plastic pipe, but there wasn't a big enough market for this pipe so they made the pipe to order, and what that meant was that if you lived in Falmouth on Cape Cod and you were in the water department and you needed to replace the water mains on Oak Street, you ordered 100 feet of water main for Oak Street and within 48 hours of the order they would paint the inside of some asbestos cement pipe and ship it off to

you, so you got freshly painted. They put it in the ground and the assumption was that it would go away. It would dry up and by the time they put it in the ground, it would all be gone. Well, that turned out to be really wrong.

Q. Has it since been remediated?

- A. So they started putting the pipe in in 1969 and by 1979 they figured this out. The amounts in the water were pretty substantial.
- Q. Sorry, the mouths in the water?
- A. The amounts in the water were pretty substantial and they had about 700 miles of this pipe and it was scattered all over the place. Oak Street might have some and then Main Street a mile away might have some for a block or two.
- Q. When you talked about substantial amounts, they were doing tests and coming up with whatever the ratios were?
- A. Yes. It was way over what EPA at that
 point allowed it, and the suggested no
 adverse response level for PCE in water was

40 parts per billion. It's now five parts per billion, and it was way over 40 parts per billion and some of the time it was thousands of parts per billion.

So the way they remediated it was to a systematic program of flushing and bleeding, so they put a tap on Oak Street where this pipe was and they just kept running fresh water through it all the time so it diluted it, basically, until they got it under the five-part per billion level, and they've been doing that ever since. It's still there.

Q. Still being flushed?

A. It's still being flushed, and the pipe is still there but a lot of the PCE now is leached out of the lining of the pipe.

So What does this have to do with us?

So I was on an advisory committee for the

Department of Environmental Quality

Engineering and this issue came before us,

what are they going to do about the pipe

and about the health threat from it. So I

actually, and that's where this flushing

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and bleeding was devised, and so I knew about this and I decided this would be a really good subject for epidemiologic investigation, and we were funded to -- you know, along with all these other possible sources of cancer, this is one that was a lot of interest to me, because one of the things you'd like to do with an epidemiological study is when you make a comparison, you like to compare like with like, and we have this natural experiment here, so we located where all the pipe was from records of the water companies and then we did a big study by comparing cancer of people who had cancer with the pipe in front of their house and people who didn't have the pipe.

Now, I've simplified a little bit because we used a mathematical model actually to estimate the amount of PCE that was leaching out of the pipe, given the diameter of the pipe, the age of the pipe, and when the person moved into their house, so it's quite an elaborate methodology, and

I think we published our first cancer paper on PCE and bladder cancer in 1993 and one of the things that you'll find, if you look at the iris assessment, is that paper is cited as one of the half a dozen with the highest quality exposure assessments.

- Q. I think you said, correct me if I'm wrong, that you're part of the study that helped devise the flushing technique?
- A. I was part of the advisory committee. It all emerged from the advisory committee and the department.
 - Q. Can't they just use different type of pipe?
 - A. Well, they would have to dig up 700 miles of pipe. That would have been the ideal solution, replace the pipe, but that was not possible.
- 18 Q. The other solution you came up with was a flushing technique?
- A. Yeah. Not ideal, obviously, but it did get the levels way down.
- 22 Q. Below the EPA level?

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- A. Yeah, substantially below, actually.
- I should explain something.

Massachusetts is not like Wisconsin or Illinois. In fact, it's not like almost any other state in the union.

Q. Well, nothing compares to Wisconsin.

- A. Well, I'm from Wisconsin so I appreciate that, but in this important respect, which is that every square inch of Massachusetts is in a city or town. There's no such thing as an unincorporated area. Counties, basically, exist only on paper. So there's 351 cities and towns and almost as many water companies, so when you have all these cities and towns on Cape Cod, it's not like you can do one thing to everybody at once. You've got all these small jurisdictions.
- Q. Small jurisdictions, municipalities?
- A. Cities and towns, and that's all there is.

 On Cape Cod it turns out there is a county
 health department but that's unusual.

Probably more than you wanted to know about this.

Q. Thank you. As part of the Cape Cod research, you're also looking at the drinking water aspect and the potential

cancer causing effect of the drinking
water?

- A. That's the study I just described to you.
- Q. It's the same one?

A. Yeah, because the PCE is in the drinking water, and it's in the drinking water if you've got that pipe and it's not in the drinking water if you don't, so that's why this is a giant natural experiment because you might have PCE in your water and your neighborhood in back of you doesn't because, and they didn't have that pipe replaced in front of their house.

So all of these studies about PCE that you see here cited on Page 6, those are all almost, I think every one of them is a study this situation of the PCE coming out of the lining of the pipe.

- Q. As an epidemiologist, you're looking at a natural setting and trying to determine if that natural setting relates to the actual event for which you're researching?
- A. Well, ideally, we like to do an experiment, which is randomly assign people to PCE

contaminated water and not. You can't do that. So you look around in the world for something that's almost like a natural experiment, and this is almost unique in PCE studies. In fact, it is unique in PCE studies because you have almost a natural experiment going on here that you can observe.

You should never ask an academic about his research. You'll never get out of here. I'll just keep talking.

Q. On that note, I have no further questions at this time. Thank you.

MS. KREIL: I have no questions.

MS. ROSS: I just have a couple of questions.

CROSS-EXAMINATION

(By Ms. Ross)

Q. I'm Becky Ross. I represent Continental
Casualty Company and Columbia Casualty
Company.

Were there any opinions that you were asked to provide that you chose not to provide?

84 1 Α. No. 2 Were there any opinions that you formed Q. 3 that you were asked not to provide? 4 Α. No. 5 Are there any plaintiffs in the Class that Q. 6 you believe have not been exposed to PCE 7 through inhalation? 8 Well, I described the information that I Α. 9 was given. On the basis of that 10 information, I can't make a determination 11 about individuals, but it's my opinion as a 12 scientist that they all have substantial 13 potential for exposure, if not actual 14 exposure. 15 That's true of the non-detects, as well? Q. 16 Α. Yes. 17 Thank you. That's all I have. Q. 18 CROSS-EXAMINATION 19 (By Mr. Condon) 20 Q. Did you ever provide the plaintiffs' 21 counsel with an itemization of your time 22 that you spent? 23 Α. No, I don't. 24 Q. Did you bill them?

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        I haven't billed them yet. I just have to
    Α.
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        remember to do that. I'm a horrible
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        business person and I don't do very much
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         litigation anymore. I'll bill him, I'm
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         sure.
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        So you haven't billed him yet. When you
    Q.
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        provide him with a bill, do you have an
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         itemized bill, this amount doing this?
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         It says one and a half days of whatever.
    Α.
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        That's how you normally do it?
    Q.
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    Α.
        Yeah.
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                Thank you.
    Q.
        Okay.
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                               Why don't we reserve
                  MR. MANZKE:
14
         and we can take a look at the transcript.
15
                  (Discussion off the record.)
16
                  MR. CONDON: Condensed and e-tran.
17
                  MS. KREIL: Same, condensed and
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        e-tran.
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                  MS. ROSS: We'll take a condensed
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         and e-mailed.
21
                  MR. MANZKE: Condensed and e-tran.
22
                  (Whereupon the Deposition was
23
         concluded at 12:16 p.m.)
24
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86 1 DEPONENT'S ERRATA SHEET 2 AND SIGNATURE INSTRUCTIONS 3 4 The original of the Errata Sheet has 5 been delivered to Atty. Edward J. Manzke. When the Errata Sheet has been 6 7 completed by the deponent and signed, a 8 copy thereof should be delivered to each 9 party of record and the ORIGINAL delivered 10 to Atty. John Busch to whom the original 11 deposition transcript was delivered. 12 13 INSTRUCTIONS TO DEPONENT 14 1.5 After reading this volume of your deposition, indicate any corrections or 16 changes to your testimony and the reasons therefor on the Errata Sheet supplied to 17 you and sign it. DO NOT make marks or notations on the transcript volume itself. 18 19 REPLACE THIS PAGE OF THE TRANSCRIPT WITH 20 THE COMPLETED AND SIGNED ERRATA SHEET WHEN 21 RECEIVED.

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87
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 1
         M.D.
         CASE: KATHLEEN McHUGH vs. MADISON-KIPP,
 2
                et al.
 3
                          ERRATA SHEET
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                I have read the foregoing transcript
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         hereby subscribe to the transcript as an
22
         accurate record of the statements made by
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         me.
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                             (WITNESS)
                                            (DATE)
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COMMONWEALTH OF MASSACHUSETTS MIDDLESEX, ss.

I, Kelly G. Patterson, a Notary Public duly commissioned and qualified within and for the Commonwealth of Massachusetts, do hereby certify:

That DAVID OZONOFF, M.D., the witness whose deposition is hereinbefore set forth, was duly sworn by me, and that such deposition is a true record of the testimony given by the witness to the best of my skill, knowledge, and ability.

IN WITNESS WHEREOF, I have hereunto set my hand and my affixed notarial seal this 15th day of February, 2013.

Kelly G. Patterson
Notary Public

My Commission expires: September 12, 2014